AUDITORY AND VISUAL SYNCHRONIZATION PERFORMANCE

OVER LONG AND SHORT INTERVALS

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A Thesis

Submitted to the School of Graduate Studies in Partial Fulfilment of the Requirements for the Degree Doctor of Philosophy

> McMaster University September 1983

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AUDITORY AND VISUAL SYNCHRONIZATION

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DOCTOR OF PHILOSOPHY (1983) (Psychology) McMASTER UNIVERSITY Hamilton, Ontario

TITLE: Auditory and Visual Synchronization Performance over Long and Short Intervals

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NUMBER OF PAGES: xiii, 350

ABSTRACT

The thesis is that the same timing mechanism is available and is used by experienced subjects in a synchronization task whether the intervals involved are relatively long (eg 2000 msec) or short (eg 200 msec) and whether the stimuli are auditory or visual.

Synchronization involves two main stimuli. One marks the start of a target interval. The second marks its end. The subject is to ∞ respond in synchrony with the second stimulus.

Previous synchronization research using short auditorily bounded intervals has strongly supported the hypothesis that the underlying mechanism is continuously adjustable, deterministic, and, despite continuous adjustability, that the main source of variability in the task is quantal. A periodic process in the nervous system controls the timing of information transfer from one processing stage to the next. Waiting state durations (before transfers) are uniformly distributed from 0 to g msec, where g is the guantum size. As an example of a timing model along these lines, suppose there is a quantal delay between sensory registration of a stimulus marking the start of an interval and transfer of the message that the interval has started to a "perfect" central clock. In synchronization, the subject waits a specified time after the stimulus, then makes a response. The message from the clock that the required interval has elapsed is subject to another quantal delay before arriving at the motor system, which carries out the response. Variability at the sensory and motor levels may add to the total timing variance in the system, but the main source

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. of variation in extremely experienced subjects is quantal.

Data from various paradigms suggest that different mechanisms govern timing of long and short intervals and that different types of, or at least differentially variable, mechanisms are involved in timing visually and auditorily marked intervals. Both hypotheses have received some support from past synchronization research though they have received much less support from duration discrimination studies that also suggest quantal mechanisms.

There were five experiments, two involving long auditory intervals, one with short auditory intervals, and two with short visual intervals.

All aspects of the thesis were supported: similar practice effects and distributional shapes are found for short auditory and visual intervals. Performance at longer intervals differs from these in variance only, which is concordant with discrimination results, though a specific quantal counting model for discrimination (Kristofferson, 1980) that has had difficulty with previous synchronization data (Hopkins, 1982) has the same difficulty with the present results.

The data support the hypothesis of a common central timing mechanism across modalities and intervals to 2000 msec, and also the idea that boundary conditions on quantal theory found applicable in any given paradigm and modality will also apply to the others.

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There are suggestions in the data that minimum reaction time has been overestimated previously. The data also indicate that a specific model for synchronization (Hopkins, 1982; Kristofferson, 1976) is probably incorrect.

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ACKNOWLEDGMENTS

I am very much in debt to my supervisory committee, A. B. (Kris) Kristofferson. Lee Brooks and John Platt. Each of them provided encouragement and solid advice as well as invaluable technical training and critical discussion during my years at McMaster.

There are many others who have helped mold my approach to research.

Among the past and present McMaster faculty, I thank Ibrahim Ahmad, Ian Begg, Abe Black, Dick Day, Jeff Galef, Woody Heron, Steve Link, and S. G. Mohanty.

Among my undergraduate teachers, I want to acknowledge the important contributions of John Deck, Kurt Danziger, John Gaito, Bruno Kohn, and Hiroshi Ono.

Many of the students and technical staff here collaborated with me and guided me in a variety of projects. In particular, I thank Dan Gallipeau, Mark Hammer, Dave Lang, Renaud Leblanc, John Lyons, Barb Markman, Don Maxwell, Itesh Sachdev, Peter Van Oot, and John Vokey.

Cy Dixon did a superb job of Keeping the machines I worked with alive and functioning, and taught me most of what I know about hardware troubleshooting.

Finally, there are family and friends (along with those mentioned above) who provided tremendous encouragement, support, and not a little guidance. I thank Merry Kaner, Sharon Hafner, Linda Toth, and last, but definitely not least, I thank my parents who also continue to play an important role as teachers.

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1. Introduction: Theories of Subjective Duration

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If any area of research could be taken as suggesting that psychologists have a talent for bringing chaos out of order, time perception must be it. As Woodrow (1930, p.473) put it, "the bewildering confusion concerning the fundamental facts of the psychology of time has existed for so many years that one might believe that psychologists accept the situation as inevitable". Woodrow denied the inevitability of this confusion and embarked on an extensive research program which he expected to "lead to a[°] resolution of the gross inconsistencies and ambiguities which characterize the findings in this field" (1930, p. 473). Unfortunately, he did not succeed. Bemoaning the state of the literature, at least two more modern works (Frankenhaeuser, 1959, p. 12; Ornstein, 1969, p. 16) introduced the field with Nichols' (1891) classic summary:

> Casting an eye backward we can but be struck by the wide variety of explanations offered for the time mystery. Time has been called an act of mind, of reason, of perception, of intuition, of sense, of memory, of will, of all possible compounds and compositions to be made up of them. It has been deemed a General Sense accompanying all mental content in a manner similar to that conceived of pain and pleasure. It has been assigned a separate, special, disparate sense, to nigh a dozen kinds of 'feeling', some

familiar, some strangely invented for the difficulty. It has been explained by 'relations', by 'earmarks', by 'signs', by 'remnants', by 'struggles', and by 'strife's', by 'luminous trains', by 'blocks of specious-present', by 'apperception'. It has been declared a priori, innate, intuitive, empirical, mechanical. It has been deduced from within and without, from heaven, and from earth, and from several things difficult to imagine as of either".

Especially over the past 20 years (since Creelman, 1962), the literature has become mathematically more formal, and the data base has grown considerably. Unfortunately, this has not led to much further agreement about the nature of perceived duration. A number of apparently mutually exclusive hypotheses have been more clearly put, and more strongly supported. My aim in the present research is to extend the realm of application of a specific theory of duration, but I do not mean to imply (and do not believe) that this is a Grand Theory. The timing mechanism explored may be one of many bases for subjective duration, and it may or may not be the main one. To place the work in context, I will begin with a sketch of the dominating perspectives in the field, highlighting what I see as the major differences in assumptions and research interest between them.

1.1 Nontemporal Cue Theories of Subjective Duration

Following a long and popular tradition, Frankenhaeuser (1959)

and Ornstein (1969) argued that our impression of the length of an interval derives from nontemporal information in the interval. There is no dedicated internal process for timing (Frankenhaeuser), or, if there is one, it doesn't play a major role, and studying it won't lead to many interesting or useful results (Ornstein). Instead, some other aspect(s) of the stimulus is used as the basis for an (unconscious) inference about the stimulus duration, such as "the amount of mental content" (Frankenhaeuser) or the amount of information encoded (Ornstein) during the stimulus interval.

The empirical case for the nontemporal cue position rests on demonstrations that differences in nontemporal characteristics of equally long stimuli (such as a different number of events occurring within the stimulus intervals) reliably produce differences in their 🛇 perceived duration. Recent demonstrations along these lines come from Avant and Lyman (1975a, 1975b), Buffardi (1971), Burnside (1971), Cantor and Thomas (1976, 1977), Curton and Lordahl (1974), Long and Beaton (1980a, 1980b, 1981), Schiffman and Bobko (1974), Thomas and Brown (19/4), Thomas and Cantor (1975, 1976a, 1978), Thomas and Weaver (1975), Underwood (1975) and Underwood and Swain (1973), among others (For more, see the reviews of Allan, 1979, Fraisse, 1963, and Poppel, 1971). Not all of these demonstrations implicate information or mental content as the mediating cue to duration, but for present purposes that is a side issue. These studies and their many predecessors demonstrate large effects on perceived duration of nontemporal differences between temporally identical stimuli. Reports of differences in duration

judgments on the order of 35% and upwards are common.

The lesson that I take from such data is that any complete theory of duration perception must include an explanation of the systematic variation of perceived duration due to changes in nontemporal characteristics of the stimulus. But other types of variability in temporal judgments must also be explained. Systematic shifts in perceived duration due to changes in stimulus duration should be quantitatively predictable. The basis of apparently random variation of perceived duration of repeatedly presented identical stimuli should also be made clear.

These latter problems have not been so well dealt with 'in the context of the nontemporal cue research to date. If amount of information encoded determines the perceived duration of an 'interval, what is the function relating information to perceived duration? What rates of encoding should we expect under different conditions, and will these vary with stimulus duration? Without answers to these questions (and the answers are not known), the relationship between stimulus and subjective duration cannot be predicted. Random variability is a more curious problem. Nontemporal approaches are typically concerned with mean perceived duration and differences in mean perceived duration across conditions. Random variation is treated as 'noise', to be minimized, perhaps estimated for purposes of testing hypotheses about mean differences, but to be ignored thereafter. This is such a common approach to research design and analysis that it would not be worth mentioning if this noise for the nontemporal cue theorist was not a

main object of study for most internal clock theorists. To the degree that a clear and accurate description of perceptual variability can be derived from a description of an internal clock, I find it hard to accept dismissals of clocks as unlikely or uninteresting, at least until nontemporal cue theories can provide descriptions of this type as well.

1.2 Internal Clock Theories of Subjective Duration

It is generally accepted that cognitive variables play a significant role in determining spatial judgments. It is not generally accepted that this implies that the visual system is a nonexistent, minor, or uninteresting supplier of information about relative spatial position and extent. It is generally accepted that cognitive variables play a significant role in determining temporal judgments. Does this imply that there are no dedicated physiological systems for supplying us with information about relative temporal position and extent?

Internal clock theorists assume the existence of at least one dedicated physiological system for temporal information, i.e. at least one internal clock. There may be more than one, just as relative spatial position is encoded rather differently by the visual and auditory systems, but there must be at least one. The key problem with this assumption is that we don't know what this internal system is. Various physiological systems have been suggested as candidate clocks, but there is little evidence, and no general agreement, that any one of these is directly involved in duration perception (see, eg.,

Kristofferson, Note 3; Ornstein, 1969).

Internal clock theories share the assumption of a dedicated process for timing but differ regarding the details of the process.¹ In the absence of an obvious and demonstrable physiological basis, description of the clock is most conveniently done in terms of a mathematical characterization of a hypothetical timing system. Such characterizations often include or lead directly to specification of the probability distribution of subjective duration associated with a given stimulus duration, and of the parametric changes in the distribution across durations or across changes in other aspects of the stimulus conditions. If a theory of performance, relating internal distributions to response distributions, can also be specified then fine-grained analyses of the "noise" in the data should provide powerful tests of the hypothesized mechanism. Examples of proposed mechanisms and of the types of distributions implied by them follow.

1.2.1 Creelman's Poisson Counter Model

Perhaps the most common theories of timing, and certainly the first to be quantitatively modelled (Stroud, 1955; Creelman, 1962) are counter models. It is assumed that some internal source produces "ticks" or "pulses" at fairly regular intervals. The count of the number of pulses emitted during the period that the observer perceives a stimulus determines her perception of the duration of that stimulus. The accuracy of the clock is determined by the average interpulse

interval (IPI) and by the variability of the IPI. If there is no variability in the IPI, we have a quantal clock (such as Stroud's), to be discussed below. It seems plausible to suppose instead that the time between pulses is subject to variation (as is, for example, the time between neural spikes), and an exponential distribution is often taken as a fairly accurate description of the distribution of waiting times between simple neural events (see, eg., Luce and Green, 1972). Creelman (1962) made this assumption, along with the assumption that the delays between pulses are strictly independent. These are the main conditions that must be satisfied for a Poisson process to apply (for the others, see Parzen's excellent presentation in 1962).

Predictions from a Poisson process model differ depending on the type of task. In some tasks, the subject is required to make a judgment about the length of a presented stimulus interval. Typically, discrimination experiments are thought of in this way (eg. Allan, Kristofferson and Wiens, 1971; Creelman, 1962), and I will refer to these as discrimination-type tasks. In other cases, the subject is required to generate an interval of a certain duration, or to indicate when a certain amount of time has passed. These include production, reproduction and synchronization tasks, and I will call these production-type tasks. Kristofferson (1977) has shown that in some discrimination settings, the subject actually treats the task as a production task. Rather than deciding, for example, how long a stimulus is, and on this basis, deciding whether the stimulus is "long" or "short", the subject generates an internal duration of a standard

length and decides whether the stimulus is long or short by noting which ended first, the produced standard or the external stimulus. This alternate strategy in some discrimination tasks will be of interest in the discussion of quantal timing below, but I will set consideration of it aside until then. For simplicity of presentation, I will also postpone, for as long as possible, consideration of the role of afferent and efferent latencies in timing. Stimuli are not perceived immediately. The delay between stimulus onset and stimulus registration is the afferent latency. The delay, in a production-type task, between the decision to respond and the completion of the movement marking the end (and/or start) of the interval is the efferent latency. If these are variable, and efferent latency certainly is, modelling of response distributions becomes more complicated.

Under the Poisson process model, the internal measure of the duration of a stimulus corresponds to the pulse count, which is Poisson distributed. Comparison of the durations of stimuli involves comparison of the magnitudes of Poisson distributed variates, and is readily dealt with using the constructs of Signal Detection Theory (Egan, 1975; Green and Swets, 1966). The distribution of produced intervals is not Poisson. In this case, the subject's goal is to produce an interval that contains the right number of pulses. The waiting time between pulses is exponential, and the sum of waiting times to the last pulse required is gamma distributed. This, then, is the production timing distribution. (See Luce and Green, 1972, for many relevant derivations.)

The Poisson and gamma distributions share the property that their variance increases linearly (proportionally) with the mean, so this linear relationship should be obtained whatever the timing task.

Creelman's theory has the potential to predict behavior in a wide range of experimental situations, on the basis of very few parameters, in the simplest case only one, the mean interpulse interval. The theory is mathematically tractable, and the assumptions seem plausible. A number of discrimination studies (eg. Abel, 1972; Creelman, 1962; Divenyi and Danner, 1977; Kinchla, 1972) have yielded results compatible either with the original version of the theory or with slightly modified versions. On the production side, Wing (1973; Wing and Kristofferson, 1973a) found that the variance of repetitive productions of an interval increased linearly with the target interval, as required by the theory. Similar results were obtained by Rosenbaum and Patashnik (1980).

There are also conflicting data, however. When the performance of well-practiced subjects has been studied, discriminability (and thus internal variability) has been found to be constant or non-monotonically (for example, stepwise) increasing with base duration, rather than linearly increasing (Allan, Kristofferson and Wiens, 1971; Allan and Kristofferson, 1974a; Carbotte, 1972; Carbotte and Kristofferson, 1973; Kristofferson, 1973, 1980, 1983; Rousseau, 1975; Rousseau and Kristofferson, 1973). In other cases variability has increased in accord with Weber's Law, which predicts a linear relation between standard deviation and base duration, rather than

between variance and duration (see below). On the production side, Kristofferson (1976) showed in a synchronization task that intervals from 160 through 550 msec could all be produced with the same variance and that the shapes of the response distributions across this range were identical. Beyond 550 msec, variability did increase, but according to Weber's Law, i.e. linear in the standard deviation, not the variance. Both findings are incompatible with a hypothesized gamma distribution. Regarding distributional shape, constancy across durations could only arise in the gamma case if enough pulses were being counted that the total waiting time to the last was normally distributed. Hopkins (1982) found that the convolution of a triangular and a logistic distribution (compatible with a quantal theory, below) described synchronization response distributions better than did the normal.

In sum, Creelman's theory is attractive, and has some support, but if it is not dead wrong, it is often not right.

1.2.2 Weber's Law

Weber's Law describes the variability of performance in discrimination experiments. It does not specify the nature of the timing mechanism involved. The original form of the law specified that the size of a just noticeable difference between a comparison and standard stimulus is proportional to the standard. This is not found in most or all modalities when extremely small or large magnitude stimuli are studied. The more recent form of the law instead specifies

a linear relationship between discriminability and base stimulus magnitude (in our case, duration.) That is, if S is the standard deviation of the probability distribution associated with discrimination judgments about a stimulus or stimulus pair, and M is the magnitude of the standard stimulus, the original form of the law specified that S = KM, for some constant K, whereas the more recent version holds that there is some minimum degree of variability, even when no stimulus is presented, and so S = kM + c, for constants k and c. The constant c reflects the minimum level of variability associated with the particular modality and task. This correction to the law has only recently been given serious consideration in the literature on temporal psychophysics (Triesman_ 1963, and, especially, Getty, 1975). (However, it is not all that "modern" an idea. Fechner, who brought the law into general notice in the first place, was also the first to propose the additive constant (eg. 1860, p. 156). Within the psychophysical literature more generally, Miller's (1947) presentation of the modification is perhaps the most often cited. It is now routidely presented in nontrivial introductions to psychophysical theory (eg. Engen, 1971; Galanter, 1963). The idea of a nonzero minimum for perceptual variability has independently gained almost unanimous acceptance in psychophysics since the advent of Signal Detection Theory (Green and Swets, 1966; Swets, 1961).

Weber's Law can be translated into a statement about the variability of performance: the function which describes the probability that the comparison will be called longer than the

standard, across all values of the comparison stimulus, is called the psychometric function. According to the original form of Weber's Law, the standard deviation of the psychometric function should be strictly proportional to the standard duration. According to the modern form, the standard deviation should be linearly related to the duration (Getty, 1975).

Getty (1975) stressed that Weber's Law does not imply that the standard deviation of internal timing distributions increases with the mean, and gave examples of very different underlying mechanisms that could give rise to the law. The variance of subjective durations might even be constant if other conditions are satisfied. However, if the law is correct, the range of acceptable models for the clock is limited, and does not include quantal clocks or Creelman's Poisson process (though Divenyi and Danner modified Creelman's theory, added a further parameter, and were able to accommodate the law in this case.)

Weber's Law is an empirical generalization about discrimination performance. It does not tell us anything directly about production. However, if the standard deviation of the clock were found, in a production setting, to be linear in the mean duration produced, we would expect that this timer, when used in a discrimination experiment, would yield results quite compatible with Weber's law.

Fraisse (1963) cites a long history of rejection of Weber's Law for duration, beginning with Nichols in 1890. Woodrow (1951) considered the case against Weber's Law conclusive. Allan and Kristofferson (1974b) and Allan (1979) also reviewed evidence against

the law. Findings such as those of Rousseau and Kristofferson (1973), who reported constant discriminability across a range of base durations from 100 to 2000 msec, are incompatible with any version of Weber's Law. There is so much evidence against Weber's Law, and there has been for so long, that one might be amazed that temporal psychologists are still willing to talk about it seriously.

There is stM1 some room for argument, however. Fraisse (1963) noted that the original form of Weber's Law (proportionality) seemed to fit fairly well in the range from 200 to 2000 msec. The problem of fit for short durations is comparable to that found for weak stimuli in modalities such as vision and hearing, which led to the linear generalization of Weber's Law (a point raised more explicitly by Triesman, 1963 and Getty, 1975). As to longer stimuli, Fraisse argued at length that intervals longer than about 2 seconds are estimated (perceived duration is determined by cognitive factors) rather than gerceived directly. The law breaks down beyond 2 seconds because a different type of timing has come into play, with different sources of variability. Working with stimuli within this range Alpern and Darwin (1982), Diveryi and Danner (1977), Getty (1975, 1976), Kristofferson (1976, 1980), Thompson, Shiffman and Bobko (1976) and Treisman (1963, who included stimuli out to 9 seconds) have all reported data which appeared compatible with Weber's Law. For present purposes, the work of Getty (1975) and of Kristofferson (1976, 1980) are most interesting.

Getty (1975) studied the performance of fairly well practiced subjects in a discrimination experiment covering base durations from 50

to 3200 msec. Weber's Law failed for durations longer than 2000 msec but accounted for shorter duration performance quite convincingly. Subjects made 330 judgments at each base duration. The base duration was constant within sessions, but was changed randomly across sessions. Kristofferson's (1973, 1980) study can be viewed as a conceptual replication of Getty's work. There were numerous procedural differences -- Kristofferson used a many-to-few procedure and provided subjects with feedback, whereas Getty used a two alternative forced choice procedure without feedback -- but the results were remarkably comparable in spite of this. Getty reported Weber ratios (standard deviation / base duration) on the order of .05 to .06. Kristofferson's ratio, for the first 5 sessions at each duration, averaged .053. The Key difference between Kristofferson's and Getty's procedure is one of practice. Kristofferson's subject stayed at each base duration for 20 consecutive sessions (6000 judgments per base duration) rather than for 1 session. While Kristofferson's results from the first 5 sessions per base duration are very similar to the results from Getty's first (and only) session per duration, Kristofferson's results for the last 3 sessions per duration are very different. After extended practice, variability no longer increased smoothly with base duration. Instead, it increased in a stepwise manner, the standard deviation doubling at base durations of 200, 400, and 800 msec. Within these subranges, the variances were not perfectly constant, but as Kristofferson suggested then, with even more practice it has been found that they become constant (Kristofferson, 1983). Allan and Kristofferson (1974a) also

stressed the need for extended practice at each base duration if results of constant or stepwise increasing variability are to be obtained.

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The other result reported by Kristofferson (1976) that is compatible with Weber's Law was obtained in a synchronization, or production-type setting, rather than in a discrimination experiment. Synchronization will be described in detail below. For now it is sufficient to note that the subject is required to produce a given interval, i.e. to wait for a specified period of time before responding, rather than to discriminate between stimuli. As in Kristofferson's (1980) discrimination experiment, the subjects practiced ad tedium, producing only one interval per session (400 times per session), for many sessions. Variability was constant in the region of 150 to 550 msec. Beyond 550 msec, the variance of synchronization response latencies increased continuously, the standard deviation being proportional to the mean latency less 550 msec. A timer whose standard deviation increases linearly with duration should yield a psychometric function whose standard deviation increases linearly with base duration in discrimination experiments, so if the timer underlying synchronization performance were also used in a discrimination setting, Weber's Law would be supported in that experiment.

In contrast with the discrimination results of Kristofferson (1980), lack of practice at specific intervals was not the cause of increasing variability with duration beyond 550 msec. The subject who

showed this effect (only one synchronized at long durations) was most practiced at 940 msec, having started the experiment with 60 sessions (24,000 responses) at this interval. Kristofferson concluded that there must be a shift in timing mechanisms used in the task at about 550 msec. For shorter intervals, a deterministic timer was used. For longer ones, a model assuming deterministic timing for the first 550 msec and variable timing for the rest of the interval provided an excellent description of the data.

Kristofferson's 550 msec transition point is of interest because a similar one was found by Michon (1967) in an entirely different type of experiment, magnitude estimation, at about the same duration. Michon found that the exponent relating judged to physical duration changed abruptly between 500 and 600 msec. Tying this to introspective reports that the subjective experience of duration changes qualitatively in the 500 to 1000 msec region, to a peak in discriminability often found in the 600 to 750 msec region, and to the common finding that intervals shorter than about 700 msec are overestimated, whereas intervals longer than this tend to be underestimated (all reviewed in detail by Fraisse, 1963), Michon argued that the timing processes used for very short (below 600 or so msec) stimuli are different from those involved in timing slightly longer ones. Kristofferson's (1976) result appears to support this suggestion strongly.

It is exceedingly curious that the same subject, studied in the same lab by the same experimenter, who used basically the same

equipment, should yield synchronization data that suggest a different timer for short and slightly longer stimuli while also yielding discrimination data (Kristofferson, 1980, 1983) that suggest precisely the reverse conclusion. Comparison of long and short duration synchronization performance is one of the foci of the present research. For now, however, I'll set that aside and return to the issue at hand. Curious it may be, but synchronization performance from an extremely well practiced subject indicates that Weber's Law describes the variability of the timer in this task from 550 to 2190 msec. Treisman (1963) and Getty (1976) also found evidence for Weber's Law in production-type tasks. In Getty's case, a number of statistics of the data were also examined for compatibility with the Poisson process. The hypothesis of a linear relationship between internal variability and duration was strongly favored over that of proportionally increasing variance with duration.

1.2.3 Quantal Theories

The final class of theories to be considered here is the quantal class. These form the theoretical context within which the present research was conducted and so will be presented in greater detail.

Quantal theories assume, as does Creelman's counter theory, that timing is determined by a periodic internal process. The Key difference in assumptions is that the interval between pulses is assumed fixed in the quantal case. The interpulse interval's constant

value is an indivisible unit of time (a time quantum), and is often represented by q. Quantal hypotheses have been around for a long time. For references to the older literature, see Harter (1967), Poppel (1971) and Stroud (1955). I will focus exclusively on the work of Stroud and of Kristofferson and his colleagues.

1.2.3.1 Stroud's Perceptual Moment Hypothesis

Stroud (1955) postulated a counter theory quite similar in conception to Creelman's later (1962) counter theory. As noted by Creelman, the only formal difference between them was that Stroud assumed no variability in the interpulse interval whereas Creelman assumed an exponentially distributed interval. In Stroud's theorizing, pulses occur every q msec, and they are used to assign internal "dates" to stimuli. Although Stroud didn't make this point explicitly, under this view the subjective duration of an interval should be directly determined by the number of pulses contained within it.

The fact that the interpulse interval is fixed does not imply that timing is free of variability. Consider a production task first. The subject is presented with a stimulus, and is to press.a button at some specified time after the onset of the stimulus. The subject achieves this by counting an appropriate number of pulses before responding. The physical response takes time to execute and this time is definitely variable (Wing, 1973, 1980) but as above, for now I will ignore efferent and afferent latencies, focussing exclusively on timing. If the subject starts counting pulses from the first one after

registration of the stimulus, he misses the delay between stimulus registration and that first pulse. The delay may be zero, i.e. the pulse might occur right after stimulus registration, or the delay might be q msec, if the stimulus is registered right after a pulse. Pulses occur every q msec independently of external stimulus conditions, so stimulus onset (and registration) may take place at any time within this range relative to the time of the next pulse. The delay is uniformly distributed, between 0 and q, as, therefore, must be the interval between stimulus onset and the time of the last counted pulse. Timing variability in a production task should thus be uniformly distributed, with a variance of $q^2/12$. (Note that this assumes, as has all discussion of counter theories above, that the counting mechanism works perfectly. If the counter misses the odd pulse, the situation becomes more complicated, a point raised by Kristofferson (1980).)

Turning to the internal measure of the duration of a stimulus, suppose as an example that the time between registration of stimulus onset and stimulus offset is 2.6q msec. If the first internal pulse occurs less than .6q msec after registration of the onset of the stimulus, three pulses (bounding 2q msec) will occur during the stimulus. If instead the delay between registration and the first pulse is longer than .6q msec, the stimulus interval will only contain two pulses (1q msec). In general, if an internal interval lasts T = nq+ m msec, where n is the largest integer less than T, and so m = Tmodulo q (by definition, the remainder of integer division of T by q),

then either n or n+1 pulses will be counted. The count distribution is thus point binomial and the probability of a count of n is m/q. The variance of the count is $m(q-m)/q^2$. This is not a direct function of the duration of the interval, but it is a function of m, the interval mod q. Examined over widely spaced values of T, the variance should appear roughly constant with duration. In fact, though, it is periodically related to T, with a period of q.

Various early applications of this model to studies of reaction time, successiveness discrimination, and other tasks appeared successful (Augenstine, 1955; Kristofferson, 1966b, 1967; Schmidt and Kristofferson, 1963; Stroud, 1955), but with complications. First, there were conflicting estimates of the quantum size. Kristofferson (1966a, 1966b, 1967a, 1967b) typically obtained values around 50 msec. Kristofferson (1969) obtained values closer to 25, as did Heath (1972). Stroud (1955) estimated q at about 100 msec. . . These differences were not due to statistical instabilities. Kristofferson obtained remarkably similar results Across' tasks, as did Stroud. Augenstine (1955) conducted a spectral analysis on response time data from two tasks and found two levels of periodicity, at 50 and at 100 msec. (A third period, of 265, was also obtained in the first task). Kristofferson (1967a) presented evidence that successiveness discrimination involved a mixture of quantum levels. Sometimes the subjects operated with a q of 50 msec, sometimes with a q of 100. This was particularly a problem with relatively unpracticed subjects. Kristofferson (1981) reports that early reaction time work of his also

yielded estimates of 25 and 50. This type of result was eventually used to advantage, as strong support for a different quantal theory (Kristofferson, 1980, discussed below), but at the time these results were essentially anomalous.

More important than the lack of agreement on a single value for q was the consistent finding that this quantal counting theory did not predict duration discrimination performance at all (reviewed by Allan and Kristofferson, 1974b). Internal duration distributions did not appear to be uniform. Instead, Allan and Kristofferson (1974a; Allan, Kristofferson and Wiens, 1971; Kristofferson, 1973) found that discrimination performance was well described if the internal distributions were assumed to be triangular, not uniform. Triangularity makes no sense in terms of the original quantal counting model. Nor does the finding that subjects can set a response criterion, or vary a response latency, across a continuous range (eg. Kristofferson, 1976, 1977). If a subject's indivisible unit of time is 25 msec, how does he shift a mean response time by 5 msec? And that without otherwise affecting the shape of the response distribution in any way?

The finding that intervals generated by subjects are continuously adjustable is a nasty one for any quantal counting theory. Some continuous process has to be accommodated within the framework of a fundamentally discrete system, and in a plausible way, if the quantal view is to be maintained. If the main function of the periodic process proposed to underlye the quantum is timing, and the main mechanism

supposed for timing is the assignment of "dates" to events in terms of these time points, continuous adjustability seems to me to be critically negative evidence. Accordingly, I will set aside this particular quantal view, Stroud's "perceptual moment" hypothesis. (See also Allan, 1975, Allport, 1968, and Baron, 1971.)

1.2.3.2 Contical Excitability Cycles

Kristofferson's (1966a, 1966b, 1967a, 1967b, etc.; Schmidt and Kristofferson, 1963) point of view was somewhat different from Stroud's. It was hypothesized that the time guantum reflected a processing limitation of the nervous system, not a dedicated process for conscious timing. According to this "cortical excitability cycle" hypothesis (reviewed in Harter, 1967), certain types of jobs can be done in the nervous system only at specific times, minimizing the total processing load at any instant. Thus, for example, attention switching might only be possible once every q msec. Sampling of input in an attended channel might also occur once every q msec, perhaps a few msec after the time at which attention switching to a channel can be completed. (This particular case was raised by Kristofferson, not by Harter. See Santa-Barbara's 1967 review of the physiological thinking behind Kristofferson's quantal theory.) The time quantum reflected in human timing is the period of whatever process it is that regulates switching and/or sampling times. The process may send out one signal per cycle that can be counted, and if it does, and these signals are counted, we have a quantal counting clock. However, that is not the
only type of clock possible under this view, and the quantum may be reflected in timing whether or not counting takes place. Suppose there is a "perfect" clock in the nervous system. Messages to the clock to start or stop timing, and messages from the clock that an interval is complete, or that an interval lasted so long, still have to be sent and received in the system. If messages to and from the clock can only take place once every q msec, then the resolution of timing is not infinite, as is this hypothetically perfect clock's, but instead it is determined by the quantum size.

The central idea in Kristofferson's quantal theorizing is that messages (eg. stimulus events) are processed in stages, and a periodic process determines when a message may proceed to the next stage. A number of different timing hypotheses have grown out of this, at least one of which is probably wrong, and more than one of which might be right, depending on the situation and the strategy of the subject. Before presenting those, I should note explicitly that EEØ alpha, with a frequency of 20 cycles per second, is very frequently discussed in presentations of periodic processes and timing. EEG figured prominently in Kristofferson's early work and thought (Kristofferson, Note 3), and in many other discussions. The research linking timing and EEG has produced a rather underwhelming case for this idea, however (for example, Holubar, 1969; Legg, 1968). The present discussion is without reference to any particular periodic process in the nervous system.

1.2.3.3 The Onset-Offset Model

The first quantal model for duration discrimination (after simple quantal counting failed), was the onset-offset model of Allan, Kristofferson and Wiens (1971). Suppose that there is time jitter of up to q msec before a stimulus is processed centrally. There are two sources of variability in stimulus presentation: the onset of the stimulus interval has to be registered, and the offset of the interval has to be registered. If the onset and offset delays are uniformly distributed, and are independent, their sum is triangularly distributed, with a variance of $q^2/6$ (see also Kristofferson, 1966b). If there is no other variability in the timing process, then subjective durations associated with stimuli should be distributed in the same way. This model also predicts that timing variance should be constant across the range of intervals over which onset and offset delays are the only sources of timing variance.

Allan et. al. found that the assumption of equal variance triangular distributions predicted the operating characteristic curves of their subjects' duration discriminations quite well, at least as well as an assumption of equal variance normal distributions, and much better than Creelman's model, which implies proportionally increasing variance in the internal durations. Quantal estimates from the data were either very close to 50 (4 between 46.1 and 52.1) or very close to 25 (5 between 23.0 and 26.5) with one exceptional value of 37.6. The numerical values thus appeared quite comparable to estimates from

experiment, the range of durations studied was rather narrow (50 - 150 msec), but results in some ways comparable to them were obtained by Allan and Kristofferson (1974) using a range of stimuli from 70 to 1020 msec. No subject worked with the entire range of stimuli, and for identical stimuli more cross-subject variability was evident than in Allan et. al. (1971). Further, variability was not (constant across the ranges studied for most subjects, and variability definitely appeared to increase with duration. However, variability did not increase continuously with duration. Instead, it seemed as though variability might be constant over subranges, with sharp increases to new levels, which would remain constant over a further range of durations. Estimates of q were again close to previous values, 25, 50, or 100, excepting data from one subject (of 5 in 1974) that yielded a q estimate close to 150. Both of these studies used light flashes as the stimuli. Rousseau and Kristofferson (1973) used the offset of a light and the onset of a tone to mark the boundary of an interval, and obtained equal stimulus discriminability across a range from 100 to 2000 msec. Discriminability in this study was poorer than previously, but the evidence of constant variability in the timing process seemed quite strong. Allan and Kristofferson (1974b) and Allan (1979) réview more of the same types of results.

The onset-offset model was explicitly dissociated from the hypothesis of quantal counting. Subjective duration was seen as a continuous variable, not a quantized one. Variance is due to periodic (quantal) delays in getting information to the clock, which,

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implicitly, is assumed to keep perfect time once it gets the message. This approach preserves the core of Kristofferson's quantal theorizing, that a periodic process is a major source of timing variability, while avoiding the complications inherent in quantal counting models.

Despite its success in accounting for some aspects of the data, there are a number of problems with the onset-offset model. First, there is a conceptual problem with the notion of independent uniform delays of the onset and offset stimuli, a problem raised in detail by Hopkins (1982). The rationale behind the quantal delay is that a periodic process determines when a stimulus representation can be dealt with centrally. Since the stimulus can be presented at any time relative to the start of a cycle of this process, we can assume that the distribution of waiting times before the stimulus is dealt with (eg. before a message to start timing or to stop timing and report the length of the interval) is uniform. But by the definition of the process, the times between successive "ready" periods are fixed. Similarly, the time between stimulus onset and stimulus offset is fixed, for identical stimuli. Once we know what the waiting time to process the first stimulus was, and what the value of q is, then for any stimulus duration we should be able to predict perfectly what the 🛹 waiting time to process the offset stimulus will be. Both delays uniformly distributed, but they can't be independent.

In the context of a model for synchronization performance, Hopkins (1982) made a more satisfactory suggestion. In the synchronization task, the subject is presented with a short stimulus

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pulse (P1) and is to make a response in synchrony with a later pulse (P2). In order to synchronize with P2, since the response movement takes time (Michaels, 1977, estimated that the triggering of the response starts about 100 msec before it is completed), the subject must estimate how long it has been since P1 and respond on the basis of that estimate, not on the basis of perception of P2 itself.

Synchronization is a timing task, not (usually) a task involving reaction time to P2. The stimulus-response chain can be divided into three stages, afferent, central (including timing), and efferent, with two quantal delays, one between initiation of timing after the stimulus is registered, and one between initiation of the response after the timer indicates that responding should begin. So far, this is identical to the thinking behind the onset-offset model, except that in that case there is no central-to-efferent delay as there is no timed response. The two delays in discrimination are afferent-to-central delays. Hopkins' contribution was the suggestion that the periodic process controlling output from the clock might(have a trivially different frequency from the process controlling input. If so, the two processes will pass through all possible phase relations, and knowing the waiting time for input to the clock will yield no information about the waiting time for transfer to the motor centers, so the sum of the waiting times will still be triangular."

Timing in the nervous system is not determined by a single crystal clock, as it is in a computer. Instead it is determined, presumably, by the actions of a number of cells. The idea that

transfer from the clock to the motor system would involve different groups of cells from transfer from the sensury system to the clock seems plausible enough. The idea that different groups of cells would be involved in periodic regulation of transfer times also seems plausible to me, and if so, the notion that the frequencies of the processes might differ seems quite reasonable. Stretching the argument back toward discrimination, if a second periodic process determines the waiting time from the clock outward, whether to the motor system or to. anywhere else, we have a second, effectively independent, quantal stage that does not require the difficult hypothesis that waiting times of time-locked stimuli into the clock are independent. I'm not fully satisfied with this argument, but this is the best suggestion available for triangularity in the discrimination task. It should be noted that not only does triangularity fit the various statistics from duration discrimination data quite well (Allan et. al. 1971; Allan and Kristofferson, 1974a; Brewster, 1983; Kristofferson, 1973, 1980, 1983), but it describes the data at least as well as or better than a normal distribution (Allan et, al., 1971; Brewster, 1983). In the synchronization situation, Hopkins (1982) compared the fit of a triangle convoluted with a logistic distribution (effecent latency variability) with that obtained from a variety of other distributions, and the triangular hypothesis was consistently supported. In sum, whether the specific mechanism proposed in the onset-offset model is reasonable, triangularity is empirically well established, and some quantal stage approach should predict it (which no other theory does).

Far more difficult for the onset-offset model is the variation observed in q. If the only source of variability is in input to and/or output from the clock, and the clock times in continuous units, then why does the variance increase with duration, stepwise or otherwise? There is no good reason for this and Allan (1979) and Kristofferson (1977, 1981) have abandoned the model.

Research since the proposal of the onset-offset model in 1971 has pointed to two Key revisions to this way of thinking about discrimination behavior. First, the onset-offset model assumes that subjects actually estimate the duration of the stimulus, and that they base their discrimination judgments on this estimate. This is called the interval measure hypothesis and it parallels standard assumptions about the basis of discrimination in any modality (eq. Thurstone, 1959; Green and Swets, 1966). At least for the tasks under study, this hypothesis is demonstrably incorrect. The alternative that does describe the data is the real-time criterion hypothesis (Kristofferson, 1977), which I describe in the next subsection. The second revision has been the acceptance of quantal counting, strongly suggested by the finding of very orderly stepwise increases in the variability of internal times. The problem for quantal counting was and is the finding that subjects can produce intervals along a continuous scale. Since the problem is critical for the hypothesis, I'll review the evidence for it, and a suggestion for its basis, before describing the evidence leading to quantal counting.

1.2.3.4 The Real-Time Criterion

In a single-stimulus discrimination paradigm, the subject is presented with, and asked to categorize, stimuli one at a time. In a given session, you might present many different stimuli, having the subject state, after each is presented, whether it is "short" or \nearrow long". The midpoint of the stimulus set is usually used by the experimenter as a dividing point, and the subject's judgment, refined with trial-by-trial feedback, is to be based on whether the stimulus is longer or shorter than that middle duration. According to the interval measure hypothesis, in this type of task, the subject compares the perceived duration of the stimulus with some criterion value, and says "long" if the duration is (too large. Now, suppose that you present a subject with a long stimulus, and you discover that she consistently responds "long" before the stimulus interval is complete. How she does this is not clear at this point, but how she does not do it is by measuring the duration of the stimulus interval, basing her judgment on (the difference between this interval measure and a criterion. (The · interval has not yet ended so how could she have measured it?) The interval measure hypothesis cannot deal with this kind of behavior, but this is what happened in numerous discrimination experiments.

Kristofferson and Allan (1973) outlined the real-time criterion hypothesis as an alternative and Kristofferson brought data to bear on the issue in 1977. The idea is that the subject turns the task into a time production task, generating a criterion interval, and bases her judgment on which ended first, the stimulus or the criterion. As soon

as one of them ends, she can immediately respond, either long if the criterion interval ended first, or short if the stimulus ended first. It is called the "real-time" criter for Decause it is assumed that the subject is working in "real" time. With training, she learns to generate an interval of the required length. How long that interval seems subjectively doesn't matter. All that matters is that it is the right length, and, given that, whether it or the stimulus ends first.

To test the idea, Kristofferson examined subjects performance in a speeded discrimination paradigm. In this case, the duration stimulus was an empty interval, bounded by two short (10 msec) auditory pulses, P1 and P2. The subject was instructed to respond "long" or "short" as quickly as possible. (To avoid response competition problems, in any given session, subjects either made "long" responses, doing nothing when the stimujus was judged short, or they made "short" responses. Many sessions of each type were conducted.) Three predictions can be made about performance in this task. First, "long" response's should be time-locked to P1, the stimulus marking the start of the interval, and the "short" responses should be time-locked to P2. By "time-locked" I mean that the distribution of response times should depend only on the time since the stimulus to which they are "locked". "Long" responses are triggered by the criterion, and should occur at the same time no matter when P2 is, so long as P2 comes after the criterion interval ends. "Short" responses are made when the stimulus ends (P2 happens) before the criterion interval ends. They should be simple reactions to P2, and should not depend on P1. The second

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prediction is related to the idea of time-locking: the distribution of "short" responses should look like a distribution of reaction times, whereas the distribution of "long" responses should look like a distribution of time production responses. The final prediction is that q should be estimable from both the response probabilities and from the response latencies, and the same estimates should be obtained in both cases.

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Kristofferson and Allan (1973) can be consulted for the details of estimating q from discrimination probabilities. As to the response latencies, Kristofferson (1977) made the following argument. If a "long" response is time-locked to P1, its latency includes the afferent latency (the delay between stimulus onset and stimulus registration), the duration of the critigrion interval, and the efferent latency (the delay between the end of the criterion interval and the completion of the response). The latency of the short response, as measured from P2, consists of the afferent latency associated with P2 plus the efferent latency, the delay between registration of P2 and the making of the fast reaction response. Assuming that the variances of the afferent latencies in both cases, and those of the efferent latencies in both cases, are the same, the variance of the internal criterion interval is simply the variance of the *long* response latencies minus the variance of the "short" response latencies. If internal timing distributions are triangular, their variance should be $q^2/6$, so the difference in variance between latencies of long and short responses should yield an estimate of q directly.

It is important to note that there is no way of filtering out afferent response latencies in the discrimination case. Estimates of variability (and thence of q) in duration discrimination are based on response probabilities, which reflect the total variability in the task. If there is afferent latency variance, the estimate of q from discrimination should be higher than the one we obtain from the response latencies.

One question that is preliminary to research of this type, by the way, is whether or not subjects <u>can</u> produce an internal criterion interval with any degree of stability. Kristofferson (1976) addressed this using a synchronization paradigm. In this case, the subject is not required to make a discrimination response. She is merely required to respond a specified time after P1. P2 is presented as a marker of that time, i.e. her¹goal is to respond in synchrony with P2, and the earliness or lateness of P2 relative to the response serves as immediate feedback on each trial. Subjects perform this task well and their response distributions look triangular.

The results from Kristofferson (1977) were as expected. My description will ignore results from stimuli that fell within one q of the base duration used (1160 msec) as examination of these would introduce complications that are not relevant here. (They are not discordant with the theory). The response distributions to the "short" stimuli were supposed to be reaction time responses to P2, and were not supposed to depend on the P1-P2 interval. They were, and they didn't. On average, responses followed P2 by 168 msec. Kristofferson (1976)

estimated reaction time to similar stimuli at about 160 msec. Response distributions to P2, at three different P1-P2 intervals, looked the same in shape as well as in mean (see Kristofferson's Figure 4, in which they are superimposed). The distributions were highly peaked, fairly symmetrical, and comparable in variance to those obtained from the same subject in synchronization (Kristofferson 1976) when the synchronization response was to follow P1 almost immediately (i.e. when P2 was in the reaction time range after P1). The distribution of "long" responses was time-locked to P1. The mean response latency from P1 was 1340 when P2 followed P1 by 1330 msec, 1343 for a P1-P2 interval of 1375 and 1347 when P2 followed P1 by 1550 msec. The distributions were again symmetrical and superimposable. The estimated q from the latencies was 95.5. From the response probabilities, the (entirely independent) estimate was 95.3 msec. To instantiate the meaning of "well-practiced subjects", since I've raised the issue of practice before, these data were collected over 114 sessions of 300 trials each, using the same base duration in each session. Data reported in the same paper from subjects who had only participated in 60 and 64 sessions, respectively, were more variable, and the subjects had clearly not attained asymptotic levels of performance, as is required by the theory for any stringent tests.

Kristofferson (1983) extended these results, on the same subject, across more base durations, obtaining highly comparable q estimates from the latencies and the response probabilities across conditions, and also obtaining q estimates highly comparable to those

found, from the same subject, in non-speeded discrimination (Kristofferson, 1980). As such, they also provided further support for the quantal counting hypothesis, which cannot be taken seriously until the problem of continuous adjustability is cleared up.

1.2.3.5 Deterministic Internal Delays

The criterion duration can be estimated from response probabilities in standard duration discrimination data, and Kristofferson (1973) noted that these seemed rather more precisely glaced than would be expected if the subjects were working in quantal (eq. 25, 50 or 100 msec) units. Without downgrading the value of such data, I'll note that inferences about the shape and mean of internal distributions that are derived from discrimination are necessarily rather indirect. All that you have to go on is a large number of binary decisions (long, short) associated with a few, sometimes very few, stimuli. Many assumptions are involved in estimation of criterion durations from such data. More direct evidence for continuous adjustability comes from production tasks. If the subject can accurately produce any duration specified, withgut changing any statistics of his response distribution other than the mean, from interval to interval, the conclusion that he is working with a Continuous scale is inescapable. Kristofferson (1976) obtained results of this type using a synchronization task.

To illustrate the problem this poses for a quantal counting theory, I'll present a hypothetical example. As above, to keep the

(discussion to the point, I will ignore afferent and efferent latencies and their variances. Suppose that q is 25 and the timing distribution is triangular. The variance of the response distribution is g2/6, or 104 msec². If we present the subject with a stimulus, P1, and ask him to respond precisely 300 msec later, he should have no trouble with this, because it is exactly 12 quanta from P1. Similarly, synchronizing with a P2 tht occurs 325 msec (13 quanta) after P1 should also be easy, the variance should be the same, as should be the response distribution's shape. Now, suppose we make the P1-P2 interval 312.5 msec. This is 12.5 guanta, but, guanta being indivisible, the subject will have difficulty counting half of one. He can produce a mean of 312.5 by counting 12 quanta half the time and 13 quanta the other half of the trials, but now his response distribution will have a flat top rather_than a sharp peak and his variance will be much larger. If his timing is based strictly on quantal counting, he cannot produce an average 312.5 msec interval in any way but by producing a mixture distribution of responses, some based on overcounts and some on undercounts. Similar problems arise, of course, for 305, 310, 315 and 320 msec intervals. (In these cases, the distributions have unique 🗫 modes but they are asymmetrical.)

Kristofferson (1976) found that synchronization response distributions had the same shape and variance from 170 through 550 msec. Variances throughout this region were about 125 msec². Many intervals studied differed by only 5 msec. The mean latencies were very close (within a few msec) to the P1-P2 intervals at every interval

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studied. I should qualify this by noting that the variance steadily increased, for one subject, as the reaction time limit was approached, and was only convincingly constant from 300 to 550 msec. The shorter the P1-P2 interval below 300, the more difficult the task for that subject, and the more variable his responses. However, small differences in the interval produced small differences in variability, and the means still tracked the intervals closely. All distributions were unimodal, bounded, symmetrical, and very highly peaked. They were not produced from mixtures of counts of different numbers of quanta.

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The response distributions were not exactly triangular, nor should they be. Efferent latency variance may be constant from interval to interval, but it is not zero. Wing (1973) developed a means, in a repetitive Keytapping task, of estimating efferent latency variability, which he manipulated by changing the required response movement from condition to condition. In well-practiced subjects, motor variance is about 10 to 50 msec². (Similar estimates come from Meijers and Eijkman, 1974, and Vorberg and Hambuch, 1978.) There is no strong evidence regarding the distribution of efferent latencies, but there are some grounds for supposing that the distribution is bell shaped. Meijers and Eijkman (1974) make a convincing case that the variability of motor delays reflects the contribution of delays from a large number of cells, in which case the delays might be normally distributed. Hopkins (1982) reviewed this literature in somewhat more detail, and used a logistic distribution as an approximation to the normal in examining synchronization response distributions, and testing the fit of the quantal model. Subsequently (during his thesis defence) he has stated that with further reading and discussion he thinks the logistic model might be more accurate than the normal. It doesn't matter all that much in the context of the present research which finite-variance bell shaped distribution we use for the efferent latencies. I will assume, with Hopkins, that it is logistic. The distribution of synchronizations under this view should be the distribution of a triangular variate added to an independently distributed (Botwinick and Thompson, 1966, for example) logistic. Plus, of course, a random variable representing afferent latency, unless that isn't variable, and it seems not to be, at least for auditory and tactile stimuli.

Synchronization variance is constant from 160 to 550 msec and there is no difference in the shapes of the distributions across this range. Delaying a reaction time range response (160 msec) by up to 400 msec has no effect on the response distribution, except in the mean. Given the lack of effect on variance and shape, it is most plausible to assume that these internal delays are not themselves variable, i.e. that they are deterministic.² But these delays are not only deterministic; they also seem to be continuously adjustable, at least for auditorily bounded durations. Quantal delays are deterministic, but they are not continuously adjustable. Where are <u>these</u> coming from?

Kristofferson (1976) speculated that it might be in the afferent latency. The introspective basis for this hypothesis was the report of perceptual "shortening" of stimulus intervals by experienced

subjects when working with relatively long intervals. At the very start of a session, the subjective duration associated with a P1-P2 interval was reported to be much longer than that associated with the same interval a few trials into the session. Once the subject was "warmed up", the interval seemed fairly short, and, subjectively, it stayed that way. This suggests that the stimulus is being delayed before it reaches the clock. As a subject who became very experienced at a P1-P2 interval of 460 msec in Joan Brewster's synchronization experiments, I should report that this does not describe my impressions at all. Nor, from reports from the two subjects in the present research who synchronized in the 1400-2100 msec P1-P2 range, does it describe their subjective experiences. However, it was striking for Kristofferson's subjects and was the basis for a plausible suggestion independently of the introspective reports.

If we parse the synchronization stimulus-response chain into afferent, central and efferent components, then a deterministic delay isn't likely to be part of the efferent latency, which is not deterministic at any level studied (see, eg. Meijers and Eijkman, 1974). If by central we mean a discontinuous quantal clock, it can't be here. If there is no variability in the afferent latency, then at this level, perhaps the stimulus could be recycled through some part of the afferent mechanisms before it is brought to the clock. If afferent latency variance is zero, this is the most likely candidate for the delay, even if the clock is not quantal (especially if the clock is variable, with increasing variance with duration).

One source of evidence for non-variable afferent latencies is the finding that in speeded discrimination the latency based estimates are the same as the discrimination based estimates. If afferent latency variance was present, q estimates from the response probabilities should reflect this, and be higher. Certainly, there is room for such variability to be missed when q is about 100, since timing variance at this level should be more than 1600 msec² (q²/6). But when q is only 23.9, as in Kristofferson (1983). variance is only 95.2, which is more constraining. When synchronization latency variances are as low as 35 msec², as obtained by Hopkins (1982; Hopkins and Kristofferson, 1980) after exorbitant amounts of practice, there is barely enough for us to \sim include efferent latency variance. If q is as low as 12.5, timing variance should be 26 msec². There is simply none left over for afferent latency variability to account for. Further, Hopkins (1982) tested the fit between these data and his model for response latencies that specified that they should be distributed as the sum of a logistic efferent latency variable plus a triangular timing variable, with no provision for afferent variability. The fit of the model to the data was excellent. Divenyi and Danner (1977) developed an extension of Creelman's (1962) theory that allowed them to estimate auditory afferent latency variability in their task. Their estimate was 1 msec². In a separate review, Divenyi (1976) has argued that it is around .2 msec2. Finally, Hopkins and Kristofferson (1280; Hopkins, (1982) were able to obtain synchronization variances to tactile

stimuli that were just as impressively low as the responses when P1 was auditory. Either tactile and auditory latency variances are non-zero and equal or the latencies are not variable and the authors considered non-variability more parsimonious.

For no reason that I could logically defend, I find it difficult to believe that afferent latencies are constant. Nor, assuming that they are, am I convinced that deterministic timing is accomplished at this level. However, the existence of deterministic timing seems well enough established, and Kristofferson's and Hopkins' plausibility arguments for this basis seem valid, as plausibility arguments, and I don't have any better suggestion to offer, so as a working hypothesis, I assume that they are correct.

Given that there is recycling at the afferent level, it can take one of two forms. In Kristofferson's (1976) study, the behavior of one subject was examined across a range from 155 to 2190 msec: His variability was constant in the 300 to 550 msec range, increasing slightly with shorter stimuli, and increasing substantially with longer ones. If it was supposed that a strictly deterministic timer was responsible for timing out to 550 msec, and that a new clock came into play from 550 msec onward, the data indicate that that new clock is a Weber's Law clock. If central timing in this task is non-deterministic, the afferent latency can be lengthened as much as 400 or so msec (the other 150 of the 550 msec latency being taken up by the time it takes to perceive and respond.) This would explain the perceptual shortening reported in the task, and the transition to the

new clock at 550 msec. Presumably, the stimulus can't be held up internally forever.

Alternatively, the stimulus might be recycled only for a few msec, and then be passed to a quantal counting clock. Recalling the example at the start of this section, if the representation of a 312.5 msec.stimulus was always held up by precisely 12.5 masec, the subject would be faced with the simple task of timing a 12 quantum interval. alf we then increased the interval to 317 msec, the afferent delay could be increased to 17 msec, and the task would again be the same for the clock, If the afferent delay, once set, is constant until it is re-set for a new interval, this adds no variability. Part of the function of training might be the development of, or development of reliance on, a constant length delaying neural pathway. This fails to explain why there is a transition at 550 msec, but, as noted earlier, Michon (1967) has made the case that there should be a transition to a new timing mechanism at this interval. On the other hand, from discrimination data to be described next, a strong case can be made that there is no shift to a new type of clock, but there is a quantal counter. If fine-tuning of the quantal counter can be accomplished by short, continuously adjustable, deterministic delays at the afferent level, the devastating argument against quantal counting, i.e. the problem of continuous adjustability, loses its force, and we are free to take evidence for such a mechanism seriously.

1.2.3.6 Quantal Counting

Throughout the discussion above, I've referred to g values that hovered around 12.5, 25, 50, or 100. The visual discrimination data of Allan et, al. (1971) yielded estimates very close to 50 in one condition, and very close to 25 in a later condition, after the experimental procedures were refined. Durations studied in this case ranged from 50 to 150 msec. Synchronization data (Kristofferson (1976) yielded estimates close to 25 in the 160 to 550 msec region of P1-P2 intervals. Converging estimates from speeded discrimination data (Kristofferson, 1977) were virtually identical, and very close to 100 * (95.3). Durations in this case were centered about 1160 msec. Hopkins (1982) used a minimum chi-square estimation technique to estimate the triangular and logistic parameters separately from synchronization response distributions. From very highly practiced subjects, he reported q estimates of 16.6, 14.8, 11.4, and 9.1, averaging 12.98. Synchronization intervals in this case were also within the 170 - 550 msec region. Other examples are distributed throughout the discussion above. What do these different values imply?

Three empirical generalizations come to mind from examining such data. First, as is evident from the numbers, q estimates form a geometric series. Each value of q is double the next smallest. Second, there is some trend for q to increase with base duration, though it is found to be constant across ranges of base durations. Third, if there is a minimum value that can be achieved at a given base or synchronization interval, there is no guarantee that it will be

achieved there. Comparing Hopkins' (1982; Hopkins and Kristofferson, 1980) results in the same range as Kristofferson's (1976) (which shared one subject, by the way, so cross-subject variability doesn't account for this), the q estimates in Hopkins' case are half of Kristofferson's. Hopkins introduced a number of 'procedural refinements into his version of the task, and practiced his subjects even more extensively, and there was evidence of continuing improvement with practice even after 100,000 trials at the same interval.(!) Thus, while there may be a minimum achievable value of q associated with any interval (there may not), the value that will be obtained in any particular task depends on practice levels and on the specifics of the task.

Some further evidence for stepwise increase in q comes from the visual discrimination data of Allan and Kristofferson (1974a). Over the range from 70 to 230 msec, q appeared to be 22. Around 400 msec, only one subject was studied. The q estimate was 59. From 500 upwards, q appeared close to 100, though for one subject it was higher. Still, there was a lot of cross-subject variability, few different base durations (midpoints of single stimulus sets) were involved, and no subject was studied across the range. The stepwise increase is suggested in these data, but it is not clearly evident unless one is looking for it.

Much more extensive data were collected with auditory stimuli by Kristofferson (1973, 1980, 1983) and the case they make is far more convincing. The 1973 and 1980 data are the same (with, I believe, more

data in 1980). The 1983 study used speeded responding whereas the 1980 study did not, and it included much more extensive practice at fewer base durations, selected on the basis of the 1980 results. In the 1980 paper performance at 13 base durations was reported. (Again, these are midpoints of single stimulus sets. The advantage of this procedure is that on a given trial, there is only one stimulus, and so only one triangularly distributed variate.) The base durations ranged from 100 to 1480 msec. The data from the first five sessions at each base duration conformed nicely to Weber's Law. Standard deviations, and thus g estimates, increased linearly with the means. With increasing practice, variability at some intervals declined substantially, while variability at others remained virtually constant. By the end of 20 sessions per base duration, a very different picture was evident in the data. Durations could be grouped into four ranges. At 100 and 160 msec, q was close to 14. From 200 to 350 msec, q was close to 25. Q was nearer 50 in the 450 to 800 msec region, and above that it was closer to 100, with some values around 150 as the interval increased. Within these ranges, q was still increasing somewhat, but at a much slower rate than found at first. Further, the differences across ranges were clearly larger than the differences within ranges. Finally, the q estimates for stimuli toward the low end of any given range had not changed much, whereas those from the longer stimuli in each range had come down considerably. With increased practice, Kristofferson (1980) argued, q values obtained from stimuli at the long end of each range should decline further. That was checked in 1983,

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with up to 70 sessions practice at durations chosen from both ends of some of these ranges, and was clearly confirmed.

The empirical generalization that emerges from such data is that the values of q can be described in terms of "doubles sets". From 100 to 200 msec, q is about 12.5. From 200 to 400 msec, i.e. double the previous range, q is 25. From 400 to 800, it is 50. Double everything again to obtain q = 100 in the 800 to 1600 region. Kristofferson (1983) also studied stimuli centering around 1800, which should be in a 1600 to 3200 range with a q of 200. The estimated q, from response latencies and from response probabilites, was very close to 204. These would be sensible values if the subject's duration estimate was based on a quantal count, with a maximum possible count of 16. If the smallest possible q is 12.5, the subject can use that to time out intervals lasting up to 200 msec. Beyond that, either the subject can give up on counting, or he will have to count in larger units. Quanta being indivisible, the next unit is 2 quanta, 25, which takes him to 400 msec, and so forth. I should be explicit here in noting that this counting is not done consciously.

Fine tuning of the system, according to the argument presented in the previous section, is done by the setting of deterministic afferent delays. Stimuli are perceptually shortened so that the clock can work with an integral number of quanta, even though the stimulus itself does not last an integral number of quanta. Kristofferson (1980) puzzled over the increase in variability within ranges, and suggested that perhaps there was some variance in the count, that

increased with the count. In this case, as stimulus duration increases beyond the minimum value for a range, so does the timing variability. Perhaps this is correct. Alternatively, I suspect that if the afferent delay hypothesis is correct, it would be much harder to delay a stimulus for a long time at this level than it would be to delay it for a short time. Perhaps the variance increase, within ranges, simply reflects the amount of practice required before a stable afferent delay can be obtained at the right value.

The hypothesis of quantal counting, with a 4 bit (16 maximum) counter, cannot be tested in all experimental settings, nor could the subject use such a counter effectively in all settings. Suppose that we randomly varied stimulus duration from 50 msec through 2000 msec within a single session, for example. (Such variation would be quite reasonable in the context of a two alternative forced choice task, in which the subject is merely to state which of two intervals is longer. This is how such tasks are usually run, in most modalities.) The subject has no way of knowing, in this case, what range of stimuli is reasonable for a given trial. If he tried to count quanta, he might be faced with the hopeless task of spreading at most sixteen 12.5 msec q's across a 2000 msec interval. Having learned a lesson on that trial, he might be faced on the next with timing a 50 msec stimulus using a 200 msec quantum. With practice, the subject would undoubtedly become expert in the use of some other timing mechanism. That other timer might be a Poisson clock, a Weber's Law clock, a "perfect" clock, or something else altogether. There is no restriction on the type of

clocks possible in Kristoffersoń's quantal theorizing. The quantum reflects the period of an information gating process, which restricts the temporal precision possible in any situation by introducing variable delays at the level of input to and output from the clock. The expectation that these delays will be relatively large enough, in variability, to be measurable behaviorally is based on the assumption that with practice the subject will learn to use the most sensitive timing strategy that he can. This will only sometimes be a quantal counting strategy.

In naturalistic settings, the use of deterministic timing strategies might be quite restricted. Quantal counting with an appropriate multiple of the minimum quantum size was possible in the discrimination experiments because the same narrow range of durations was used for session after session until well after the range of stimulus intervals possible was extremely well known to the subject. The types of situations in which people might use quantal timing regularly might include timing out a short interval before striking the next Key on a piano, by a professional musician, or timing the required delay between the pitch of a baseball and the swing of the bat by an experienced ball player (hopefully resulting in good synchronization between the arrival time of P2, the ball at the plate, and the response).³

1.2.3.7 Visual/Auditory Modality Differences

If timing is centrally done, and if the quantum reflects a

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periodic process determining when input will progress to central processing, and when processing will progress to the next stage after, the last thing that should affect timing variability is the modality of the stimulus markers, especially if afferent latency variance is zero. Why should it matter whether the stimulus used to mark the onset and offset of an interval is visual, tactile, or auditory? Yet it seems to. Returning to the discrimination data, comparison of the visual results of Allan et. al. (1971) and Allan and Kristofferson (1974a) to the auditory results from Kristofferson (1980) (comparable values were obtained from Brewster, 1983, at 240 msec), and averaging results. across subjects, conditions, and measures, I obtain the following (all units in msec):

	Interval	Auditory q	Visual q	-
	100	14	22 (1974) 34 (1971)	
	150	16	24	
	200	25	29	
	350	31		
	400		59	
	450	48		
	570	60	105	
	740	60	127	
•	910	113	112	

There are some problems with this comparison because the subjects in the visual experiments participated in fewer sessions, in total, than the subject of Kristofferson (1980) and those of Brewster (1983, who obtained q estimates as low as 17 in the 240 msec region, but using a task too complicated to describe in detail here.) This is especially problematic in the 1971 visual study, in which conditions

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were varied in random order, from session to session, which may have reduced the benefits of practice. Further, some procedural refinements affected q, yielding a bimodal distribution of q estimates clustering around 50 and, in the second experiment, 25. In the 1974 study, however, subjects stayed at the same base duration for 10 to 21 (usually 20) sessions (288 trials per session), which isn't very different from the 20 sessions (300 trials each) at each base duration used by Kristofferson (1980). This small difference in practice shouldn't account for the virtual doubling of q values in the visual relative to the auditory data.

There is also evidence of a difference in the estimated duration of visually and auditorily marked intervals. Visually marked intervals are typically perceived as shorter. Goldstone and Lhamon (1972, 1974) and Lhamon and Goldstone (1974) review dozens of experiments of theirs (many with Goldfarb) that employed a wide range of psychophysical procedures, including production, reproduction, discrimination, and category rating. Their findings are quite consistent, and they cite many independent replications. *Interestingly, Lhamon, Edelberg and Goldstone (1962) found a much

smaller difference between auditory and tactile stimuli than between auditory and visual ones. Goldstone's studies have typically employed lots of subjects, making judgments about lots of stimuli, but not making many judgments about any particular stimulus. The only cross-modal study that I am aware of that involved many judgments per stimulus was conducted by Tanner, Patton and Atkinson (1965), who

studied only three pairs of durations, 500 vs 600 msec, 1000 vs 1100 msec, and 1500 vs 1600 msec. Only one pair of intervals was presented (512 times) per session. Either of the pair of stimuli could be auditory or visual. Visual stimuli were judged as longer (not shorter) than auditory stimuli when the base duration was 500 msec, but not otherwise. The authors suggested that they failed to obtain the effects reported by Goldstone and his colleagues because of the difference in practice of their subjects. Goldstone's response has included many more experiments, including discrimination studies (whichhad not been run previously that used many subjects, many stimuli, and not many responses per stimulus. He still gets the effect, which, despite the occasional failure to replicate (eq. Brown and Hitchcock, 1965), seems pretty reliable for unpracticed subjects (see also, Walker and Scott, 1981, who report even more of the same). Eisler (1977) reviewed and reanalyzed studies which indicated that in the psychophysical function for duration can be represented by a power function, the exponent for auditory durations is smaller than that for visual durations. Again, these involve few judgments per stimulus. The one effect arising from this literature for practiced subjects, which is all that I'm interested in for present purposes, is the finding by Tanner et. al. (1965) that pairs of auditory stimuli in their experiment were discriminated more accurately than pairs of visual stimuli, which is what the comparison of the visual and auditory data above also shows. In this case, however, the subjects all have the same level of experience with the stimuli.

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There have been no extremely long term studies of visually presented duration of the class available for auditory durations from Kristofferson (1973, 1980, 1983). Comparisons of fully asymptotic discrimination behavior across modalities is not possible. There has been one_attempt to compare long-term synchronization performance across modalities (Hopkins, 1982).

Hopkins (1982) started with a comparison of synchronization performance when P1 and P2 were tactile to performance when the stimuli were visual. I'll spend some time discussing the specifics of his approach, partially because an understanding of these is prerequisite to an understanding of his interpretation of visual to auditory comparisons, and also because many of my analyses, and an important aspect of my procedure, were directly based on his work.

In the synchronization work of Kristofferson (1976) and Brewster (1983), trials were paced by the computer that controlled the experiment. That is, at the end of a trial, the computer waited a fixed period of time before starting the next trial with a warning stimulus, a loud auditory. "honk". Following the honk was a silent interval, the foreperiod, that lasted a fixed interval of time, and it was followed by P1 and thence P2. Hopkins (1982; Hopkins and Kristofferson, 1980) and Michaels (1977) used self-paced trials instead. Subject to a very short minimum intertrial interval, the subject was free to start the next trial at any time she wanted. To start a trial, she placed her finger on a brass button, closing a very weak electrical circuit between the button and a brass plate on which

she rested the heel of her hand throughout the session. The subject kept her finger on the button until she was ready to respond, in synchrony with P2. Each trial, then, included two responses, a trial initiation response (finger down) and a synchronization response (finger up, breaking the circuit). Hopkins' synchronization stimulus was presented almost immediately after the trial initiation response. The foreperiod in this case, the interval between trial initiation response and onset of P1, was usually fixed at a constant value of 91 msec.

The act of pressing the response button is a source of tactile stimulation in this situation." If the auditory P1 stimulus is perfectly timelocked to the initiation response, as when the foreperiod is a constant 91 msec, the subject doesn't <u>have</u> °to listen to P1 in. order to perform the task. She could just as well start timing from the moment she feels the button, which is a very salient stimulus under the circumstances. Hopkins' goal was to determine which stimulus was controlling her behavior, auditory P1 or tactile feedback from the response. He started by inserting a uniformly distributed delay of up to 20 msec into the foreperiod. The variance of the interval between initiation response and P1 is now 33 msec², which would be trivial except that the variances of the synchronizations were well under 100 msec². P2 was still fully time-locked to P1, i.e. the time between P1 and P2 (the "synchronization interval") was kept constant. . If the subject responded on the basis of the time of P1, this manipulation should have had no effect on her behavior. If she responded on the

basis of the time of her response, P1 and thus P2 were randomly delayed relative to that response, so the extra variance should have shown up in synchronization latencies, which are measured from the time of P1. Hopkins found no difference in performance due to insertion of the variable foreperiod, indicating that P1 was controlling behavior.

Hopkins then removed P1 from the stimulus sequence, forcing the subject to rely on feedback from her response. After practice, when ' the foreperiod was constant the subjects' variances looked exactly the same as they did when P1 was present (the average difference in variances was 0.15 msec²). This indicates that subjects can use their initiation response as a temporal marker and that they can use it as precisely as an auditory stimulus. Hopkins also inserted a uniformly distributed delay into the foreperiod when P1 was not presented to the subject. In this case latency variance, relative to P2, increased by 33 msec². This was simply a check, to confirm that if the subjects were using the initiation response in the presence of auditory P1, the addition of a random delay between that response and P1 would have had a detectable effect on their performance.

Hopkins' results tell us three things. First they tell us that subjects can indeed do what they're instructed to do in the auditory case, i.e. start timing the P1-P2 interval from B1. Second, they tell us that the subjects don't have to do this. They can perform just as well if they base their time estimates on the tactile stimulation obtained from their trial initiation response. Finally, if afferent latencies associated with, auditory and tactile stimuli are variable, the variances in both modalities must be the same. Otherwise the synchronization latency variances in these conditions would have differed.

Hopkins proceeded to compare visual and auditory performance in the task, using some new and some auditorily experienced subjects with visually presented P1 and P2 stimuli. Performance was somewhat more variable with the visual stimuli, compared to performance with auditory stimuli at the same stage of practice, through much of acquisition, but eventually the variances settled down to a level somewhat comparable to the auditory levels. In this case, however, insertion of a random delay between initiation response and the visual P1 introduced variance into the subjects' response latencies, as measured from P1. Further, performance with visual P1 present was the same as performance with visual P1 absent, whether or not the foreperiod was random. This indicates that the subjects were relying on tactile feedback from their initiation responses in both cases, i.e. that they had learned to ignore visual P1.

Ignoring visual P1 is a suboptimal strategy when the foreperiod is variable, unless the variance associated with processing the visual stimulus (such as afferent latency variance) is greater than that associated with tactile stimuli. If visual afferent latencies are more variable than tactile latencies by more than 33 msec², the internal time of visual P1 is less reliable as a predictor of the time of P2 than is the internal time associated with the initiation response, even if the delay between that response and P2 is experimentally subjected

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to jitter of up to 20 msec. Hopkins argued that visual afferent latencies are variable, and are more variable than tactile and auditory latencies by at least 33 msec².

Hopkins also studied the effect of removing P2 on performance For short P1-P2 intervals, the only difference between synchronization and many reaction time paradigms is the presence of the marker, P2, signalling when, exactly, the subject is to respond. There is another feedback stimulus, presented after the response, indicating whether the response was early or late relative to P2, and by how much. The temporal asynchrony between P2 and the response serves as immediate feedback about the accuracy of the synchronization response. The later feedback serves as delayed feedback. Comparable delayed feedback has been used in simple reaction time experiments (experiments without a P2), in which the subject is instructed to respond within a certain small range of times after the stimulus (eg. Saslow, 1974; Snodgrass, 1969; Snodgrass, Luce and Galanter, 1967). Response latency variances in those tasks were much higher than those obtained from synchronization, a point discussed at length by Kristofferson (1976; Hopkins and Kristofferson, 1980; and Hopkins, 1982). The goal of this modification to the simple RT paradigm was variance minimization. The obvious difference between synchronization and these modified RT tasks is the presence of immediate feedback (P2), so it appears that the presence of P2 is important for acquisition and maintenance of low variance synchronization performance. Hopkins found that removal of P2 had an effect (averaging 35 msec²) on the subjects' variability

when P1 and P2 were auditory. Removal of P2 when it was visual had no effect on the results from well practiced subjects, suggesting again that they had learned to ignore the visual stimulus markers. The effects of removing both P1 and P2 in the visual case depended on the amount of the practice of the subject, suggesting that the subject had used visual markers early in training but, again, that they were not used later on.

I should note that Hopkins reported that his subjects (himself included) fully believed that they were relying on the visual stimulus markers when they were present, even though all of the data indicate that they were not. This was not a mere strategy problem that could have been changed with better or repeated instructions.

By the end of the experiments, visual variances were comparable to auditory variances, but were still higher. Asymptotic performance of the one subject studied in both visual and auditory stimuli, when the foreperiod was constant, was 53 msec² in the visual case and 34 msec² in the auditory case. The performance difference may have been due to the ineffectiveness of visual P2 in the visual condition.

Hopkins' conclusion, that subjects learned to ignore visual P1 and P2, seems inescapable. His suggestion that they did so, unconsciously, because of increased afferent latency variance in the visual system, is less certain. There is evidence, as Hopkins pointed out, that visual latencies are variable. Hopkins cited a study by Zacks (1973), who studied the discriminability of temporal order of light flashes. Two spatially separated flashes/were presented to the

subject, who had to judge which started first. Using analyses derived from Signal Detection Theory, Zacks estimated the variances in the task as ranging from 56 to 151 msec² (or 182 msec² with less bright stimuli). As Zacks (1973) pointed out, these represent upper limits on afferent latency variability. Levick (1973) estimated response latency variances of retinal ganglion cells at various intensities, using a number of procedures (see also Levick and Zacks, 1970). In this case, latency variance estimates were substantially lower, ranging from 1 to 36 msec². It is difficult to interpret cell variabilities, however, because the final-result may be more or less variable than that of component inputs. This point was made at length by Meijers and Eijkman (1974), who found variances on the order of 400 msec² in the responses of pyramidal tract neurons whose output eventually lead to motor responses. Variances of the responses themselves, measured behaviorally (eg by Wing) or by examination of the EMG (Meijers and Eijkman) are closer to 20 msec² despite the much larger variability associated with the component inputs. So perhaps Zacks' (1973) psychophysically based estimates are the ones we should rely on.

There are real problems associated with an afferent latency variance in the visual system of 56 to 151 msec², though. Harking back to the temporal discrimination results of Allan et. al. (1971) and Allan and Kristofferson (1974a), we find q estimates for visual duration stimuli that are reliably as low as 22. These are based on response probabilities, based on judgments of the durations of visual stimuli. Thus, the variance of the judged durations is about 81
msec². If variation in the afferent latencies perturbs perceived onset with a variance of 56 to 151 msec², the variance left to be accounted for by timing is remarkably low, ranging from 25 msec² (q = 12.2) to -70 msec², which is an unlikely value for a variance. Note that this ignores the afferent latency variance associated with the offset of the stimulus, which would cut our estimates of q even further. From these data, then, afferent latency variance couldn't be more than 50 and is probably much less.

There is another way of interpreting Zacks' (1973) results. Zacks made the argument that the stimuli were widely enough separated that they could be considered, in effect, to be processed along separate input channels. That is, presentation of one has no impact on processing of the other. In this case, temporal order judgments fit within the standard framework of a quantal gating theory. Successiveness judgments have often been found to be describable in terms of a uniform distribution (Kristofferson, 1967a; Schmidt and Kristofferson, 1963, for example), representing a one-stage quantal delay in processing. Suppose we interpret these variances along those lines and estimate q (variance = $q^2/12$). The three variances reported by Zacks were 56 (q = 25.9), 151 (q=43.3) and 182 (q=46.8), close to 25 and 50 as is typical of quantal estimates. Zacks' variances need not reflect afferent variability at all.

If there is a 25 to 50 msec² afferent latency variance associated with visual stimuli, this could explain the discrepancy between auditory and visual discrimination performance when q is found

to be close to 12.5 in the auditory case and close to 25 in the visual case, as the calculations of the second last paragraph show. However, they do not explain discrepancies at longer intervals, when we are dealing with larger values of q. At 570 msec, for example, q was estimated at 60 msec when stimuli were auditory and at 105 msec when stimuli were visual. The variances associated with these values are 600 and 1838 msec² respectively. The difference of 1238 msec² cannot be accounted for by a 50 msec² afferent latency variance.

In sum, there is evidence for a difference in temporal discriminability as a function of stimulus modality, but this may instead be due to a difference in practice levels of the subjects. It is not due to a small difference in afferent latency variance. There is evidence for some difference in estimation of intervals as a function of marker modality, though it may disappear with practice. There is some evidence of afferent latency variance in the visual system. Afferent variability appears less likely in the auditory and tactile systems. There is clear evidence that experienced subjects. don't use visual synchronization stimuli under Hopkins' conditions, and there is an argument that this is because of higher variance in the visual system than in the auditory system. There are no long term comparisons of auditory and visual discrimination or synchronization performance. I don't count Hopkins' data as a long term comparison because, as he so carefully demonstrated, the subjects didn't use the visual stimuli, so his data don't tell us what would have happened if they had used the stimuli. There are grounds for concluding that

timing of visual and auditory stimuli differ, but this conclusion is not inescapable.

I find Hopkins' arguments difficult to fully credit more because of my extended experiences with synchronization as a subject in Brewsters' research and in pilot studies leading to the present research than because of my theoretical predilections. There are differences between auditory and visual stimulus conditions which are not captured by dry descriptions of experimental procedures. Here are some of my introspections. Many of these have also been volunteered by subjects in the present experiments. They are not strictly idiosyncratic.

The auditory stimulus is a 68 dB 10 msec tone, delivered over headphones. It is quite salient. Since it is delivered over headphones, the intensity does not change with the posture of the subject. Subjects were free to vary their posture in these experiments. (Uncomfortable subjects probably wouldn't produce such low variances). The result of this is that a subject can change the position of his head by over a meter relative to the position of the stimulus light used for delayed feedback or as P1 and P2. Sometimes the subject is hunched over the desk on which the response pad (holding the button and the brass plate) rests. Sometimes the subject leans back with his feet up, pad on his lap or on a different desk. The distance of the visual stimulus varies over at least a twofold range, i.e. the intensity varies by a factor of four or more, and it can vary from trial to trial. Latency is definitely a function of intensity, as

is routinely found in reaction time and other studies (see Grice, 1968; Kohfeld, 1971; Kohfeld, Santee and Wallace, 1981; Levick, 1973; Teichner, 1954, and their many references). Free movement of the subject in the visual case introduces afferent latency variability that has nothing to do with latency variance for identical stimuli. (It is tempting to argue that subjects in the visual case should be forced to use bite bars, but I not only doubt that many subjects would be willing to participate in the experiments over long periods of time in this case but I also think the discomfort involved, especially toward the end of a session, would induce more variance than would be saved by the control. My experience with bite bars dates from employment as a research assistant in Hiroshi Ono's laboratory at York University, who used them routinely for visual research.)

The visual stimulus used by Hopkins was a 5.08 mm light emitting diode which produced a 3.2 mcd stimulus when driven at 20 mA. Rise time for led's is very short, so this intensity was attained even for the very short 10 msec duration of P1 and P2. Hopkins (and Brewster, (1983), who used a similar light) called this stimulus "salient". It was salient in the sense that if you were looking at it and it came on, it was clearly on. But if you weren't looking directly at it, you could miss it. In pilot work with the light, I found blinking a problem. It wasn't a big problem because I didn't blink at just the wrong time very often. But relative to a variance of 33 msec², almost any problem can be a big problem. As an auditory subject, to aid concentration, I often stared off into space, totally

ignoring the delayed feedback light that Brewster (and Hopkins and I) used to indicate whether the response was early or late. Other times, I would close my eyes, or stare at my finger as I responded. Again, this reduced the value of delayed feedback, but that didn't seem to matter, a point made by some of the present studies' subjects (who did the same things). Hopkins (1982) demonstrated that delayed feedback indeed did not matter once subjects were experienced. In the auditory case, we could afford to ignore the visual stimulus. Closing your eyes may be conducive to concentration on auditory stimuli but it is not the best of tactics when P1 and P2 are visual. Thus, many of the things that I could do, and often did do as an auditory subject, I could not do as a visual subject. I missed them in pilot work.

Even when I was looking at the light, I would not say that it was as salient a stimulus as the tactile feedback from pressing the response button on Hopkins' (and my) pad to start a trial. Pressing that button was a voluntary behavior and I expected to feel it. The auditory stimulus was, to me, more salient than the visual one. I would rate the tactile feedback as more salient still, but only slightly, and others might disagree in either direction in the auditory vs. tactile case.

Hopkins reported that the subjects in his experiment (including himself, in visual and auditory conditions) felt that P1 came immediately after the trial initiation response, with no delay, and, when the delay was variable, they did not notice it. He knew the delay was variable, and he reports that he still didn't notice it. In such a

case, I doubt that that variability was behaviorally effective. My suspicion is that in Hopkins' visual task, the clearly more salient stimulus, i.e. the tactile feedback, is the one that controlled the subject's behavior. The stimulus was so closely connected with P1 that, according to Hopkins, the subjects felt they were using visual P1 even though they weren't. The distinction between the stimuli appears to have been insufficient for the subjects to notice that they were not using P1, but were instead using the more salient stimulus, and to

Separation of the trial initiation response by a longer and more variable foreperiod would probably go a long way toward aiding the subject in separating them subjectively, making it easier for him to use the visual stimulus. A brighter and/or larger stimulus would probably solve some of the salience problems. Unfortunately, it still wouldn't solve all of them unless the stimulus was unreasonably bright or large. (By unreasonably I mean that some of the stimuli I tried in pilot studies gave me prolonged headaches. Other stimuli, which I did note try, would have been excessively expensive.) The subject still has to be looking at the stimulus, which rules out staring blankly, staring at one's finger, closing one's eyes, and blinking. Even with the stimulus that I eventually settled on, I would expect acquisition of low variance levels to take longer than it takes in the auditory case, because the auditory stimulus, to me, seems more salient. This expectation is stronger for Hopkins' stimulus light.

Summing up this section, the issue of timing dependence on \Im

modality seems unclear, at least for practiced subjects. Asymptotic performance levels might or might not be equivalent. For unpracticed subjects they are not. Nor are they for fairly well practiced subjects. The difference in exponents, in time estimation studies, and, to a lesser degree the difference in perceived duration of lights and tones in various studies, nearly all obtained from relatively unpracticed subjects, suggests that there might be a difference in processing mechanisms that goes beyond a difference in variability. Afferent latency variance in the visual system accounts, to my mind, for neither effect, and I am not convinced that the evidence for afferent latency variance in this syster is conclusive. Finally, the evidence for afferent latency variance from synchronization is, I would argue, wholly inconclusive. Asymptotic performance levels might not differ at all and even if they did this might reflect other differences in task difficulty and latency variance due to variance of the intensity of the visual stimulus, caused by postural changes by the subjects.

1.3 Boundapy Conditions

The four approaches to internal clocks discussed above make clearly different predictions about the variability associated with timing. Yet if we ask whether the variance or the standard deviation increases linearly with duration or if, instead, the increase is stepwise or nonexistent, the answer we get back is "yes". Competently run experiments have provided evidence for all of these relationships.

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It appears that people can use a variety of different mechanisms for timing, in which case we should be studying the circumstances under which each is used, not trying to decide which one is the mechanism. There is a theoretical risk associated with an eclectic viewpoint: new mechanism can be invoked to account for the*results of each new experiment. One way around that is to focus on a single mechanism that seems well established and to determine how much behavior can be explained in terms of it. Ideally, systematic examination of a number of different models would lead to acceptance of domains of applicability of each, probably with some indistinct boundaries separating different domains. Systematic work along these lines has been done for the quantal theory, and the intent of the present research is to explore what appear to be fuzzy edges along the deterministic quantal timer's boundaries. In this section I will briefly recap the description of the quantal applicability domain, leading to a statement of the present thesis in terms of it.

Variability in timing is subject to an irreducible minimum, the time quantum, which is the period of an internal gating process. Processing of information passes through stages. The periodic process(es) determines when the information may be passed to the next stage of processing. Delays at each stage are uniformly distributed, ranging from 0 to q msec, and are effectively independent. Waiting time until processing is complete can thus be represented as the sum of a series of identically distributed uniform variates. In some tasks, such as successiveness discrimination, only one quantal delay may be

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involved. In many tasks involving timing, two stages are involved, and in these timing variability is triangularly distributed.

Quantal delays are assumed to be involved in all tasks but the relative contribution that they make to the variability of judgments or behavior may sometimes be negligible. Quantal delays are relatively short, so their variance is relatively small. Variance from other sources can mask the contribution of these delays. For example, if the standard deviation of an internal timer increases proportionally with timed duration, the quantal contribution might only show up as a positive intercept in the function relating discriminability (whose variance sources include the timer plus delays into and out of the timer) to duration. Evidence for guantal delays, then, and data suitable for estimation of their magnitude, should only be expected to be obtainable under highly réstricted situations, in which all other sources of variability are brought under control. In the optimal case, the only central source of variability will be the periodic gating process. This can be achieved in a timing task only if the subject uses a fully deterministic timer. It is remarkable that people are capable of using such a timer at all, and we should not expect that such a mechanism will be usable under all circumstances. It appears, from the data at hand, that at least five variables limit the range of circumstances under which a deterministic timer can be employed consistently by the subject, to the exclusion of other mechanisms, . these being the range over which the turrent stimulus can vary, the degree of experience that the subject has had with the specific stimuli

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involved in the task, the type of task, the duration of the stimuli, and, perhaps, the stimulus modality.

From Kristofferson's (1980, 1983) discrimination results it appears that deterministic timing includes a quantal counter, i.e. one which increments once per gating period. In auditory tasks, this appears to be a_{γ} four bit counter, i.e. it runs to 16 at most. If intervals longer than 16 guanta are to be counted, the guantal intervals must be concatenated, so that the count advances once per n quanta, where n is a nonnegative integral power of 2. These larger values themselves serve as quantal units. A counting strategy will fail in any task in which the same guantal unit cannot be used for all of the stimuli that might be presented on a given trial. Subjects might use the q value necessary to time the longest interval to time all intervals, but at some stage, such as working with a q of 200 msec when stimuli lasting 20 msec are possible, this is not an optimal. timing strategy and it will probably be discarded. Quantal counting should not be expected in typical scaling tasks, such as magnitude estimation, magnitude production, and so forth (described in Stevens, 1976), nor should it arise in discrimination experiments involving roving standards, i.e. cases in which pairs of similar stimuli are to be evaluated during each trial but pairs may differ widely across trials. In such cases the first stimulus of a pair could not be timed accurately, so there-would be no basis for comparison with the second. The 4 bit quantal counter describes auditorx-discriminationprobabilities and latencies extraordinarily well, but visu

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discrimination data and auditory synchronization latency data both suggest limits to its generality. In the visual case, q also appears to increase in steps, but values of q are larger at many durations than they are when stimuli are auditory. If stepped increases are due to a maximum possible count, the visual maximum must be lower. On the other side, Hopkins' results indicate that a q as small as 12.5 can be used when auditory synchronization intervals are 460 msec long. Even if we subtract 150 msec from the 400 to take afferent and efferent delays into account, central timing would involve 310 msec, or a count of 24.

Practice may not make timing perfect but it helps, and it continues to help all the way out to 100,000 synchronizations at the same interval and, apparently, beyond (Hopkins, 1982). Different amounts of practice seem necessary for different stimuli, at least in: discrimination. Kristofferson (1980, 1983) found that little practice was needed for asymptotic performance to be reached at the low end of a quantal step but that much longer practice was needed for durations at the high end of the step, i.e. for durations which required a large count of the smallest usable (count < 16) quantum level. Early in practice, performance conformed to Weber's Law (see also Getty, 1975). Only after thousands of trials with stimuli at the high end of their respective steps did performance fully conform to a quantal counting model. Also evident in those data, but not in the synchronization data of Kristofferson (1976) and Hopkins (1982; Hopkins and Kristofferson, 1980), is the need for extended training at the specific interval under study. There is relatively little transfer of training in

discrimination across durations.

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In at least one task in which subjects have been trained for long periods at the same interval, timing never appears to be quantal. This is the repetitive tapping task, in which the subject is to tap-tap-tap-etc. with a fixed inter-tap interval. Wing (1973) demonstrated that the variability of well practiced subjects in this task increases with duration and provided one well cited model (Wing and Kristofferson, 1973a) to account for this. Vorberg_and Hambuch (1976) showed that different models describe performance as the interval is increased over a fairly modest range. There are complexities in this behavior which are most surprising given its . subjective simplicity. Kristofferson (1981) has interpreted the task as involving "pure motor timing". Many tasks are executed under the control of motor programs which, once started, continue without the need of peripheral (to central feedback (Schmidt, 1982). Perhaps this is one of them, in which case no opportunity for deterministic central timing would arise.

Regarding stimulus duration, internal clock theories have typically been restricted to short intervals. Poppel (1971), a quantal advocate, indicates that he expects the quantal mechanism to break down for intervals longer than 10 seconds. Kristofferson has proposed no specific upper limit but in discussions we have had, he has made it clear that he expects some upper limit. Relevant research in his lab has generally been restricted to about 2000 msec and less. Fraisse (1963) has argued that judgments of intervals longer than 2 seconds;

must involve higher order cognitive processes. It is interesting that Getty (1975) found Weber's Law inapplicable for durations longer than 2 seconds, and this in fact replicates much older work (reviewed by Fraisse). Most research involving cognitive processes and time has involved relatively long stimuli, with the notable exception of Thomas' studies. Thomas (eg. Thomas and Brown, 1974) used stimuli on the order of 100 msec and has repeatedly found evidence of cognitive mechanisms in conjunction with an internal timer. Thomas' case, which I used to find convincing, has been weakened by the argument of Long and Beaton (1980a, 1980b, 1981) that such effects may be due to visual persistence. Bigger, brighter, more complex stimuli last longer on the retina than smaller, duller, simpler ones. There may well be higher-order determinants of this low-level physiological process, but if there are, this is an altogether different role for "cognitive" variables in duration perception.

There is some indication of a second transition point in timing, in the 500 to 700 msec region. Michon (1967) reviewed and presented evidence along a number of lines but it is not clear that those data have any relevance to well-practiced behavior in tasks in which we would expect to find, deterministic timing. (Though some workers in the field, such as Carbotte (1973) believe that they, should be relevant.) There is no hint in discrimination results of a change in mechanism, just a change in q, with increasing diration. However, there is clear evidence of a shift in timers at 550 msec in synchronization. Kristofferson (1976) found that Weber's Law described

performance from 550 to 2190 msec in the task despite extensive practice at 940 msec.

Finally, we come to modality. Some modality restrictions should be obvious: subjective duration of tastes or odors would probably be too variable to be of interest. Alternatively, there appears to be no difference between practiced timing of auditory and t_{a} of the stimuli (Hopkins and Kristofferson, 1980), and very little difference in relatively unpracticed estimation across the modalities (Lhamon et, at, 1962). There is a mass of confusing data for auditory vs. visual timing. 🔀 For more, see Allan's 1979 review.) If Hopkins' (1982) conjecture that visual afferent latencies are variable is correct, and if Kristofferson's (1976) conjecture that continuously addustable deterministic delays are established via recycling through the afferent systems is also correct, then we might expect visual synchronization pergormance to show increasing variability with duration, as the length of time over which the stimulus is recycled increases. Alternatively, if minimum length delays are used, i.e. if 4 stimuli are only held in the system until the learned remainder of their duration is an even multiple of q, we should expect to find a periodic increase of variance with duration, reaching a maximum when the stimulus is infinitesmally shorter than an integral multiple of q.

1.4 Statement of the Thesis (

My goal in the present research is to demonstrate that the same timing mechanism is available and is used by experienced subjects in a

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synchronization task whether the stimuli are long (to 2100 msec) or short (to 150-msec) and whether the stimuli are auditory or visual.

I expected to find that synchronization with long auditory stimuli yields performance comparable to that obtained with short stimuli either in all respects or in all respects except absolute level of variance once extended practice across a range of long stimuli has been completed. I made no predictions about initial performance with long auditory stimuli, except that later performance would be much better. I expected asymptotic performance to yield constant variance or stepped increase in variance with increasing stimulus duration across a range of long duration stimuli.

I expected to find that visual and auditory performance are comparable in distributional shape, in showing similar practice effects, in showing continuous adjustability, and in the constancy of variance across stimulus ranges or subranges, without any obvious periodic relationship between duration and variance in either modality. I hoped, but did not expect, to find that visual and auditory variance levels would be the same. I very much hoped, but did not expect, that visual stimuli would yield asymptotic performance levels as quickly (relatively speaking) as auditory stimuli.

1.5 Overview of the Research

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Five closely interrelated experiments were conducted, using very long term subjects.

The first experiment studied auditory synchronization in the

1400[']- 2100 msec range.

The second experiment examined auditory synchronization performance in the 150 - 350 msec range, to allow comparisons with longer interval auditory data and with visual synchronization performance.

Experiment 3 was intimately connected with Experiment 2 in execution but had a stronger conceptual link with Experiment 1. In this experiment (3), I reduced the P1-P2 interval of subjects who had been working with 150 to 200 msec intervals to below 100 msec. It should be impossible for subjects to use P1 effectively as a response trigger because absolute minimum reaction time is estimated at 100 to 105 msec in the literature (see, eg., Kohfeld's, 1971 review, and some further discussion below, in Chapter 4.) Accordingly, they should be timing from the warning stimulus or the trial initiation response. Given a long foreperiod (close to a second here), this is a covert long auditory synchronization-condition, and it allows within-subject comparison of long and short auditory synchronization performance, which was otherwise highly impractical.

Experiments 4 and 5 involved visual synchronization in the 275 - 500 msec region. Experiment 4 used new subjects. Experiment 5 used the one subject-from Experiments 2 and 3 who was willing and able to participate further in the research, hopefully allowing within-subject comparison of auditory and visual performance.

The procedure is discussed in detail in the next chapter. It was quite close to Hopkins', but with a few key revisions. The visual

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stimulus was larger and brighter than Hopkins'. The foreperiod was longer and more variable, hopefully making it easier for the subject (and for the experimenter) to distinguish timing from the Warning Stimulus (WS) from timing from P1. The larger variance and a geometric, distribution of foreperiods rather than a uniform distribution were used to minimize the effectiveness of reliance on WS (or, equivalently, on the trial initiation response) instead of P1. Finally, Hopkins studied performance at very few intervals, relying on very extensive practice at a given interval to produce asymptotic performance at that interval. Kristofferson (1976) showed that apparently asymptotic performance at a given interval could be substantially improved upon if the subject was trained at other intervals and then returned to the original one. Suspecting that this would be a more efficient means of achieving minimal variance levels, and because I was interested in performance across a range of durations in each Experiment, I used Kristofferson's "scanning" technique (practice across a range with extensive training only at a few points) rather than studying performance at only one interval per condition.

2. Methodological Details

2.1 The Session in Brief

The sequence of events in a trial was: trial initiation response, warning stimulus, foreperiod, P1, silent period, P2, silent period, feedback, intertrial interval. The subject was to respond in synchrony with P2, which was time-locked to P1. There were 100 trials in a block, with four blocks per session. Subjects were advised that the first block of each session was a "warm-up" block and that only the data from the last three blocks would be analyzed. Discussions of all analyses below refer only to the last 300 trials of each session.

By "P1-P2" interval I mean the interval from P1 onset to P2 onset, which was constant for all trials in a session. The "synchronization interval" was usually the P1-P2 interval, and I often use them interchangeably. In some cases I explicitly distinguish between them below: the synchronization interval is the interval between the stimulus from which the subject times (usually P1) and P2.

2.2 Computer Control

Stimulus presentation, response collection and all timing was controlled by a computer, originally a PDP-B/e minicomputer and, for the majority of the sessions, an Apple II microcomputer. The PDP-8-was retired after a series of equipment failures, including an intermittent timing problem. There was a six week delay between shutdown of the PDP-8 and completion of testing of the Apple program and a variety of

new peripheral devices introduced with it. Some aspects of the procedure were modified at this time, with, so far as I can tell, no effect on the subjects' performance. These changes are noted below.

The PDP-8 was programmed in a version of Focal, Tenhanced mainly by Reese, Michaels and Hopkins to allow real-time control. Durations were specified in msec in this language and the interval from the trial initiation response to the end of the interval preceding feedback had to be restricted to 4096 msec to avoid timing variability. Timing of critical events within a trial was accurate to within a millisecond, typically to within narrower limits. The main Apple program was written in 6502 Assembler. Input of session parameters at the start of the session and data analysis and output at the end were done in Basic, which was co-resident with the Assembler program. Durations were specified in units of 200 microseconds, with typical timing accuracy to within 200 microseconds and a maximum possible error from trial initiation to the end of the interval before feedback of 600 microseconds. I spent about 8 months designing and testing this system and, since it was brought on line, no "bugs" have shown up and no modifications to the assembler program have been made. Session parameters, such as device addresses and output voltage levels, were specified in a separate data file on disk, read at the start of each session. I made a few blunders in describing the stimuli to the computer, delivering a 1000 Hz auditory stimulus to the subject at about 60 dB rather than a 2000 Hz, at 68 dB for 🕏 few sessions. These were caught and fixed early on. Maximum trial duration in this

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program, when timing is done in 200 microsecond units, is just over 13 seconds, rather than 4 seconds as on the PDP-8. This eliminated the need for some otherwise arbitrary restrictions on some intervals.

2.3 The Response Pad '

Responses were made on a response pag, basically an armrest with a metal button near the front end and a metal plate toward the middle. The subject rested the heel of her hand on the metal plate throughout the session. To start a trial, she rested her finger on the button, completing a very weak electrical circuit from the plate to the button. Circuitry in the pad amplified this signal and sent it to the computer. The subject kept her finger on the button until she was ready to make her synchronization response, which she did by moving her finger away from the button, usually upwards, breaking the circuit. Schematics for the original bad were given by Hop (1982). The original pad wore out and was replaced with a similar one on which the metal was copper rather than brass. Some minor details of the circuitry were changed with the changeover to the Apple and, later, to improve the pad's signal to noise ratio.

There is no travel time on the response but on and only a minimal degree of pressure on the button was required to maintain the electrical connection.

2.4 Warning Stimulus and Foreperiod

Hopkins' (1982) finding that subjects started timing at the art of the trial, rather than from P1, when P1 was visual, is

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reminiscent of the common problem in reaction time (RT) experiments that subjects time from the warning stimulus rather than reacting to the reaction stimulus (equivalent to P1). In fact, while some timing work has been done with synchronization-like paradigms (eg. Bartlett and Bartlett, 1959), most of the development of this procedure was done in an effort to bring this problem for RT measurement under control by indicating to the subject when she should respond in reaction time tasks (Naatanen, Muranen and Merisalo, 1974; Saslow, 1974; Snodgrass, 1969; Snodgrass, Luce and Galanter, 1967; and see the reviews in . Kristofferson, 1976; Hopkins and Kristofferson, 1980). When the P1-P2 interval is very short, the subject must respond immediately to achieve γ coincidence with P2, so synchronization is a reaction time task. Kristofferson (1976) showed that delayed reactions, i.e. synchronizations up to 400 msec past reaction time to P1 (about 150 msec), looked the same in variance and distributional shape as synchronization responses when the P1-P2 interval was at the reaction time limit.

Kristofferson (1976) and Brewster (1983) initiated a trial with a warning stimulus, a low frequency, noisy, 68 dB (A-weighted) "honk", delivered to the subject over an external speaker (rather than headphones), which lasted for 100 msec. P1 was presented 1000 msec later. Michaels (1977) also used a 1000 msec foreperiod but the subject's trial initiation response replaced the honk as the signal that the trial had begun. I prefer self-pacing (not done by Kristofferson or Brewster), but-with a long foreperiod it helps to have a/stimulus that signals that the constrained picked up the

initiation response and that the trial has now started. This may be one of the reasons that Michaels' results were so much more variable than the others, though his procedure differed from the others in many other respects too. Hopkins_used an extremely short foreperiod (91 msec when it was constant) and was the first to use a variable foreperiod to determine from which stimulus the subject started timing. In his case, subjects didn't even notice the foreperiod, and they used their trial initiation response, rather than P1, when P1 was visual. Wanting the subjects to time from P1, I used a long foreperiod to separate P1 and trial initiation. I also increased its variability, to 90 msec², partially to make it easier for me to tell which stimulus controlled behavior and partially to make pre-foreperiod events less reliable for the subjects as predictors of the time of P1 and P2. As did Kristofferson and Brewster; I preceded the foreperiod with the "honk" warning stimulus.

Hopkins' foreperiods were uniformly distributed, over a 20 msec range. In the reaction time literature, uniformly distributed foreperiods are quite common, and the typical finding is that reaction time is negatively correlated with foreperiod duration (Klemmer, 1956, 1957; Ollman and Billington, 1972; Naatanen, 1970; Niemi and Naatanen, 1981). The intention behind adoption of random foreperiods in these settings is to make the time of the reaction stimulus sufficiently unpredictable that the subject will give up on a strategy of timing from the warning stimulus. Uniformly variable foreperiods don't achieve this goal, partially because their hazard functions are not constant. The hazard function is a conditional probability function:

given that the stimulus has a set been presented, how likely is it that it will come in the next few msecs? For uniformly distributed foreperiods, which by definition can only last so long, 'this conditional probability increases as the foreperiod drags on, and this seems to provide usable information to the subject. Geometrically distributed foreperiods have constant hazard functions. 'Knowing that the stimulus has not yet happened gives you no further information about when it will happen. Very few reaction time studies have used geometric, or approximately geometric, foreperiods, but those that have have found smaller correlations between response latency and foreperiod duration (reviewed by Niemi and Naatanen, 1984) With long and variable foreperiods I was worried about this, and adopted a geometric rather than a uniform foreperiod distribution.

On the PDP-8, foreperiods ranged from 890 to 990 msec, with a mean of 900 msec, rather than 1000 msec, the foreperiod length of Brewster, Michaels and Kristofferson. I kept the interval shorter, and restricted its range, because of the 4096 msec trial duration limit on the PDP-8. Those extra milliseconds were needed in the long synchronization interval work to allow the subject plenty of time to make a late response. This mean was inconvenient for Experiment 3, however, and on the Apple it was changed.

In Experiment 3, I covertly forced subjects to time from the warning stimulus (or from their initiation response; to which it was time-locked) by making the P1-P2 interval impossibly short. Subjects' variances at those levels suggest that they were operating with a q of about 50, which is operative in discrimination experiments over a 400

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to 800 msec range. With a foreperiod of 900 msec (on average) and a P1-P2 interval of 100 msec (or less), if subjects time from the warning stimulus the actual synchronization interval is at least 1000 msec. It is probably 1100 msec, the time between the onset of the honk and the onset ϑ_{f} P2. Subtracting 150 msec as an upper estimate of the time taken by afferent and efferent delays, we find that the subject is timing an interval between 850 and 950 msec in duration, barely into the q=100 range, which, in discrimination, is operative across base durations from 800 to 1600 msec. The claim that subjects can use a g of 50 for intervals longer than 800 msec could be more convincingly made with a longer foreperiod. On the Apple, the mean foreperiod was 1000 msec rather than 900 as on the PDP-8, yielding net timed intervals. of about 950 to 1050 msec rather than 850 to 950 msec. These are still at the short end of the 800 - 1600 msec range but a larger change may have produced unwanted side effects. As far as I can tell, this 100 msec shift had no effect on any of the data.

Both computers were involved in Experiment 3. Mean foreperiod duration is graphed and tabled on a session by session basis below, and this indicates which computer each session. There was no specified maximum foreperiod duration on the Apple, as there should not be for geometric variates. The variance was kept constant across computers at 90 msec².

Finally, different algorithms were used to generate the geometric foreperiods on the PDP-8 and on the Apple. The LabFocal random number generator does not pass standard tests of randomness. I used the generator to select (without replacement) members of a previously generated list of candidate foreperiods, which appeared in the list in proportion to their geometric probabilities. This yielded a distribution which approximated the geometric well in terms of 4 frequency (as tested by the Kolmogorov-Smirnov and Chi-Square tests) but which suffered from a small serial dependence of one foreperiod duration on those preceding it. The dependence was smaller than would have been obtained from a more direct use of the LabFocal generator, and I don't think it was noticeable, but it was statistically detectable.

The random number generator used on the Apple, for foreperiod generation and for the simulations noted below was developed by John Vokey and myself. This was a traditional mixed linear congruential generator (Knuth, 1981), i.e. one which generates successive numbers, *i* RINJ, RIN+1J, according to the algorithm,

 $R[N+1] = (A R[N] + B) \mod M$,

where A, B and M are constants and M, if A and B are properly chosen, is the period of the generator. In our case, M, and the period, were 240 (just over a trillion). The values of A (we developed three generators of this class) were selected from about 38,000 candidates tested by the full-period Spectral Test (Knuth, 1981), essentially a search for nonrandom patterning of doublets, triplets, etc., in the output of the generator using a Fourier analysis across the entire period. This is the most powerful test Known for random number generators. Large values are good, with 0.10 being a "pass" and 1.0 being a "pass with flying colors" according to Knuth. Our lowest value (examining doublets through sextuplets) was 2.37. Values of B, which

have no effect on the Spectral Test results but which do affect serial correlations, were chosen to minimize full-period serial correlations, lags 1 through 20, and our worst correlation was less than 0.00000002. The values of A for the two generators that I relied upon most heavily were 27182819621 and 8413453205. The respective values of B were 3 and 99991. In simulations, these were often used in combination (for example, using the McLaren-Marsaglia Algorithm, commonly called Algorithm M, again see Knuth) to break up some theoretically remaining (if statistically indetectable) sequential dependencies. The third generator was rarely used because it performed less well than the two above on other tests for randomness that they passed. An extended discussion of the testing of pseudorandom number generators is beyond the present scope. For an extended tutorial-level discussion of our @ approach and results see Kaner and Vokey (Note 1), and for a thorough general treatment?, see Knuth (1981).

2.5- P1 and P2 .

These stimuli were identical in a session. They were either visual or auditory and they Pasted for 10 msec.

If P1 was auditory, it was a 2000 Hz stimulus presented binaurally to the subject over headphones. When on continuously, the intensity of the signal coming from each headphone was 68 dB (re .0002 "dynes/cm²). To avoid "clicks" at onset and offset of the stimulus, the stimulus was gated at zero crossings and was presented with a 2.5 msec rise and decay time. The stimulus was at peak intensity only for the middle 5 msec of P1 and P2. Shaping of the wave envelope was not

done during the first few weeks after the Apple was brought on line.

Light flashes were presented over a pair of light emitting diodes (l.e.d.'s: Hewlett-Packard #2785) glued together to make a single larger stimulus and mounted in a rectangular grey metal box. These l.e.d.'s are rectangular, but a circular stimulus, subtending about 1 degree visual angle, was desired. Only part of the l.e.d. pair was made visible, through a circular aperture cut in the box, measuring 2 cm in diameter. With variation in the subject's head position, the actual visual angle subtended by the stimulus ranged from about .7 to about 1.5 degrees. The stimuli emitted a yellowish green light (585 nm dominant wavelenth), at 7 mcd, when driven at 6 mA. Background lighting was dim but in the photopic range.

Subjectively, the stimulus was easily detected and was seen as reasonably bright. The stimulus was much more salient than the small l.e.d. used by Hopkins and it seemed better than various other stimuli that I tried out in pilot work would rate it as less salient than the auditory stimulus.

Visually bounded intervals were subjectively different from auditorily bounded intervals due to visual persistence of the visual markers. P1 and P2 were physically 10 msec but they lasted much longer subjectively. (See Efron, 1970, Haber and Standing, 1970, and Long and Beaton, 1980a, 1980b, 1981, for more on visual persistence.) An auditory P1-P2 interval of 150 msec was subjectively clearly divided into three parts, P1, silence, P2. Analogously, the visual interval should look like P1, darkness, P2. Instead, with a P1-P2 interval of 150 msec, the visual stimulus typically looked to me like a single

flash that varied in intensity over time. While I always saw three events (P1, dark, P2) when P1-P2 was 250 msec or more, subject DK reported that early in the experiment (but not later) she sometime's saw only one flash even when the interval was as long as 300 msec. Under these conditions, however, the 300 msec interval still looks shorter to me than an auditory 300 msec interval sounds. The relationship across different synchronization intervals between the subjective durations of the visual and auditory P1-P2 stimulus sequences appears nonlinear to me. The goal of Experiment 4, the main experiment in which visual stimuli were used, was demonstration that the response distributions associated with visual stimuli were the same across a 275 - 500 msec synchronization interval range. This result was in fact obtained, with variance levels comparable to those obtained with auditory stimuli in the 150 to 350 msec range. We know from Kristofferson (1976) (that auditory synchronization distributions are the same from 150 to 550 msec, or at least from 300 to 550 msec (depending on which subject's performance is examined). Given this constancy across ranges for both modalities, in the context of the present work it seems not to matter that there are real problems of commensurability with the subjective durations of the auditory and visual stimuli.

The degree of visual persistence is a function of various characteristics of the stimulus. Hopkins' stimulus yields less persistence and so is usable across a wider range of durations, though it is less salient.

2.6 Events Past P2

Following P2 there is a silent interval during which the subject may respond if she has not already responded. The exact length of this interval varied on the PDP-8, depending on the P1-P2 interval, but it was never shorter than 400 msec, which is much longer than reaction time to P2. On the Apple, the interval was fixed at 1000 msec. On both machines there was a short delay between the end of the period in which responses could be made and the start of feedback (shorter and less variable on the Apple, and less than 100 msec, at the most, on either system). During this period, the computer determined which type of feedback was to be presented, on the basis of the response latency.

Feedback came in three varieties. If the subject's response came after P2, a 2000 Hz, 68 db stimulus was presented. It lasted for as many milliseconds as the response was late, where a perfect response would be one synchronous with the onset of P2. If the response was early, the same stimulus was presented, the duration indicating how early instead of how late, and it was accompanied by a light flash of the same duration. The feedback light+was a 5.08 mm diameter l.e.d., easily detectable if the subject was looking in its general direction, but less salient than the P1 light. Responses which coincided with the onset of P2 were originally given no feedback, which was perceptually quite distinct. The subjects repeatedly complained that lack of feedback on these trials broke their rhythm and disrupted performance on the next few trials. I changed the computer program so that it would treat perfect responses, for feedback purposes (only) as though they were 5 msec early.

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The final type of feedback was "blown trial" feedback. If the subject responded before the onset of P1, or if she failed to respond, the trial was considered "blown" and was re-run. Data from these trials was kept and early responses were analyzed when the P1-P2 interval was less than 200 msec, for in such cases early responses were sometimes "real", generated by subjects who were timing from the warning stimulus rather than from P1. On the PDP-8, blown trials were marked by ithree 300 msec presentations of the "honk", separated by 100 msec from each other. On the Apple, the feedback was instead a simultaneous 500 msec presentation of the honk, the P1 light and the feedback light. Both were obviously different from normal feedback. They were also quite effective for waking dozing subjects, which was sometimes necessary in this rather boring task.

2.7 Instructions and Other Information Available to the Subjects

Subjects were told to minimize their variance while responding as closely to P2 as possible. They were advised that I considered mean synchronization latencies which differed from the synchronization interval by more than 10 msec to be unacceptable.

At the end of each session, the computer printed out the latency means (measured from P1 onset) and variances for each block (including the first) and for all blocks combined (excluding the first). It also printed a histogram of the response distribution for the session (excluding block 1). Subjects were aware that I expected the distributions to be highly peaked and symmetrical. They were never explicitly told that the distributions should look like triangles, but

some of them read Kristofferson's previous papers, so they probably figured this out for themselves.

Subjects were advised that the foreperiod was variable and were told that the variability was there to make timing from the warning stimulus a poor strategy. They were instructed to time their responses from the onset of P1. Occasionally I mentioned to some subjects that they were relying on the warning stimulus, not on P1, and requested that they pay more attention to P1. The subjects thus knew that I examined the relationship between foreperiod duration and response latency, but they were not told how, nor what I intended to make of this. To their knowledge, my examination of foreperiod effects was done simply to monitor their performance. Especially in Experiment 3, in which responding from P1 was made impossible, some subjects expressed the feeling that they might be timing from the warning stimulus. In such cases, I agreed that the task was difficult, but asked them to try harder to respond on the basis of P1.

Subjects were allowed to examine each others' computer printouts and were not asked to avoid talking with each other about the experiment. A spirit of friendly competition was encouraged throughout the experiment.

During the very first session, I instructed the subject verbally about the stimulus order and then went into the experimental chamber with the subject to demonstrate how to perform the task. The subject made about 30 responses on the first day, with me in the booth to answer questions as necessary. These data were discarded. Session 1 in the tables refers to the first complete session, which was always

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actually the subject's second.

Subjects were paid at a pre-tax rate of \$5 per session. Occasional rewards (such as a pitcher of beer at the graduate pub) were given to some subjects from time to time in recognition of excellent performance.

Sessions lasted about 45 minutes on the PDP-8, with about 8 minutes of that required for generation of the random foreperiods distribution at the start and for analysis, printing and punching out of the data onto paper tape at the end. Sessions lasted about 35 minutes on the Apple, due to faster generation of the random foreperiods at the start of the session, a faster printer, and to the use of floppy disk, rather than paper tape, data storage.

2.8 Scanning Procedure

All of the experiments below are concerned with performance across a range of durations. It is impossible to fully practice subjects at every interval within the ranges. Extended practice can only be-carried out at a few intervals.

Figure 1 is a reproduction of Figure 5 from Kristofferson (1976). In it the variance of synchronization latencies from one subject are plotted as a function of duration and as a function of practice with other stimuli within the range. The subject started with 60 sessions at 940 msec, reaching a stable asymptotic variance of 497 msec². Synchronization intervals were then reduced by 10 msec per session until the lowest interval, 160 msec, was reached. Intervals were then increased, by varying amounts, and were kept constant for 3

FIGURE 1

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Synchronization variances from Kristofferson, 1976 (reproduction of his Figure 5). The figure shows mean within-block variance as a function of mean latency. Filled circles are from the initial descending series, with 300 latencies per point. Open circles are from the ascending series, after the first session at each interval, and are based of a minimum of 900 latencies per point.



to 10 sessions. Study of performance over the range as the interval is decreased gradually can be called "scanning down" the range. The graph shows the results from an initial scan down (filled circles) followed by a scan up (empty circles).

Variances of this subject appear constant in the 300 - 550 msec region and appear to increase as the interval is changed in either direction outside this range. Variance levels from another subject were constant from about 170 msec to 550 msec, at about the same level as obtained by this subject from 300 to 550 msec. The effect of scanning was to reduce variability across the range, but especially to reduce it in the regions in which it was a non-constant function of duration. One might speculate that with even more extensive practice, the relationship between variance and duration might flatten out completely, across the 160 to 940 msec range shown. The reduction in variance at 940 msec was from an apparently stable asymptotic level of 497 msec² to a final level of less than 300 msec², due to practice at durations other than 940 msec.

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Kristofferson (1983) studied discrimination performance as the separation between pairs of stimuli on opposite sides of the midpoint (base duration) was varied. The estimates of q were based on performance with these stimuli and with others left alone from condition to condition (given a constant midpoint) in this single stimulus paradigm. Initially, the estimated value of q varied with stimulus separation even though base duration was unchanged. Scanning up and down the range of separations had to be done twice before these estimates of internal variability appeared constant, which was

eventually obtained.

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Scanning across a range of values appears to be a highly effective training strategy for reducing variances and it may be more efficient than further training at the same interval, once asymptotic / performance at this interval appears to have been reached.

I made extensive use of a scanning strategy in all of the experiments. After initial stabilization training at a specific interval for at least 50 sessions, synchronization intervals were gradually increased or decreased toward the other end of the range for that experiment. Typically extended practice was also given at one or a few intermediate intervals within the range. Scanning was typically carried out to extreme durations just beyond the hongest or shortest intervals at which extensive practice would be required, to minimize any end effects. In general, I was interested in performance levels at the well-practiced intervals. Except in Experiment 4 (vision), I was only concerned with performance at these intervals, spending sessions at the others only as a means of improving performance at the Key ones.

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The details of the scanning procedures used varied from experiment to experiment and, especially, from subject to subject. Such variation between subjects was absolutely necessary. Some subjects were willing to participate in over 600 sessions. Most were unwilling to commit themselves beyond 200 sessions. Interested in asymptotic performance levels, I adopted what seemed to be the most thorough training schedule possible given the number of sessions that could be expected from the subject and the subject's rate of improvement in the task.
2.9 Data Truncation and Lost Data

The raw data from some sessions were lost due to equipment problems. The PDP-8 paper tape punch generated a number of unreadable tapes, leaving the printout of means and variances at the end of the session as the only record of the data. The Apple disks were substantially more reliable but the computer "crashed" a few times on encountering bad sectors on disks. Further, one of the drives needed more frequent speed calibration than it originally received, and it wrote some unreadable files. Eventually'I solved this and other difficulties with the Apple by training at Apple Canada to be a Level 1 Service Technician, after which I calibrated the drives and did other preventative maintenance myself on a regular basis. In the session by session graphs of the data, lost sessions are indicated by gaps. When mean latencies and their variances were obtainable from printouts, these were included in the appropriate graphs.

A number of other problems arose which introduced outliers (extreme latencies) into the data. On the equipment side, electrical transients produced when P1 was presented sometimes caused the PDP-8 to record responses at the time of P1, even though the subject responded much later, near P2. A mixer through which the auditory P1 and P2 stimuli were passed before being presented to the subjects in the PDP-8 setup was a source of intermittent noise for an undetermined number of sessions. The problem was corrected once it became regular enough that the subjects and I realized were was a problem. The effect was that

sometimes a short noise burst produced by the mixer was interpreted by the subject as P1 or P2. Noise bursts occured at random times. The PDP-8 suffered a variety of timing problems, all intermittent, at different times. The typical result was a random reduction in recorded response latency. The original response pad occasionally generated its own responses, or responded to movements of the hand in the vicinityof, but not on, the button. The replacement was originally less sensitive than the first one, and contact with the button could be maintained without maintenance of closure of the electrical circuit. Even with the fixed pad, some subjects required Redux Cream, an electrolytic cream, to increase the conductivity of their hand. Use of too much cream made the button sticky, which interfered with performance. The Apple also failed from time to time, requiring over the course of the experiment a new power supply, two new motherboards, and various memory and other chip replacements. Intermittent and total failures to present stimuli were uncommon but they arose. DK's session 395 has been excluded from analysis because P1 and P2 were not -presented during the last hundred trials, due to a Aaulty connection. In a few other sessions (3 or 4), the same thing happened-but the subjects terminated the session early, leaving no data to exclude from analysis. In perhaps 5 further sessions, P1 and P2 were not presented for very few trials. These data were kept: I relied on the truncation algorithm described below to automatically remove the #abberant responses obtained in these trials.

On the subjects' side, sneezes were the main cause of outliers. Subjects also occasionally leaned back on the chair at a strange angle,

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ending up on the floor, making an unintentional response by removing the hand from the response pad in the process. Some subjects deliberately "blew" the odd trial, withholding a response in order to hear or see the interval. This was usually no problem since no-response trials were discarded. Occasionally, however, the subject would accidentally lift her finger during the post-P2 interval, generating an accidental and spuriously long response.

There were other problems, but this is a sufficient list of examples. The vast majority of the data were uncontaminated but over the course of a million trials such problems should be expected.

The problem of transients in the PDP-8 system arose in Hopkins' and Brewster's work as well, and had been so for some years before the present research began. In their data, it appeared as a measured latency of 10 msec, i.e. as a response coincident with the end of P1, and they routinely eliminated 10 msec latencies from their data. This was the earliest latency that their programs could report. Three or four such latencies were often obtained in a single session. In my situation, for other technical reasons, response-latency timing was from the start of the foreperiod, so reports of latencies shorter than 10 msec, including negative ones (responses preceding P1) were possible. Remember the task here. The subject is to time an interval that begins with P1 and ends with P2, responding in synchrony with P2. If the response occurs at or before P1, according to the computer, this may reflect a transient or a sneeze, but whatever the difficulty, this is not a synchronization response. I decided to deal with this ⁵ directly, on an <u>a priori</u> basis, as part of the design of the

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experiment. A response which the computer claimed occurred at or before P1 offset was flagged immediately with "blown trial" feedback, the abberant latency was stored in a separate data array in the program, and the trial was re-run. My goal in this was to obtain 300 good trials per session and to remove the "bad" trials (the "blown" " ones) from view completely. Confident that extremely short and negative latencies could not be "real", I had no intention of including them in any analyses.

For P1-P2 intervals longer than 200 msec, I am still confident that this reasoning was correct, and these "blown" trials have not been analysed.

For (P1-P2) intervals of 200 msec or less, I am not confident that all "blown" trials should be removed automatically. The difficulty is clearest when P1-P2 is a mere 85 msec, which was used in Experiment 3 (designed long after Experiments 1 and 2 were started). In this case, the subject is almost certainly timing from WS (the warning stimulus), rather than from P1. For example, with a mean foreperiod duration of 900 msec, the subject might try to wait 985 msec from offset of WS before responding, independently of the time of occurrence of P1. I will consider this strategy in Chapter 4. The point here is that if this is the subject's strategy, and if there is an extraordinarily long foreperiod in a given trial, the subject's response time relative to the long delayed P1 could indeed be 10 msec or less. When P1-P2 is very short, the subjects' response distributions have long negative tails. There is little justification for truncating these tails when they extend past 10 msec.

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Even with the addition of foreperiod variance in the response distribution, when timing is from WS rather than from P1, there is a limit on how far back "plausible" latencies can extend. A latency of 10 msec might be believable when P1-P2 is 85 msec. It is less believable when P1-P2 is 150 msec, especially when the low variances and symmetry of the response distributions at this interval are considered, except for subject KC. Even at 200 msec, KC's performance was highly variable and her latencies were strongly correlated with . foreperiod duration, suggesting that she was timing from WS rather than from P1. It seemed improbable, but not sufficiently extremely improbable, that responses at or before P1 offset could have been part of her response distribution. KC's behavior in the task was sufficiently abberant that I eventually asked her to leave the experiment, a point to which I'll return in Chapter 4. Her data were analysed, and summary tables appear in Appendix 1. To avoid missing latencies that might have been "real", I included in the analyses any response that preceded P2 by 200 msec or less (her longest P1-P2 interval was, 200 msec). To avoid inconsistency of treatment of the data from different subjects, I adopted the same rule for all of them.

To summarize, on an <u>a priori</u> basis, I decided to exclude from analysis all responses which occurred at or before P1 offset. This decision was made before Experiment 3 was designed and the criterion was inappropriate for that experiment. I modified it as follows: all responses which occurred at or before P1 offset <u>and</u> which preceded P2 by more than 200 msec were excluded from analysis. They are not reflected in any tables or graphs below, nor will I refer to them again

in the thesis. This restriction on truncation left many trans.ent-based outliers in the data of Experiments 2 and 3.

I spent considerable time thinking about how to remove these remaining transient-based short latencies and other outliers which arose from the problems discussed at the start of this section. I object in principle to data elimination on the basis of the "eyeball test". Further, the notes about problems that subjects often made on their sessions' data printouts were not an adequate basis for a consistent elimination criterion, though I made much use of them in checking the effects of different outlier removal criteria that I considered.

After a number of false starts, I settled on a kurtosis criterion for removal of outliers. If M is the sample mean of a distribution of responses, R_i , and V is the sample variance, the sample kurtosis is defined as b_2 where

 $b_2 = \Sigma (R_1 - M) 4 / V^2$.

This statistic is quite sensitive to extreme values, becoming very large when one or more outliers are present in the data. The kurtosis criterion for removal of outliers is used as follows: Calculate the sample kurtosis. If it exceeds a critical value, exclude the datum most distant from the sample mean. Recalculate the Kurtosis, continuing to remove the most extreme of the remaining values until the kurtosis drops below the critical value. This is a common and powerful test for outliers (see Barnett and Lewis, 1978, for more details.)

The difficulty with this procedure, for present purposes, lies in choice of a critical value for the kurtosis. Simulation-based

estimates of the Kurtosis distribution are available when the response distribution is normal (eg. D'Agostino and Pearson, 1973), but they are not available for most other distributions. Accordingly, I ran a small scale simulation to obtain critical values for this procedure. If the subject is timing from P1, the response distribution is known to be symmetrical and highly peaked. Pseudorandom variates were generated in batches of 300, to make up a simulated session, and 200 such sessions of data were examined for each distribution studied.

To approximate the response distribution when the subject is timing from P1, I used normal (population Kurtosis of 3.0), triangular (Kurtosis of 2.4), and logistic (4.2) distributions, and three convolutions of triangular and logistic distributions (Kurtosis ranging from 2.4 to 4.2, depending on the relative variances of the two distributions). Across the 1200 "sessions" of data so examined, sample kurtosis never exceeded 8.3. On the basis of this, I conclude that if the subject is timing from P1, it is exceedingly unlikely that the sample kurtosis of his data would exceed 10 by chance. For this case, then, I use a kurtosis criterion of 10.

If the subject is timing from WS, his response distribution includes the variability of the foreperiod. If we measure response latency from WS, the distribution should be symmetrical and peaked, as it is from P1 when the subject times from P1. However, if we measure latency from P1 and the subject is timing from WS, we are subtracting the foreperiod duration from the interval the subject is actually timing. This is a geometric variate, with a variance of 90 msec². I used the same base distributions (triangle, logistic, convolution of

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the two, normal), and subtracted geometric variates to simulate the response latency distribution to be expected in this case. The Kurtosis of such distributions depends on the relative variance of the timing and foreperiod distributions, so I varied these as well. In a total of 17 distributions (3400 Kurtosis values calculated), 10 was exceeded five times. That is, a Kurtosis of 10 should be rare if the subject times from WS and we measure the latency from P1, but it is not extremely rare. The same results would be expected if the subject timed from P1 and we measured latency from WS, adding the geometric foreperiod to the latencies, rather than subtracting it from them.

It is not always possible to decide beforehand whether a given subject at a given P1-P2 interval will time from P1 or from WS. To avoid arbitrary decisions here, I used a compound criterion. If the Kurtosis of the latency distribution exceeded 10 when latencies were measured from P1 onset, and if it also exceeded 10 when latencies were measured from foreperiod onset, then I eliminated the most extreme latency. In principle, one of these two latency distributions should be symmetrical and highly peaked while the other should be skewed and less peaked due to the addition or subtraction of a geometric variate, whether the subject is timing from P1 or from WS. Values as large as 10 from the symmetric distributions were never obtained in the skewed distributions were this large.

I consider this compound criterion quite conservative. Further, from examination of the histograms of the response distributions printed for each session, and from the subjects' notes, I

am satisfied that the latencies which were eliminated should have been eliminated. A few outliers remain in the data, which can be directly traced to known equipment problems, but relaxing the Kurtosis criteria to allow elimination of these responses would have resulted in elimination of others that should not be removed from the data.

As a final note on the simulations, I derived the population kurtosis for each of the distributions studied, and compared this to the estimate obtained by simulation (along with estimates of other statistics of the distributions). The theoretical and obtained values agreed well, as they must for the simulation to be considered valid.

Appendix 1 includes tables of a number of statistics of the response distributions for each subject and session. Included in this is the number of trials per session that were included in the analysis. If any trials were eliminated from that session's data, the number of these is given in Brackets beside the number included. A total of 186 responses were eliminated, from 3,116 sessions' data (or about 1 response per 5,000). Reflecting the string of problems with the machine, 146 of the 186 eliminated responses were from data collected on the PDP-8, which controlled only 756 of the 3,116 sessions.

3. Experiment 1: Long Auditory Intervals

In this experiment, synchronization intervals were varied across a range from 1400 to 2100 msec. On the basis of the review in the introduction, three plausible results can be envisioned:

From Kristofferson's (1976) synchronization experiment, we should expect variability to increase continuously as the P1-P2 interval is increased. Kristofferson found that beyond 550 msec, the standard deviation of synchronization latency distributions increased linearly with duration. (More precisely, the standard deviation of the added delay, in excess of 550 msec, was directly proportional to the mean of the added delay.) Only one subject was studied in that experiment. His variances in the 1400 to 2100 msec region ranged from 888 msec² at 1390 msec to 3283 msec² at an interval of 2090 msec.

On the basis of Kristofferson's (1980, 1983) discrimination research (which involved the same subject), we should expect Weber's Law compatibility early in the experiment but after extended practice the variance should increase in a stepwise manner with duration, rather than continuously. Kristofferson (1980, 1983) estimated q, the parameter of the internal triangular distribution, as 100 msec across the range from 800 to 1600 msec, and as 200 from 1600 upwards.

Translation of the discrimination results into predictions about synchronization latency variances involves a few steps. First, the breakpoint should probably not be at 1600 msec because some time is

taken for the subject to perceive P1 and to respond to it. If we parse the stimulus-response chain into three stages, afferent (input), central, and efferent (output), and assume that the internally timed interval starts after the stimulus is registered and ends with the initiation of the response, we should subtract afferent and efferent latencies to estimate the duration of the internal interval timed. Following Kristofferson (1976), we can estimate the non-central latencies as 150 msec, close to the mean latency of simple reaction time responses when using 2000 Hz 68 dB auditory stimuli. (The parameters are important because RT varies with intensity and frequency.) In this case, we should expect a break from g=100 to g=200 at a synchronization interval of about 1750 msec. The triangular variances (given by $V = q^2/6$) are 1667 msec² and 6667 msec² respectively. To these, a further 10 to 50 msec² should be added to reflect efferent latency variance of well practiced subjects (Wing and, Kristofferson, 1973). Wing's (1973) results depended on the type of response demanded of the subject. Hopkins (1982) estimated efferent latency variances using the same equipment and response as were used in the present experiment. His estimates averaged approximately 25 msec². Accordingly, we should expect the variances to be about 1690 msec² from 1400 to 1750 msec (1600 + 150) and about 6790 msec² for intervals ranging from 1750 to 2100 msec⊁

These values are substantially larger than many long interval synchronization variances reported by Kristofferson (1976), so a prediction of complete compatibility with the discrimination results o Kristofferson (1980, 1983) is not plausible. We might still anticipate a stepwise increase in variance, however, with a break at 1750 msec.

Kristofferson's (1980) explanation of the stepwise variance increase was in terms of a quantal counting model, with a maximum count of 16. Hopkins (1982) reported synchronization variances as low as 33 msec² for intervals as long as 460 msec. With a q of 12.5 msec (close to Hopkins' estimates) and subtracting 150 msec from the 460 to reflect input and output delays, we still obtain a count of over 24 q. If we maintain a quantal counting model, we must drop the maximum of 16 for the count. This restates the difficulty noted in the last paragraph: the variances are too small to be fully compatible with Kristofferson's (1980) discrimination findings.

Brewster (1983) showed, in a task involving both synchronization and discrimination responses, that variability of each type of response could be manipulated independently of the variability associated with the other. She interpreted this as indicating that different timing mechanisms or strategies were involved in the two tasks. These results give us some leeway to ignore the discrimination findings. The third outcome that I will raise as "plausible" is that the variances may stay constant across the entire interval.

My expectation in this experiment was that the results would be similar to those obtained in most other very long term experiments on subjective duration and/or timing. With respect to variability, I expected that after sufficient practice the relation between synchronization interval and variance would be constant or stepwise increasing, not monotonically increasing. This is not a foregone

conclusion. Kristofferson (1976) included very extensive practice at 940 msec, but despite that practice the variance at this interval was higher than that obtained at 890 msec, and lower than that gbtained when the synchronization interval was 990 msec (the two surrounding intervals in this experiment). It appeared from that experiment that Weber's Law would describe asymptotic performance. I considered this result anomalous and, as discussed in the introduction, I did not expect to obtain it.

There is more to "similarity of results" than similarity in terms of constant or stepwise increasing variance. My thesis is that the same timing mechanism is available for long interval synchronization as is used for short intervals. If this is true, we should find that estimates of q in this experiment hover about 12.5, 25, 50, or 100 msec, after much practice, and that the distribution of responses is symmetrical and peaked. From Hopkins (1982), I would expect that convolution of a triangular (timing) with a logistic ((efferent latency) variate should yield a theoretical distribution that describes the response latencies. Population statistics of this and other distributions are given in Appendix 2. I evaluate the quality of this description by examining the coefficients of symmetry, $\sqrt{b_1}$, which should be 0, and of Kurtosis, b_2 , which should be between 2.4 and 4.2.

I will not specify the mechanism beyond the statement that it is quantal and that it yields response distributions as described above. All that we have to work with from this task are the response distributions, and different quantal mechanisms could generate the same

results. Hopkins' model, involving effectively independent uniform delays in getting information into and out of the clock, perfect timing by the clock, plus an independent logistic variate describing motor variance, is the one I adopt as a working model. There is no hint of quantal counting in this model. I am confident that a counting model could be constructed to predict exactly the same results in this task.

3.1 Method

3.1.1 Subjects

Two male subjects, DG and GH, participated in this experiment. DG was a senior undergraduate, GH a graduate student. As is true for all of the subjects in this thesis, neither had participated in synchronization experiments previously. Both had some familiarity with the timing literature and had participated in various other psychological experiments.

3.1.2 Procedure

The general procedure was described in Chapter 2. DG started at 2000 msec and practiced extensively at this interval before moving toward 1500 msec, in steps of 25 msec. There was only one session per interval for most of the intermediate intervals. I scheduled 5 sessions per interval at 1900, 1800, 1700 and 1600 msec to slow the rate of change somewhat. My intent in designing the schedule was to change the interval to 1500 msec quickly but not so quickly that the benefits of practice at the longer intervals would be lost at the shorter intervals. After 5 sessions at 1500 msec, the interval was reduced in steps of 25 msec to 1400 msec, to avoid any end effects at 1500. The interval was then lengthened, always in 25 msec steps, to 1500, where DG stayed for 30 sessions. The interval was then brought up to 1750 msec, and after extended practice it was lengthened to 2000 msec where it was kept for the final 52 sessions of the experiment.

The schedule for GH was similar, except that he started at 1500 msec, worked with intervals as long as 2100 msec, and finished at 1500.

In both cases, attention was paid to performance at 1750 msec, as compared to that obtained at 1500 and 2000 msec. My intention was to study the region around 1750 msec in detail if it appeared that there was a variance step between 1500 and 1750 or between 1750 and 2000 msec.

The exact schedule for each subject is displayed visually in numerous figures below. Appendix 1 provides tables of descriptive statistics for each session of each subject, and these list each session's synchronization interval.

3.2 Results

_3.2.1 Vatencies

Figures 2.1 and 2.2 show the relationship between synchronization interval (lower panel) and mean synchronization latency (upper panel) for each subject. All sessions are plotted. Clearly, response latencies are linearly related to the synchronization intervals. It is not evident in the graphs, but can be seen in the first two tables of Appendix 1, that DG's mean latencies were typically within 10 msec of perfect synchrony. This is typical of

synchronization performance, in the data of this dissertation and in

the other works cited previously. GH's responses typically occurred more than 10 msec before P2. The means were often 20 to 30 msec early. Still, increases and decreases in GH's latencies followed increases and decreases in the duration of the P1-P2 interval quite closely.

3.2.2 Variances

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The upper panel of Figures 3.1 and 3.2 show DG's and GH's variances for each session. The middle panels again show synchronization intervals. The lower panels show the mean foreperiod durations, near 900 msec on the PDP-8 and near 1000 on the Apple. There is no indication of a change in performance due to the change of equipment.

Figures 4.1 and 4.2 show variance levels at "practiced" intervals. I refer to statistics of "practiced intervals" frequently below. By this, 1 mean all intervals with which the subject worked for two or more consecutive sessions. The figures show averages of the within-session variances of the last five sessions, or of all sessions if there were fewer than five at this interval.⁴ These averaged variances are presented in tabular format in Tables 1.1 and 1.2, along with the intervals, averaged estimates of q, the mean symmetry and kurtosis coefficients, and the mean correlation between response latency and foreperiod duration. There are clear practice effects in the data, so these tables list the results in the order in which they were obtained. The final few rows in each table are the ones of major interest.

DG's and GH's data contain some striking dissimilarities, so I



Upper panel: Mean synchronization latencies, in milliseconds, of each session.

Figure 2.1, subject DG. Figure 2:2, Subject GH.

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FIGÙRE 3

Upper panel: Response latency variances, in msec², of each & session. Note that there are two lines marking the top of the panel. Variances shown as extending to the topmost line were too large to show accurately on this scale. For exact values, see Appendix 1.

Middle panel: Synchronization interval, in milliseconds, of each session.

Lower panel: Mean foreperiod duration, in milliseconds, of each session. The means were approximately 900 msec when the PDP-8 controlled sessions and 1000 msec when the Apple controlled sessions. Gaps in the figure (ostensibly 800 msec means) are due to loss of the raw data, making calculation of that session's mean foreperiod duration impossible.

Figure 3.1: Subject DG. Figure 3.2: Subject GH.



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Mean within-session response latency variances (msec²) as a function of mean response latency (msec). Filled squares show the descending series variances. Filled cirles show ascending series variances. The variances plotted are listed in Table 1.

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Figure 4.1: Subject DG. Figure 4.2: Subject GH.

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TABLE 1

Mean within-session variances (msec²), q estimates (msec), coefficients of symmetry (ROOT B1), of Kurtosis (B2) and of correlation between response latency measured from P1 and the foreperiod duration. Only data from intervals at which the subject spent two or more consecutive sessions were averaged for this table. The averages are based on the last five sessions at this interval before the subject moved to the next, or, if there were fewer than five, on all sessions at the interval.4

Estimates of q were obtained from the triangular variance relationship, $q^2 = Variance/6$, after subtraction of 25 msec² as an estimate of efferent latency variance. That is, if V is the sample variance, q was estimated from

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 $q^2 = (V - 25)/6.$

Table 1.1: Subject DG. Table 1.2: Subject GH.

SUMMARY STATISTICS FOR DG AUDITORY RESPONSE LATENCY MEASURED FROM P1

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INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH F	P
2000	1604	97.34	0.28	4.84	-0.03	
1925	1098	80.23	0.00	3.15	-0.05	
1900	1042	78.11	0.07	3.12	-0.01	
1880	988	76.01	0.14	3.09	-0.08	
1700	752	66.04	0.09	3.02	-0.05	
1600	813	68.77	0.19	3.08	0.01	
1500	798	68.11	-0.02	3.24	-0.03	
1400	750	65.94	-0.03	3.78	-0.03	
1425	729	64.98	0.00	3.49	-0.02	
1450	690	63.18	-0.09	3.37	-0.02	
1475	699	63.60	0.00	3.27	-0.05	
1500	619	59.68	0.05	5.67	-0.03	
1525	611	59.28	0003	2.98	-0.06	
- 1500	463	51.29	0.05	3.84	-0.04	
1750	457	50,90	-0.03	2.98	-0.05	.:
2000	594	58.42	0.02	2.99	0.02	•

SUMMARY STATISTICS FOR GH AUDITORY RESPONSE LATENCY MEASURED FROM P1

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INTERVAL	VARIANCE	Q	ROOT B1	82	CORR WITH FP			
1500	1054	70 50	0 00	2.20.				
1000	1004	18.37	0.02	3.28	0.04			
1600	1214	84.47	0.05	3.25	0.01			
1700	1786	102.79	0.04	3.52	0.03			
1800	1920	106.62	-0.02	3,25	0.04			
1980	2569	123.54	0.09	3.11	0.03			
2000	3152	136.98	0.02	3.13	0.04			
2100	3660	147.68	-0.04	3.13	0.10			
2000	1475	93.28	0.15	2.89	0,.04			
1750	784	67.49	-0.02	3.26	0.03			
1500	674	62.42	-0.05	3.60	-0.01			
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will discuss their results separately.

By about session 140, DG had achieved a stable level of variability, hovering about 500 msecf. The variances at 1500 and 1750 msec are quite close (averaging 463 msec² at 1500 and 457 msec² at 1750, for the final five sessions). There is a slight increase at 2000 msec, to a final five session average of 594 msec², but there is no hint of a stepwise increase. He and I both felt that with further practice (impossible because DG had to leave to attend graduate school in California) this variance would have dropped further.

Figure 4, based on the variances reported in Table 1, highlights the effect of "scanning" on the data. DG's variance at 2000 msec appeared roughly stable at 1604 msec² before he practiced at the shorter intervals. After working with the shorter ones, his variance at 2000 was substantially less than half this value, and it was so even during the first sessions of the final series at 2000 msec (see Figure 3.1).

DG's q was very close to 50 msec at 1500 msec (51.3) and at / 1750 (50.9). It was not much greater than 50 (58.4) at 2000 msec.

If DG was timing these intervals by counting quanta, by these estimates, at 1750 and at 2000 msec he was counting about 34 of them. There is absolutely no support in these data for the hypothesis of a maximum possible count of 16 q.

The Weber ratios (standard deviation/synchronization interval) for the final 5 sessions at 1508, 1750 and 2000 msec are 0.014, 0.012 and 0.012 respectively. Hopkins (1982) reported ratios as low as 0.016

in his subjects' synchronization performance. The lowest value value obtained previously was 0.023, from Wing and Kristofferson (1973). Weber ratios obtained in discrimination tasks typically range from 0.06 to 0.10 according to Hopkins' review. Compared to these values, DG's performance is quite impressive. It is equally impressive when compared to the synchronization variances reported by Kristofferson. (1926), over the same range of intervals (888 to 3283 msec²).

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GH's performance was substantially more variable, in all respects. I think that part of the problem was that GH spread the sessions over a much longer period (he took almost a year longer than DG to complete the experiment). Further, he reported as late as session 187 that he was still experimenting with new response strategies, whereas most of the other subjects had settled into a routine approach much earlier in training. Whatever the problem, it is not clear that GH ever achieved a stable performance level. Some of his worst, and some of his best sessions occurred in his final ten.

Figure 3.2 paints a confusing picture of the relationship between synchronization interval and his variance. There are striking practice effects. The variance at 2000 msec dropped by more than half with gractice. Variability declines from 2000 to 1750 msec, but this appears to be a continuous decline. Perhaps with more practice this would flatten out and we would see a step. Perhaps not. From 1750 msec downwards, the variance appears roughly constant.

Given the day to day high variability of GH's performance from day to day, I decided not to schedule a further series of sessions to examine the 1750 to 2000 msec region for a step in variance levels. Even if there is one, without years' more data collection, I doubt that it would be evident in the results. $\sqrt{2}$

Table 1.2 and Figure 4.2 give the variances at the final five , sessions at 1500, 1750 and 2000 msec intervals. It is hard to interpret the values at 1500 and 1750 msec. I think they are fortuitously low. The variance of the second last 5 sessions at 1500 msec was 1737 msec² (q = 101.4 msec), compared to 674 msec² in his fast 5. At 1750 msec, the second last group of 5 yielded an average variance of 1142 msec² (q = 81.9 msec), compared to 784 msec² for the last group. I'm not saying that these are in any way "better" or "more representative" variance estimates. Instead, I'm saying that I'm not sure that values ("representative" of asymptotic performance levels can be found in these data. Nor am I confident that further data collection would lead us to better data from GH.

If we do look only at the last 5 sessions at 1500, 1750 and 2000 msec, the Weber ratios are 0.017, 0.016 and 0.019 respectively. While not as good as DG's, these are as good as those reported by Hopkins' (1982) and are better than any reported previous to them. They are very respectable results. Further, GH's variances throughout the descending series (from 2100 msec down to 1500 msec) were typically better than those Peported by Kristofferson (1976).

3.2.3 Weber's Law

Kristofferson (1976) reported that increases in synchronization wariance beyond 550 msec followed Weber's Law, but there was very little practice at most of these intervals. The same was true for intervals in the 150 to 550 msec range, but there, constant variances were found, suggesting a transition in mechanisms. Kristofferson (1980) found that Weber's Law described discrimination performance quite well if only the first five sessions at each base duration were considered. In the present data, for both subjects, variance appeared to increase with duration before the subjects had practiced at the intervals. Perhaps Weber's Law should describe their unpracticed performance as well.

I checked this notion by examining the variances obtained during the first session at each interval studied during the first series (ascending for GH and descending for DG). I also excluded from this analysis data from the first interval studied (2000 msec for DG and 1500 for GH) because variances during the very first sessions were high due to the newness of the task, independently of the interval. I also excluded those sessions from which the raw data were lost (due to paper tape or disk failures) because the usual truncation procedures could not be carried out on these data. This sometimes resulted in artificially inflated variances, at least relative to the others, which would have clouded any relationship that-exists between duration and variability.

To examine the fit of Weber's Law to the data, I fit a simple linear function to the mean latencies and the associated standard deviations. According to the modified form of Weber's Law,

Standard Deviation = k (mean latency) + c where k and c are constants.

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Table 2 presents the estimated Weber fractions, k, and the

Pearson correlations between standard deviations and mean latencies, for DG, GH, and for the main subjects of the later experiments, whose intervals ranged from 150 to 525 msec.⁵ From the results of Kristofferson (1976), we should expect the correlations from the shorter interval experiments to be close to zero, whereas we should expect high correlations between standard deviation and mean latency for -DG and GH.

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Also presented in Table 2 are the correlations between mean latency and latency variance across the same sessions. I calculated these out of curiousity about comparisons between Creelman's theory and Weber's Law. Getty (1976) and Kristofferson (1976) reported that their standard deviations increased linearly with duration and that the variances increased nonlinearly. Other reports (eg. Creelman, 1962) indicate that variances can be described as increasing linearly with the mean, but to the best of my knowledge none of these explicitly tested for a nonlinear relationship between standard deviation and mean. The question is not sufficiently relevant to the data or to the main questions at hand to justify a simulation or other extended study here, but what I am curious about is whether, with noisy data, if both relationships were tested, it might appear that both are true. In that case, evidence of a linear increase in variance without concomitant evidence of nonlinearity in the standard deviations might provide only equivocal support for a Poisson process. As Kristofferson has stressed in discussions with mey the data might be equally compatible with Weber's Law, for which there is more supporting evidence.

Table 2√

		 Correlat 	ions Betwe	en Response Lat	encies				
	and Their Standard deviations and Variances								
	- · · -								
	Subject	interva 🛏 Weber		Correlation Between Mean an					
		Range	Fraction	Stand, Dev.	Varji ance				
	DG	1408-2000	.016	.823	.815				
	GH	1500-2100	.046	.942	.934				
	RW	150-210	.081	. 798	.788				
/	-JB)	150-200	.040	.277 1	.305				
(DK 💛	300-500	.022	.453 1	.437				
1	IS	300-500	.030	.626	.628				
	LL	300-500	.000	.013	og 12				
	RW Visual	300-400	.040	.649	.643				

The correlations between mean latency and either measure of variance are always positive, indicating that variance is an increasing function of duration for unpracticed performance. For the short intervals these correlations are not large, which suggests that neither the relationship between mean and standard deviation nor the relationship between mean and variance should be taken as linear.

The correlations are larger for the long durations (DG and GH), but here, as for the short P1-P2 interval data, the correlations between mean and standard deviation and between mean and variance are the same. Perhaps one of these relationships is linear, but it is impossible to tell which one. Looking at graphs for both subjects (not presented here but easily constructed from the data in Appendix 1), I am not convinced of linearity in either case. I should stress, though, that there is a great deal of random fluctuation in the variances, which are only based on a single session's data at each interval. Given this noise, I think the correlations are large enough to support

the claim that the long interval data are compatible with (if not strongly suggestive of) Weber's Law, and with linearity between mean and variance.

The results of these analyses also provide some further grounds, albeit modest ones, for querying whether results thought of as compatible with a Poisson process model might not also be compatible with Weber's Law. The similarity of correlations between mean and vaciance and between mean and standard deviation.is guite striking.

3.2.4 Correlation Between Response Latency and Foreperiod Duration There is no reason for the subject to time from the warning stimulus when the P1-P2 interval is 1400 msec or longer, so there should be no correlation between latency and foreperiod duration. The correlations are of interest in the later experiments so for comparison I report them here as well. Figures 5.1 and 5.2 display the correlations (upper panel) for each session. These are listed in Appendix 1 as well. Table 1 presents some averages across sessions, at practiced intervals.

There is no relationship between correlation and synchronization interval, but the correlations, while very small, are consistently non-zero for both subjects. DG's correlations are negative, which would arise if he was timing from the warning stimulus (though they would be larger if he did this at all consistently). GH's correlations are fairly consistently positive. The correlations are quite small, and they account for very little variance. I will . generally ignore them in discussing the data.

FIGURE 5

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Upper panel: Correlation between response latency (from P1) and foreperiod duration for each session. Some sessions' raw data were lost and correlations could not be calculated. Gaps in the figure are due to lost sessions.

Lower panel: synchronization intervals, in milliseconds, of each

Figure 5.1: Subject DG. Figure 5.2: Subject GH.

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FIGURE 6

Upper panel: Symmetry coefficients of the response latency (from P1) distributions of each session.

Middle panel: Kurtosis coefficients of the response latency distributions of each session. Coefficients shown as extending to the topmost line of the figure were too large to be represented accurately at this scale. For exact values, see Appendix⁶1.

Some sessions raw data were lost and symmetry and kurtosis coefficients could not be calculated. No values are plotted for these sessions. The lost sessions are indicated by gaps in the figures.

Lower panel: synchronization intervals, in milliseconds, of each ζ session.

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Figure 6.1: Subject DG. Figure 6.2: Subject GH.



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3.2.5 Distributional Shape

Table 1 presents symmetry and Kurtosis coefficient averages for the practiced intervals. Session by session listings are available in Appendix 1. Figures 6.1 and 6.2 show these visually.

There is no apparent relationship between skewness and synchronization interval for either subject. The distributions of each subjects responses are trivially but fairly consistently skewed positively rather than being perfectly symmetrical. In the histograms of the response distributions printed for each session, these distributions look quite symmetrical. Hopkins (1982) also reported a small positive skew in his data, with b_1 averaging .025 across all subjects and conditions.

There might be a small trend for DG's Kurtosis to increase as stimulus duration decreases, but the Kurtosis is at least roughly constant across the range, for both subjects.

Kurtosis coefficients of both subjects are typically slightly larger than 3 but they are usually 4.2 or smaller, which is consistent with the hypothesis that the response distribution can be described by a convolution of a triangular and a logistic variate. The very minor but consistent asymmetry suggests that this is not the perfect model for these data, but the imperfection of description may be due to the small foreperiod effects or to sequential effects in the data (as found by Kristofferson, 1976 and Hopkins, 1982).

Since the Kurtosis of the response distributions is almost always larger than 3, a normal distribution would not describe these

data. Nor would the convolution of a triangular variate with a normal variate, rather than a logistic, since the Kurtosis of that convolved distribution should be between 2.4 and 3 (see Appendix 2). Individually, the Kurtosis coefficients are generally close to 3 so this lack of fit would probably not show up in traditional goodness of fit tests such as chi-square. However, the consistency of the slightly too large Kurtosis values weighs against this description.

Despite the fact that the Kurtosis values fall within the range expected for a convolution of a triangular and a logistic variate, their magnitudes are difficult for Hopkins' model. The problem is that the Kurtosis of such a distribution depends on the relative magnitudes of the triangular and logistic variances. The formulae for the variance and Kurtosis of this distribution are given in Appendix 2. Using the notation of that appendix, let b^2 be the logistic variance and let V be the response latency variance, which is assumed to be the variance of the convolution of independent logistic and triangular random variables. Table 3 shows how the Kurtosis of this convolution varies with the ratio b^2/V , i.e. with the proportion of the variance that is contributed by the logistic variate.

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Table 3

Kurtosis of the Logistic + Triangular Distribution As a Function of b^2/V

-	b2/U 0.0 0.1 0.2 0.3 0.4 0.5 0.6 0.7 0.8 0.9	Kurtosis 2.40 2.53 2.66 2.81 2.98 3.15 3.34 3.53 3.74 3.97 4.20	
	1.0	4.20	

Suppose, for example, that the response latency variance is 500 masec² and that the Kurtosis is 3.15. These are common values in DG's data. According to the table, the logistic variable contributes 50% of this variance. Under Hopkins' model, this means that 50% of DG's variance (250 msec²) should be attributed to variability in the response (the efferent latency) itself. The Kurtosis coefficients from GH's data are about the same, so the distributional model would lead us to estimate GH's efferent latency variances as 350 msec² or more. These are not very plausible values.

Hopkins could not encounter this difficulty because his subjects' variances were so low that any reasonable proportion of them would yield logistic variances lower than 50 msec². He reported 14 kurtosis coefficients (1982, p. 180). Their average was 3.045 which is in line with the values reported here.

I am not arguing here that a convolution of a logistic variate and a triangular variate would not fit the data. I have every

confidence that traditional goodness of fit tests would not reject this model. Further, I would not be attall surprised if traditional tests failed to reject a model of this type even if the logistic variance was restricted to low values during parameter estimation.

Hopkins (1982) used a minimum chi-square parameter estimation procedure to estimate q and b in his data. He pointed out that the parameter space associated with this procedure was quite flat. To see just how bad this situation was, I constructed an artificial data set (1500 responses, or 5 sessions' data pooled) that fit a logistic-plus-friangle distribution perfectly according to a chi-square test when the logistic variance was 50% of the total. I then varied the specified proportion of the variance contributed by the logistic, estimated q (and thus the triangular variance) by minimizing chi-square, and noted the final chi-square value obtained. The test, indicating an excellent fit, even when the logistic, which actually contributed 50% of the variance, was specified as contributing only 10%.

On the basis of this result, and of Hopkins' maps of the parameter spaces he studied, I concluded that chi-square testing of symmetrical distributions with reasonable (less than 4) kurtosis coefficients would not be very informative, and I have not conducted such tests on these data. From comparison of the Kolmogorov-Smirnov and chi-square statistics in a different context (Kaner and Lyons, Note 2), I doubt that a standard Kolmogorov-Smirnov test would be any more useful when parameters are estimated from the data. Simulations could

be run to obtain better critical values for either test, given minimum chi-square or minimum Kolmogorov-Smirnov parameter estimation, but by my calculations it would take a few years of Apple computer time to do this properly. Instead of spending (I think wasting) this time, I am quite satisfied to accept the obtained values of symmetry and Kurtosis as evidence of reasonable fit to some convolution, and thus to many convolutions, of logistic and triangular variates. Further, the kurtosis coefficients themselves point out the key problem, which might or might not become evident in a minimum chi-square estimation context:the convolution might describe the data acceptably, and this might be true even if the logistic variance is restricted to a value smaller than 50 msec², but it appears that the logistic variance should be larger, much larger, than this upper limit. On that basis, I suspect that something is wrong with this model.

3.3 Summary

A number of patterns in the data are in general agreement with those found in other experiments whose data were well described by guantal models.

Early in training, there was an increasing relationship between variance and duration, though it is not clear whether Weber's Law provides the correct description of this relationship.

As training continued, DG's variances declined substantially and are roughly constant from 1500 to 2000 msec. The variance at 2000 msec is slightly higher than the final variances at 1500 and 1750 msec, comparable to that obtained at 1500 msec earlier in training. The

value of q appears close to 50 across the 1500 to 2000 msec range, after extended practice.

GH's variances appear to be roughly constant from 1500 to 1750 msec though there is much variability in performance from session to session. The variances increase beyond 1750 msec. The large changes in variance from session to session make it difficult to arrive at a final estimate of q. Past estimates in the literature were based on stable data, which is what quantal theory requires. GH's variances were not stable. It is not clear that the q estimates obtained are meaningful.

The variances of both subjects are lower than those previously reported for long duration synchronization performance, and they are much lower than we would expect from discrimination results at similar intervals. They are higher than those reported by Kristofferson (1976), Hopkins (1982), and Brewster (1983) for shorter intervals, and we will see below that they are higher than variances obtained in the present work at shorter intervals. There may be a step in variance level's somewhere between 150 and 1500 msec. However, as also found by Hopkins (1982), a quantal counting theory with a maximum count of 16 cannot predict this step. If the subjects are counting quanta, they are counting many more than 16 of them.

The response distributions of DG and GH are similarly shaped, as measured by coefficients of symmetry and kurtosis, and the coefficients are quite close to those reported by Hopkins (1982), the only other source of symmetry and kurtosis statistics in the synchronization literature. The distributions appear symmetric and

peaked. They can be described in terms of a convolution of a triangular and a logistic variable, as also found by Hopkins. However, the variance of the logistic appears to be much larger than it should be if we are to interpret this as reflecting motor variance.

The basic phenomena: constant and low variances across a range of intervals, and symmetric distributions with thin tails appear confirmed in these data, for both subjects. The particular model suggested, which allows for two independent quantal stages (the triangle) and an independent motor component as the only sources of variability, may not be correct for these data. If it is not correct for these, given the similarity in shape between these distributions and those reported by Hopkins, it may not be correct for the short synchronization interval performance either.

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4. Short Auditory Intervals: Experiments 2 and 3

These two experiments involved P1-P2 intervals from 85 to 350 msec. It is difficult to draw a rigid boundary between the experiments. Both involved the same subjects. Half of what I call Experiment 2 was conducted before Experiment 3, and half was conducted after it. From the subjects' point of view, this was one integrated experiment. I separate them here because of a conceptual difference between them.

Experiment 2 dealt with subjects' performance in this particular variant of the synchronization paradigm when P1-P2 intervals are short, but not so short that subjects cannot do the task as instructed. It is a control experiment, allowing comparison of variance levels obtained with this procedure to be compared to those obtained by Kristofferson (1976), Michaels (1977), Hopkins (1982; Hopkins and Kristofferson, 1980) and Brewster (1983), who all studied performance at short intervals. Experiment 2 was intended as a bridge between the rest of the literature and Experiments 4 and 5, on visual synchronization.

Experiment 3 was designed to study synchronization performance when the P1-P2 interval is so short that the subject cannot do the task as instructed. It is generally accepted that there is an absolute minimum reaction time to a stimulus. Reaction time to auditory stimuli depends on stimulus intensity. When extremely loud signals are used it is possible to obtain mean reactions as short as 105 msec. Based on

such data, absolute minimum RT has been estimated as 100 to 105 msec (see Kohfeld, 1981, for a recent review). RT to 2000 Hz, 65 dB stimuli has been estimated at approximately 150 msec (Kristofferson, 1976). Presumably, as the P1-P2 interval is reduced below 150 msec, the subject will find it ever more difficult to respond in synchrony with P2 if he waits for P1 to signal P2's imminent occurence. When the P1-P2 interval is as short as 100 msec, this should be impossible for the subject. Instead, the subject should have to start timing from some event preceding P1 in order to respond at the same time as P2.

Hopkins (1982) showed that the subject can use his trial initiation response instead of P1 and that this adds no variance to his responses if there is no variance in the foreperiod. In the present situation, the subject may use that response, or the onset of the 100 msec auditory "honk", the warning stimulus (WS) which follows the response almost Instantaneously, or he may use the offset of WS. All of these events are time-locked in this experiment so it is impossible to tell which controls behavior. All of these events occur before the foreperiod, which contains no other events that the subject could use as a response trigger, and which ends with P1. In the present context, the most conservative assumption is that if the subject starts timing from some event preceding P1, he is timing from the offset of the warning stimulus, rather than from the onset or the initiation response, which precedes it by 100 msecly. To avoid redundant and awkward constructions below, I will restrict attention to timing from the offset of WS, ignoring timing from the initiation response and from

WS onset, when considering timing from events preceding P1.

The interval between WS and P2 contains a long foreperiod. If the P1-P2 interval is 100 msec then on average the WS-P2 interval, the "true" synchronization interval if the subject times from WS, is 1100 msec (1000 msec on the PDP-8). This is not too far from the intervals of Experiment 1, which were as short as 1400 msec. While Experiment 1 gives us grounds to question the applicability of discrimination-based variance estimates to synchronization performance, if we do follow Kristofferson's (1980) results and reasoning we should expect timing variance of practiced subjects to be the same at 1100 msec it is at 1500 msec. The finding of constant variance across the range from 1500 to 2000 msec for DG (his variance at 1400 was slightly higher, due to less practice at this interval), and of constant variance from 1500 to 1750 msec for GH lends credence to the idea that variance at 1100 msec is the same as that obtained with intervals a few hundred msec longer.

In effect, then, Experiment 3 is another study of long duration synchronization performance, with intervals which may be in the same range, so far as variance is concerned, as those of Experiment 1. The Key difference between Experiment 1 and Experiment 3 lies in the instructions to the subject. In Experiment 1, the subjects knew they were timing long intervals. In Experiment 3, the subjects knew they were supposed to be timing short intervals.

In Experiment 1, DG's variances averaged about 500 msec². GH's variances were higher, as were those reported by Kristofferson (1976) for long intervals. DG's lowest variances were just above 300 msec². Variances at shorter intervals are typically smaller.

Kristofferson (1976) and Brewster (1983), who used Kristofferson's procedure, usually obtained variances between 100 and 200 msec². (One of Brewster's three subject's variances generally exceeded 200 msec²). Under Hopkins' (1982) procedure, variances were lower still. Michaels (1977) reported higher variances, generally between 400 and 1000 msec², but his subjects were less practiced and his feedback stimuli differed substantially from those used in other work. As will be seen shortly, the short duration variances from Experiment 2 were about 200 msec². These are larger than most previous results, but they are still substantially smaller than the variances of Experiment 1.

It appears that there is one variance level for synchronization intervals from 150 to 550 msec, the exact value depending on procedural details, and a different level for intervals ranging from 1500 to 2000 msec. We might suppose that somewhere between 550 and 1500 msec there is a stepwise variance increase, just as steps were found in discrimination performance. If variances are constant across these two ranges, it seems unlikely that variability would increase smoothly between 550 and 1500 msec. A stepwise increase seems more plausible. It's not clear why there should be a step, however. Kristofferson's (1980) quantal counting hypothesis accounted for steps in discrimination performance by assuming a maximum count of 16 quanta, but this maximum does not apply to synchronization. We could entertain a similar notion for synchronization, with a different maximum count, but why should the maxima differ across tasks?

One hypothesis about the difference at short and at longer

intervals makes no reference to a quantal mechanism or to any fundamental internal limitations, Perhaps the difference reflects different expectations on the parts of subjects about how variable their responses "should be" at short and at longer durations. There is a common expectation that measurements of big things will be more variable than measurements of small things. In many psychology courses it is routinely stated that internal measures of big things, i.e. perceptions of large maghitude stimuli, are more variable than internal measures of small things. The subjects of synchronization (and many other long term) experiments mare rarely newcomer's to psychology, or "to this type of thinking. With particular reference to this experiment, different subjects volunteered the "explanation" for the obviously higher variances at long intervals that timing of longer intervals should be more variable than timing of shorter ones because the long intervals are longer (i.e. bigger) than the shorter ones, so of course the variances will be greater.

I will not attempt to detail a psychological model relating expectation and performance levels. I suppose that people have internal standards about what they consider to be acceptable levels of accuracy in their work, that those standards may vary across tasks, and in particular that larger synchronization errors might be considered more acceptable when the interval is long than when it is short. If so, asymptotic performance at longer intervals might be less accurate and more variable simply because the subjects are satisfied that this performance is fairly good, whereas at short intervals the subjects might try asbit harder to do better.

The subjects of Experiment 3 learned in Experiment 2 that small changes in the P1-P2 interval had no effect on their variance. This is what they saw in printouts of their own performance, for many sessions. In Experiment 3, I simply made further small changes in their P1-P2 intervals, reducing them in steps of 5 msec toward 100 msec and below. This should not have led subjects to expect and accept higher variances in the same way that increasing the interval in big steps to 1500 msec might.

From the subject's point of wiew, Experiment 3 involved a range of intervals from 85 to 200 msec. Given that the subjects cannot use P1 as a response trigger when the P1-P2 interval is very short, the actual range of <u>synchronization</u> intervals was probably closer to 150 -1150 msec, with a steep shift somewhere between P1-P2 intervals of 85 and 150 msec. But this manipulation was not obvious to the subjects. If variances are higher at long intervals because subjects expect long interval mariances to be higher, or due to other similar cognitive factors, then we should expect variances from 85 to 200 msec (that is, 150 to 1150 msec) to be constant. Alternatively, if some internal constraints limit accuracy and consistency as stimulus duration increases, we should see an increase in variance across this range, to a level comparable to that obtained from 6H or D6.

In sum, my plan for Experiment 3 was to study synchronization at long intervals that the subjects thought were short intervals, avoiding the effects of any biases that the subjects might have about how good performance should or can be at long intervals.

It would be inaccurate to say that the subjects were totally

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unaware that they relied more and more on WS as P1-P2 was decreased. They volunteered that they did this. They also complained that the task was getting more and more difficult as the interval was reduced. I commisserated with them, complimented them on their efforts, asked them to please rely on P1 as much as possible, rather than on WS, agreed that they might need more practice before variances at 100 msec matched those at 200, and asked them to keep trying. The subjects were aware that they relied somewhat on WS, and that the task was difficult, but they tried very hard, extremely hard, to obtain the same variances at 100 msec as they produced at 150 to 200 msec. On this basis, I think this was a successful manipulation of their expectations about variance levels, and I think that they took the idea that they might achieve very low variances at what were in fact long intervals much more seriously than did D6 and GH.

4.1 Method

4.1.1 Subjects

Four subjects started in this experiment: RW, JB, KC and KF. KC had just completed her first year as an undergraduate. RW was a male graduate student studying psychology. JB and KF were senior male undergraduates, interested in, but not majors in, psychology. These subjects were somewhat less familiar with the timing literature than the subjects of Experiments 1, 4 and 5. Excepting RW, I think they were almost completely unfamiliar with it. Excepting RW, they had participated in significantly fewer psychological experiments than the subjects of the other experiments. Only RW and JB completed the experiments. KF left the experiment shortly after he dropped out of the university. KC left the experiment at my request. After 76 sessions, her variances were still quite high, her response distributions were skewed, and there were many extreme responses. There was no apparent trend toward improvement. I reluctantly concluded that her data would not provide a useful basis for comparison with long auditory or short visual results, the point of Experiment 2. Further, since she was unable to achieve low variance levels at 200 msec, it appeared pointless to look for low wariances at shorter intervals, the goal of Experiment 3. In the context of these experiments, KC's data appeared uninterpretable, so I stopped collecting them.

4.1.2 Procedure

The general procedure was described in Chapter 2. Some further information on instructions in Experiment 3, has already been noted in this chapter. JB and KF started at 200 msec, while RW and KC started at a 150 msec P1-P2 interval. KC found 150 msec extremely short and difficult. I switched her interval to 200 msec after 9 sessions. She stayed at this interval until she left the experiment.

RW, JBrand KF scanned the range from 150 to 200 msec. I then started Experiment 3, reducing the P1-P2 interval to 100 msec, in steps of 5 msec. KF left during the scan down to 100 msec, leaving RW and JB in the experiment. RW completed the first part of these experiments much earlier than the other subjects and was available to participate in more sessions than the others. After a few sessions at 100 msec I brought his interval back to 150 msec to check that performance at such short intervals had no deleterious effects on performance at longer ones. There appeared to be no problem, and his interval was again reduced to 100 msec, in steps of 5 msec. Both subjects practiced extensively at 100 msec and less extensively at shorter intervals. After this, I brought the interval back to 150 msec, ending Experiment 3, and continued increasing it, to 350 msec, with extended training at 300 msec, as the final part of Experiment 2, and mainly, as preparation for Experiment 5, in which the subjects were to transfer to visual synchronization. (RW actually did transfer. JB, who had left the university by this point, decided to stop at the end of Experiment 2.)

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The exact schedules for all subjects, and various summary statistics for each session, are listed in Appendix 1. JB's and RW's schedules also appear visually in numerous figures below.

4.2 Results

As noted above, KC's results were poor. There is not much else to say about them, and they will not be discussed further. KF's performance was excellent but he left after only 82 sessions, during the early stages of the scan down to 100 msec. All appeared that his results would parallel those of RW, merhaps with lower variances. Unfortunately, there is not enough data for any firm conclusions, so his data will also be set aside. The tables in Appendix 1 list various statistics of each session's data for $a \neq 1$ subjects, including KF and RC. The reader who wishes to compare their performance to RW and JB's types of graphs and summary tables for these subjects as those presented below for RW and JB:

4.2.1 Latencies

Figures 7.1 (RW) and 7.2 (JB) show the relationship between synchronization latency (upper panel) and P1-P2 interval (lower panel) for each subject for each session. The relationship in each case is linear. Both subjects were typically able to produce mean latencies within 10 msec of the P1-P2 interval, even at the very short intervals.

4.2.2 Variances

Figures 8.1 and 8.2 show RW's and JB's variances for each session. The middle panels again show P1-P2 intervals. The lower panels show the mean foreperiod durations, which indicate which computer was used when. Performance was better when the experiments were controlled by the Apple, but this appears to be natural improvement with practice. There is no abrupt transition due to the change in the equipment.

Tables 4.1 (RW) and 4.2 (JB) give averaged variances, q estimates, shape statistics and correlation coefficients for each subject across the "practiced" intervals, the intervals at which the m subjects spent two or more consecutive sessions. As in Table 1, if the subjects spent more than 5 sessions at an interval, only the last 5 were included in averaging.4

The tables list the same types of statistics twice for each



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Upper panel: Mean synchronization latencies (relative to P1), in milliseconds, of each session.

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Lower panel: P1-P2 intervals (msec) of each session.

Figure 7.1: Subject RW. Figure 7.2: Subject JB.

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FIGURE 8

Upper panel: Response latency (from P1) variances, in msec², of each session. Note that there are two lines at the top of the panel. Variances shown as extending to the topmost line were too large to represent accurately on this scale. For exact values, see Appendix 1.

Middle panel: synchronization intervals, in milliseconds, of each session.

Lower panel: Mean foreperiod duration, in msec, of each session. The means were approximately 900 msec when the POP-8 controlled sessions and 1000 msec when the Apple controlled sessions. Gaps in the figure (ostensibly 800 msec means) are due to lost raw data, making calculation of the sessions' mean foreperiod durations impossible.

Figure 8.1: Subject RW. Figure 8.2: Subject JB.







SESSION: 30 60 90 120 150 180 210 240 270 300 330 360 390 RH AUDITORY





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TABLE 4

Mean within-session variances (msec²), q estimates (msec), and coefficients of symmetry (ROOT B1), Kurtosis (B2), and correlation between response latency and foreperiod duration. Only data from intervals at which the subject spend two or more consecutive sessions were averaged for this table. The averages are based on the Tast five sessions at the interval before the subject was transferred to the next, or, if there were fewer than five, on all sessions at the interval.4

Table 4.1: Subject RW. Table 4.2: Subject JB.

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SUMMARY STATISTICS FOR RW AUDITORY RESPONSE LATENCY MEASURED FROM P1

INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH FP
150	191	31.54	-Ò.46	4.38	0.05
175	163	28.77	0.15	3.86	0,11
200	240	35.91	0.36	4.07	0.11
200	224	34.55	0.36	3.58	0.11
100	483	52.42	-0.79	3.86	-0.22
150	147	27.03	-0.30	3.50	-0.07
100	255	37.18	-0.97	5.16	-0.22
95	363	45.00	-1.09	5.76	-0.37
90	336	43.22	-0.97	4.69	-0.40
85	409 -	48.00	-0.95	5.04	-0.34 -
150	142	26.44	0.06	4.26	0.05
200	188	31.31	0.19	3.01	0.11
250	364	45.11	0.16	2.97	0.06
300	241	36.01/	0.13	2.95	0.07
350	290	39.84	-0.01	3.05	0.03
300	205	32.87	0.05	3.00	6.06

SUMMARY STATISTICS FOR RW AUDITORY RESPONSE LATENCY MEASURED FROM WARNING STIMULUS

INTERVAL ·	VARIANCE	Q	ROOT_B1	B2	CORR WITH FP
150	301	40.67	0.26	4.71	0.61
175	288	39.73	0.85	5.50	0.66
200	361	44.88	0.61	3.74	0.59
200	339	43.44	0.61	3.80	0.59
100	479	52.21	-0.56	3.79	0.20
150	218	34.07	0.21	3.78	8.58
100	274	38.66	-0.51	4.16	0.35
95	322	42.23	-0.43	3.76	0.17
90	288	39.73	-0.46	3.73	,0.16
85	364	45.12	-0.82	4.94	0.12
150	257	37.32	0.81	5.80	0.66
200	309	41.31	0.52	3.47	0.63
250	478	52.11	0.26	3.05	0.49
300	358	,44.68	0.55	4.15	0.57
350	388	/ 46.69	0.36	3.92	0.50
300	306 /	41.08	0.39	3.51	0.58
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SUMMARY STATISTICS FOR JB AUDITORY RESPONSE LATENCY MEASURED FROM P1

INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH FP	-
200	409	48.01	0.06	3.55	-0.02	
175	341	43.56	0.14	5.84	-0.06	
.150	394	47.04	-0.73	5.99	-0.17	
108	> 295	40.25	-0.87	5.75	-0.15	
95	411	48.11	-1.09	5.60	-0.19	
150	144	26.71	-0.16	4.19	-0.03	
200	. 151	27.52	0.13	2.93	-0.03	
250	265	37.92	0.48	4.02	-0.01	
300	175	30.04	-0.11	3.45	-0.04	
350	210	33.34	-0.08	3.18	0.00	
300	200	32.40	-0.1i	4.05	-0.14	

SUMMARY STATISTICS FOR JB AUDITORY RESPONSE LATENCY MEASURED FROM WARNING STIMULUS

INTERVAL	VARIANCE	Q	ROOT B1	_B2	CORR WITH FP
200	494	53.07	0.36	3.88	0.41
175	409	47.98	0.42	5.80	0.41
150	421	48.74	_ -G_ 23	5.52	0.31
100	333	42.97	-0.41	5.04	0.37
95	422	48.84	-0.56	4.41	0.26
150	223	34.48	0.36	3.78	0.59
200	244	36.21	0.62	3.88	0.61
250	348	44.05	0.57	3.92	0.49
300	247	36.51	0.19	3.82	0.54
350	299	40.55	Ó.33	3.59	0.54
300	248	36.62	0,17	4.77	0.46

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group of sessions. The upper section of each table gives statistics computed on latencies measured from P1. The lower section of each table gives statistics computed on latencies measured from WS. The <u>distinction between latency from WS and latency from P1 is a tritical</u> one for comprehension of the rest of this chapter. Here is a numerical example. Suppose that in a given trial the foreperiod is 950 msec, the P1-P2 interval is 100 msec, and the response occurs 10 msec after P2. If we measure the latency from P1, it is 110 msec. If we measure the latency from the offset of the warning stimulus, it is 1060 msec. The difference between the two values is that the foreperiod duration (here, 950 msec) is added to the latency as measured from P1 (110 msec) to give us the latency from WS (1060 msec).

The foreperiod duration is random, geometrically distributed. Many of the statistics of the latencies should differ substantially depending on whether the subject is timing from WS or from PL. At this point, I will focus on the variances. The variance of the foreperiod is, theoretically and on average, 90 msec². The actual variances per session are listed in Appendix 1.

If the subject is timing from P1, his performance should be unaffected by the foreperiod duration and foreperiod variability should not show up in the variance of latencies measured from P1. Hopkins (1982) demonstrated this empirically for uniformly distributed foreperiods with a 33 msec² variance. If the subject is timing from P1 and we measure his latency from WS, we add an independent source of variance to his responses. On average, the variances

Now suppose that the subject is timing from WS. If the average foreperiod is 1000 msec and the P1-P2 interval is 100 msec, the subject waits for an average 1100 msec after WS before responding. The response is time-locked to WS, independent of the time of P1, which occurs too late to influence the response. As usual for synchronization latencies, the distribution of latencies as measured from WS should be symmetrical and highly peaked. Latencies as measured from P1 should not look like this however. If the subject times from WS and we measure latency from P1, we are subtracting a geometric variate (the foreperiod duration) from the waiting times generated by the subject. The latencies from P1 should, in this case, be more variable than those measured from WS, again by 90 msec².

These are not the only two ways that the subjects could perform the task, though these alternatives do describe the two response strategies obtained by Hopkins (1982) if we replace "timing from WS" with "timing from the initiation response" in his case. I will very briefly consider some alternative models later, after showing that in fact the subjects did not time exclusively from WS, ignoring P1, even when the P1-P2 interval was less than 100 msec. For now I will consider only these two alternatives. They are the clearest ones, I considered them the most likely, and I want to make sure that they are understood before rejecting them:

Table 5 briefly characterizes the distributions that we should obtain under these two strategies, depending on whether we measure latencies from WS or from P1. Supporting derivations are given in

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Appendix 2. Numerical values and formulae are presented in the text as

they are needed.

Table 54

Response Distributions if Timing is From WS or From P1

Latency Measured From:

The Subject Times from:

- WS WS & Low variance Symmetrical Low Kurtosis No correlation between latency and foreperiod duration
- P1 Higher variance Negatively skewed Higher kurtosis Negative correlation between latency and foreperiod duration

P1 Higher variance Positively skewed Higher kurtosis Positive correlation between latency and foreperiod duration

Low variance Symmetrical Low kurtosis No correlation between latency and foregeriod

4.2.2.1 Experiment 2 Variances

When P1-P2 intervals are 150 msec or longer, the variances of latencies measured from WS are substantially larger than those of latencies measured from P4. The differences are not exactly 90 msec², indicating some effect of the foreperiod even when the intervals are as long as 300 msec. RW's differences are larger than 90 msec², which we would expect if foreperiod duration and latency from P1 were positively correlated, which they are. JB's differences are smaller than 90 msec², reflecting a negative correlation between latency from P1 and foreperiod duration across the range. I'll return to the correlations later. They are consistent, and they are larger than those obtained from DG and GH, but they are not very large, 'especially in the final ascending series, which reflects the most practiced and best performance. For now, I will set aside these small correlations.

From 150 to 200 msec, at which both subjects practiced most extensively, final variances were generally between 150 and 200 msec². At the longer intervals , they were initially larger but with practice the variances of both subjects at 300 msec dropped to 200 msec². These are much higher than Hopkins' variances. They are slightly larger than those reported by Kristofferson (1976) and than the results from two of Brewster's (1983) three subjects. Kristofferson, Brewster and I all used 1000 msec foreperiods. Hopkins' foreperiods were an order of magnitude smaller & In simple reaction time, short foreperiods lead to faster and lower variance reactions than long foreperiods (see Niemi and Naatanen's, 1981 review). Perhaps this is at the root of the difference between Hopkins' results and the others, including the present ones. As to the difference between the present results and those of Brewster and Kristofferson, their foregeriods did not vary. The small foregeniod - latency correlations in RW and JB's data indicate that foreperiod variability has some effect on their performance and that they are not always simply timings from P1. It is impossible to tell whether Brewster's and Kristofferson's subjects always timed from P1. Since all stimulus events in their experiments were fully time-locked to each other, subtle shifts in strategy from trial to trial may have bed no impact on varianies. I suspect that the relatively high foreperiod variance

the present case is the main contributing factor to the difference in variance between RW, JB and the subjects of Kristofferson's and Brewster's research.

4.2.2.2 Experiment 3 Variances

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As the P1-P2 interval was reduced, variances increased and sizable negative correlations were obtained between response latency as measured from P1 and foreperiod duration, indicating an increasingly * important role for the warning stimulus. The correlations do not approach -1.0, which would indicate perfect and nonvariable timing from WS rather than from P1, but since there is timing variance the correlations should not be -1 when subjects time from WS. A formula and examples of expected values for the correlations will appear in the next section. For now, it suffices to say that the correlations are negative at short intervals, larger than those obtained at slightly longer intervals, and that this is the general pattern we should expect if WS eventually takes over as the stimulus controlling behavior as P1-P2 shrinks.

Examining the variances, RW's and JB's variances at 100 msec are actually smaller when latencies are measured from P1 than when they are measured from WS. The difference is not very large but it is in a surprising direction. It suggests that the response latencies are more heavily determined by the time of P1 than by the time of WS even when P1-P2 is a mere 100 msec.

However we measure the latencies at 100 msec, JB's and RW's variances declined to about 300 msec² after much practice. This is
about 200 msec² better than DG's performance. Variances at P1-P2 intervals of 100 msec, which <u>should be</u> variances at <u>synchronization</u> intervals of 1100 msec, are substantially lower than those obtained at 1500 msec. RW's variances are comparable to his performance at 300

Both subjects worked with shorter intervals than 100 msec, though they did not practice at these intervals as much as they did at 100 msec. At 90 msec, RW's variances as measured from P1 finally took on larger values than his variances measured from WS, suggesting that the time since WS was a more important determinant of his response latencies than the time of P1. Even at 85 msec, though, his variances were smaller than those obtained in Experiment 1. Whether we measure latencies from P1 or from WS, the variances are definitely greater than those he produced at longer intervals.

If the subjects were timing from WS and were ignoring P1, as did Hopkins' visual synchronization subjects, these low variances would indicate that at synchronization intervals as long as 1100 msec, variances even lower than DG's at 1400 and 1500 msec can be obtained. The variances are larger than those obtained in Experiment 2. This could be interpreted as suggesting that variances at 1100 msec are intermediate to those we should expect at 150 - 550 msec and at 1500 -2000 msec, or it could be interpreted as suggesting that the variances obtained at 1500 msec. from DG and GH could be improved upon if they eracticed further or if their expectations in the task were changed. However, it appears that the subjects did not base their responses exclusively on the time since WS, even when the P1-P2 interval was as

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short as 85 msec. Presumably, some of the responses were reactions to P1 and some were based on the time since WS. If simple reactions are less variable than long interval synchronizations, the intermediate variance obtained might be due to a mixture of a high variance and a lower variance distribution. Under that view, it could be argued that internal timing variance at 1100 msec might indeed be the same as that at 1500 - 2000 msec. I will return to this notion, and to prixture hypotheses, below.

4.3 Correlations Between Response Latency and Foreperiod Duration

Figures 9.1 and 9.2 present RW's and JB's correlations for each session. The upper panels show the correlation between foreperiod duration and response latency as measured from the warning stimulus. The middle panels show the correlation between foreperiod duration and latency as measured from P1. The bottom panel again shows the P1-P2 interval.

If the subject bases his response exclusively on the time since. P1, and if we measure response latency since P1, or if he bases his response exclusively from the time since WS, and we measure latencies from WS, then the correlation between latency and foreperiod duration should be zero. What should the correlations be if the subject times from P1 and we measure latency from WS or if the subject times from WS and we measure the latency from P1?

A brief derivation of the population correlation values is given in Appendix 2. The magnitudes are the same in both cases, and they depend on the relative variance of the timing distribution

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FIGURE 9

Upper panel: Correlation between response latency (from WS) and foreperiod duration for each session. Correlations extending to the topmost line of the figure were too large to represent accurately at this scale. For exact values, see Appendix 1.

Middle panel: Correlation between response latency (from P1) and foreperiod duration for each session.

Some sessions' raw data were lost and correlations could not be calculated. No values are plotted for these sessions. They are indicated by gape in the figures (sometimes most easily seen as breaks in the top lines of the figures.)

Lower panel: synchronization intervals, in milliseconds, of each session.

Figure 9.1: Subject RW. Figure 9.2: Subject JB.





(unknown) and the foreperiod distribution (90 msec²). Let C be the population correlation coefficient and let VT and VF be the timing (including motor variance) and foreperiod variances respectively. The magnitude of the correlation is given by:

Absolute Value of C =
$$\sqrt{\frac{VF}{VT+VF}}$$

If the subject is timing from WS and we measure latency from P1, C is negative. If the subject times from P1 and we measure latency from WS, C is positive. Table 6 presents some sample correlation magnitudes, for comparison with the statistics reported in %ables 4.1 and 4.2.

Table 6

Expected Correlations as a Function of Timing Variance

(f Variance Correlation (Absolute Value)

100	nsec≥		•1	.688
125	mset ²			.647
150	msec ²			.612
175	msec ²	•		.583
200	msec ²			.557
225	msec ²			.535
250	msec≥			.514
300	msec²			.480
500	mse,c ² ,	Ð		.391
750	msec2			.327

Consider first the data of Experiment 2, i.e. the results from -150 to 350 msec. In RW's case, the correlations between latency (from P1) and the foreperiod are small but positive. The correlations, between latency measured from WS and foreperiod duration are quite large, generally larger than .50 and often larger than .60 (see Table 4). While there seems to be some effect of the foreperiod on the

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latencies, the responses are mainly determined by the time of P1.

In JB's case, there are negative correlations between latency from P1 and foreperiod duration across the range. (I should note that the final five session average correlation at 300 msec, of -.14, is stronger than the value obtained during most sessions at 300 msec. See Appendix 1's listings.) If we examine correlations between foreperiod duration and latency measured from P1, from 150 to 350 msec these are typically large and positive, always exceeding .40 and usually' exceeding .50. They are smaller than the correlations obtained from RW and they are smaller than we would expect if JB timed exclusively from P1, since his response latency variances are generally around 200 msec² or less. There is some indication of a consistent effect of the foreperiod duration on latency, but it is not a very sizeable effect.

Now let us examine the data in the 95 to 100 msec P1-P2 interval range. The variances are larger here, but from Table 6 we should still expect to find correlations of at least -.45 between foreperiod duration and latency measured from P1 and and we should expect to find negligible correlations between the foreperiod and interview measured from WS. That is, this is what we should expect to find if the subject times exclusively from WS. These expectations are the mirror image of those of Experiment 2. We know from Experiment 2 that such correlation magnitudes are not merely hypothetical. The right correlations were obtained, at least approximately, for intervals Tonger than 150 msec. Examining the Experiment 3 results, the correlations between latency (from WS) and foreperiod duration do not 1 drop to zero. Nor do they drop to levels comparable to the correlations between foreperiod and latency from P1 obtained in Experiment 2. There are consistent positive correlations between foreperiod and latency (from WS) even when P1-P2 is 85 msec. At an interval of 100 msec, the correlations are fairly large, .35 for RW and .37 for JB: The correlations between foreperiod duration and latency from P1 are negative but they are too weak. RW's values reached -.40, but JB's didn't even reach -.20.

The correlations between latency from P1 and foreperiod duration are too weak and those between latency from WS and the foreperiod are too strong for me to accept the hypothesis that when the P1-P2 interval is very short the subject will time an interval from WS and will respond exclusively on that basis. The time of P1 appears to play an important role in the subject's behavior, even when the P1-P2 interval is as short as 85 msec.

It seems that to describe this behavior we need a model that gives both WS and P1 a role in determining the time of the subject's response. There are models of this class for simple reaction time, developed because subjects' reaction times are correlated with the duration of the foreperiod preceding presentation of the stimulus. Niemi and Naatanen (1981) review that very extensive literature in detail. One line of work within that literature led to the development of the synchronization procedure. Rather than instructing their subjects to respond as soon as they perceived a stimulus, some experimenters instructed their subjects to respond at some time, or within some narrow range of times, after that stimulus. This line of

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work includes research by Snodgrass (1969, Snodgrass et. al., 1967), Naatanen et. al. (1974), and Saslow (1974). Kristofferson (1976) reviewed the relationship between synchronization as a procedure for studying timing and synchronization as a procedure for studying reaction time. Further discussion can be found in Hopkins and Kristofferson (1980) and Hopkins (1982).

I am reluctant to venture into that literature in any great detail here. Detailed consideration of the microstructure of reaction time <u>per se</u> is beyond the scope of the present work (and of the main stream of my research interests). Extensive further empirical work would be required in the development of a satisfactory model for data such as these: there is not an appropriate model extant in the RT literature. As an example of the difficulties, and of what appeared to be a very promising idea, consider the following approach, suggested by Kornblum (1973; see also Ohlman and Billington, 1972).

In a reaction time task, the subject's goal is to respond as soon as possible after the presentation of the reaction stimulus (analogous to P1 when the P1-P2 interval is extremely short). If he waits until he perceives P1 before responding, he will respond some time after P1, because of afferent and efferent/defays. Alternatively, he could use the warning stimulus. (Niemi and Naatanen, 1981, make an effective argument that there is always some stimulus in RT situations that the subject can use as a warning stimulus. Usually, the experimenter provides an explicit one, to prepare the subject to respond.) If the subject knows that the reaction stimulus will occur 1000 msec after WS, he can time and interval from WS and at some point

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after WS offset he can initiate a response which should be completed just as P1 is presented. If he could time the required interval perfectly, and if there is no variance in the foreperiod, he could produce any apparent reaction time desired. Subjects' timing is not variance free, however, and foreperiods are often variable. If the subject tries to appear to have too short a reaction time, his responses will sometimes precede the reaction stimulus, revealing the strategy. With practice and feedback from the experimenter that discourages early responding, the experienced subject may refine the strategy, trying to respond a bit later, at a point when the response can always or almost always be expected to occur after P1, but not much _ after P1. For example, rather than trying to respond 1000 msec after WS, if P1 never follows WS by more than 1050 msec, the subject might try to resignd 1050 msec after WS even though this would yield some late reactions if P1 occurs much earlier than this. Now suppose, that the subject uses this strategy but on one trial, while waiting out the 1050 msec, P1 is presented 900 msec after WS. There's no point waiting another 150 msec# The subject should respond immediately instead.

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Under Kornblum's analysis, if the foreperiod is variable, we should expect two types of response triggers. The subject generates a criterion interval internally, initiating the response at the end of this interval if he has not initiated the response already. If P1 is perceived before the internal interval ends, he responds immediately to that stimulus instead.

Kornbl 's hypothesis is quite similar, to Kristofferson's

(1977) theory of discrimination performance. In that case, the subject generates a criterion interval and decides whether a stimulus interval is long or short on the basis of which ends first, the criterion or the stimulus. The criterion can be studied by asking the subject to respond as soon as he can. Kristofferson found that 'long' responses (arising when the criterion ends before the stimulus) were time-locked to the stimulus initiating the interval (analogous to WS). "Short" responses were time-locked to the offset of the stimulus interval (analogous to reactions to P1). In the reaction time situation, the subject is not required to make a discriminatory response, but he is required to speed the response that he does make and under Kornblum's model the basis of the response is the same as in the discrimination setting. The subject responds as soon as the criterion interval ends or as soon as he perceives the end of the interval between WS and the reaction stimulus (by detecting P1), whichever comes first.

If the subject does the task this way, we should still see a negative correlation between response latency and foreperiod duration. Kornblum's model was explicitly designed, as was Ollman and Billington's (1972), to explain this correlation. If the foreperiod is long, P1 will be delayed relative to the end of WS. The subject's response, based on the time since WS, will be early relative to the time of P1. If the foreperiod is short, the subject will respond to P1 itself. Since the subject will always respond to P1 if he perceives it before the criterion interval ends, his responses to P1 should never have a shorter latency (measured from P1) than his responses based on the end of the criterion interval. There may be a few exceptions due

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to motor, and timing variability but in general responses triggered by P1 should occur later, relative to P1, than responses triggered by a criterion interval following WS. The longer the foreperiod the earlier the responses triggered by this interval will be, relative to P1.

It is not easy to predict the exact magnitude of the megative correlation. When P1 occurs before the end of the criterion interval, the response time to it should be approximately constant. Across the range of foreperiods over which P1 will occur before the end of the interval, response latency and foreperiod duration, should be uncorrelated. Combining these latencies with the negatively correlated latencies time-locked to WS, we should obtain a smaller correlation than would be obtained if all of the subject responses were time-locked to WS. The exact correlation will depend on the variance and range of the foreperiod, the variance of the subject's criterion, and the mean of the criterion. Whatever the exact value is, however, it should never be positive.

Kornblum's hypothesis appears plausible as an explanation of JB's and RW's relatively low correlations between foreperiod duration and latency from P1 for very short intervals. But from 140 msec upwards, RW's correlations are neither negative nor zero. They are positive. Is a different mechanism responsible for RW's small positive correlations at longer intervals than that responsible for his small negative ones at shorter intervals? Perhaps. But if so, and if Kornblum's model describes the mechanism underlying the negative correlations, this isoonly half of the story for these data. Given that RW's correlations appear to vary continuously as the P1-P2

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interval is changed (see Figure 9.1) a complete model would in this case require specification of a slow transition from one mechanism to another that induces positive correlations, and would include a range of P1-P2 intervals over which both mechanisms or strategies were operative in the same sessions.

A further difficulty posed by the present data that is not escaped by Kornblum's model lies with the actual reaction times obtained. Suppose that some of the subject's responses are in fact simple reaction responses to P1. These should take a long time relative to an 85 msec P1-P2 interval. Yet RW's longest mean response time when the interval was 85 msec was 91 msec, and means shorter than 85 msec, such as 82 msec, were also obtained. None of RW's latencies (from P1) was longer than 140 msec when P1-P2 was 85 msec, and exceedingly few were longer than 120 msec. Most were shorter than 100 msec. According to the standard summaries, reaction time varies with stimulus intensity and to reliably obtain reaction times as short as 🌤 105 msec you should present stimuli on the order of 110 dB. These are extremely intense. In terms of sound pressure, they are more than 100 times as intense as the 68 dB stimuli used in this experiment. Reaction time to 68 dB stimuli has been estimated at 150 msec for practiced subjects. How can RW be emitting simple readtions to a 68 dB P1 this quickly? (And JB's performance at 95 msec is not much less striking.) I don't know. Perhaps such short reaction times can be obtained with moderately intense stimuli under optimal conditions and perhaps these were optimal conditions. Certainly, synchronization yields much lower variances than those obtained in any other type of

reaction time task (see Kristofferson, 1976; Hopkins and Kristofferson, 1980). Perhaps this procedure can elicit faster responses as well. That conclusion is certainly suggested by these data, though I am not convinced that it is forced by them.

4.2.4 Distributional Shape

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Figures 10 show RW's and JB's symmetry coefficients for each session. Figures 11 show the Kurtosis coefficients. The upper panels are based on latencies measured from the warning stimulus. The middle panels are based on latencies from P1. The lower panels show the correlations between latency (from P1) and foreperiod duration. The coefficients should vary as a function of this correlation. In Appendix 2, I derive expressions for population symmetry and Kurtosis coefficients for convolutions of Jogistic and triangular distributions and for convolutions of logistic, triangular and geometric distributions. The latter convolution, of the three distributions, should describe performance if the subject times from WS, with the usual triangular + logistic latency distribution assumed for synchronization latencies, and we measure latency from P1 rather than from WS. Equivalently (except that the geometric foreperiod distribution was subtracted in the convolution above and will be added in this one), it should describe the latency distribution as measured from WS if the subject is timing from P1. The coefficients all depend on the relative variances of the logistic, triangular and geometric variates convoluted. Some sample values are calculated in the appendix. Having rejected this simple model of the subject's behavior.

FIGURE 10

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Upper panel: Symmetry coefficients of the response latency (from WS) distributions of each session.

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Middle panel: Symmetry coefficients of the response latency (from P1) distributions of each session.

Some of the symmetry coefficients were too large in absolute magnitude to represent accurately in these figures. Where there are two lines at the top or bottom of a figure, and a symmetry value extends to the outermost line, the actual value should be shown as extending further. For exact values, see Appendix 1.

Some sessions' raw data were lost and symmetry coefficients could not be calculated. No values are plotted for these sessions, which are indicated by gaps in the figures.

Lower panel: Correlation between response latency (from P1) and foreperiod duration. Repeats the middle panel of Figure 9.

Figure 10.1: Subject RW. Figure 10.2: Subject JB.



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FIGURE 11

Upper panel: Kurtosis coefficients of the response latency (from WS) distributions of each session.

Middle panel: Kurtosis coefficients of the response latency (from P1) Addistributions of each session.

-Some of the Kurtosis coefficients were to large to represent accurately in these figures. Kurtosis values shown as extending to the topmost line of a figure are larger than they appear. For exact values, see Appendix 1.

Some sessions' raw data were lost and kurtosis coefficients could not be calculated. No values are plotted for these sessions, which are indicated by gaps in the figures.

Lower panel: Correlation between response latency (from P1) and foreperiod duration. Repeats the middle panel of Figure 9.

Figure 11.1: Subject RW. Figure 11.2: Subject JB.



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I will not detail those results here. The graphs, however, show up a few aspects of the data that should be mentioned.

Consider first the data of Experiment 2, from 150 to 350 msec. The distributions of both subjects' latencies do not appear symmetric from these figures. Early in practice, RW₂s latencies (from P1) were clearly negatively skewed, and substantially so. After much practice, this relationship changed sign. JB's symmetry coefficients change sign more frequently. In both cases, the distributions seem more skewed than those obtained by Hopkins (1982) and than those obtained in Experiment 1. There is some tendency for the skewness coefficients to follow the correlation coefficients, in sign and in magnitude, and this makes sense. If latencies are positively correlated with foreperiod duration, and the foreperiod duration is positively skewed, the latency distribution should be positively skewed. If the correlation is negative, long foreperiods translate into short latencies, so the latency skew should reverse. Whatever model explains (the correlation between latency and foreperiod duration should have no difficulty with these asymmetries in the data.

The Kurtosis coefficients of both subjects were also a bit large in Experiment 2, but this may again be due to foreperiod effects. RW's Kurtosis averages in the 200 to 300 msec range were close to 3 (see Table 4), which matches other synchronization results nicely. JB's averages were higher but from Figure 11.2 (or from Appendix 1) we can see that kurtoses near or smaller than 3 were not unusual in his data.

The Experiment 3 symmetry coefficients (85 to 150 msec

intervals) are more curious. They are not due to computational errors in the computer program, which I rechecked after examining the graphs. When (as in Experiment 2) the correlation between response latency (from P1) and foreperiod duration is positive, zero, or only slightly below zero, we should expect foreperiod duration and latency from WS to be strongly positively correlated. Since the foreperiod distribution is positively skewed, we should expect the latency (from WS) distribution to be positively skewed and it is. The curpus finding is what happens when the correlation between latency (from P1) and foreperiod duration becomes quite negative. If the subject is timing from WS then his latency distribution, measured from WS, should be symmetrical. The correlations (foreperiod and latency from P1) are too weak to support the hypothesis that he is timing from WS exclusively. ÷. so we should not expect perfect symmetry. The correlations between foreperiod duration and latency from WS are still positive, though less large, so we should find that the latency distributions (from WS) are positively skewed, though less skewed than at the longer interval. But they are not positively skewed, nor symmetrical. Instead they are very negatively skewed. I do not know why these data look this way. I think this will prove to be a very difficult problem and it may be the critical feature of the data for discriminating between alternative hypotheses designed to describe foreperiod-latency correlations.

4.3 Concluding Comments

The variance levels in Experiment 2 were higher than those typically obtained in synchronization experiments, probably because of

the long and variable foreperiod. This will pose some difficulties in interpreting the visual synchronization results, though given those data, the difficulties will not be as great as they could be. The problem is whether we should compare the visual variances with these data or with variances from the rest of the literature. For future reference I will note that RW's best variances were 123 msec² (twice) and JB's were 113 and 125 msec². (KF's were 156 and 160.)

The correlations between latency and foreperiod across the 85 to 350 msec range appear to be reflected in all aspects of the latency distributions' shapes (variance, symmetry, Kurtosis) but not always in easily predictable ways.

These data raise more questions than they answer. In particular, is minimum response time for moderately loud stimuli subtantially shorter than is currently believed? Should we still accept 100 to 105 msec as an absolute minimum on reaction time, given that reaction time to these less than maximally intense stimuli may be at this level or shorter? Finally, what are the contributing processes or strategies that yield negative, zero, or positive correlations between latency and foreperiod, depending on P1-P2 interval? Unambiguous answers to these will require much further work and thought but they are important questions for those interested in determinants of reaction time.

Estimates of q in Experiment 3 were generally between 40 and 50, however the latencies were measured. These are smaller values than those obtained with long auditory intervals and larger than those obtained with short intervals.

Experiment 3 involved a trick and a gamble. The trick lay in manipulation of the subjects' conscious attitudes. If they could be convinced that their synchronization variances should be at very low levels, at levels they knew they could attain under conditions they were led to consider similar, then perhaps they could achieve those levels. The gamble was that the subjects would not be so resourcefyl that they would develop uninterpretably complex strategies for achieving this.

The trick worked: the subjects tried very hard to attain low variances and their performance at P1-P2 intervals of 10D msec is impressive.

The gamble failed. I opened the discussion of possible results for this experiment with three alternatives: subjects might time these long WS-P1 intervals at the same q level as they used with short but timeable P1-P2 intervals (such as 200 msec), or their internal timing variability might be intermediate to the levels seen at the short and long intervals, or variability might rise to the same level as found in Experiment 1, suggesting a stepwise increase between 550 and 1100 msec, with constancy from 1100 (or so) msec out to 1750 or 2000 msec. None of these hypotheses is incompatible with the present data.

Suppose first that the subjects in fact operated at a q level of 25 or 30, i.e. at the level typical of short interval auditory performance. If this is true then the measured latency variances include more than just internal timing variance and a low level of motor variance, i.e. for our purposes the measured variances are inflated. The measured variances almost certainly are inflated. Here

are five extra sources of variance that I consider likely, and there are probably more:

(1) The subjects sometimes timed from WS and sometimes timed from or reacted to P1. If we measure latency from WS and the subject times from P1, or if we measure latency from P1 and the subject times from WS, we add the variance of the foreperiod to the variance of the response latency itself in estimating latency variance. The degree of inflation of our estimate depends on how often the subject timed from the stimulus we assume he ignored.

(2) There must be some lower limit on mean reaction time and the assumption that it is as low as 100 msec at 68 dB flies in the face of much reaction time data. If it is not this low, then mean latency (relative to P1, at a P1-P2 interval of 100 msec) when the subject is timing from WS should be different from mean latency when the subject is reacting to P1. When the subject is timing from WS, he can typically achieve a 100 msec latency (relative to P1). When reacting to P1, he typically cannot. Distributions of a mixture of both types of responses should be bimodal, with high variance, even if the variances of the component distributions (timing from WS, reactions to P1) were identical and low. The degree of variance inflation should depend mainly on the separation between the two means. The expectation of bimodality is a natural outgrowth of analysis of the task demands along the lines suggested by Kornblum. I cannot say that this expectation is forced by that line of thought, but it is highly compatible with it. I have examined the histograms of the response distributions and was surprised that this is not apparent in them.

There are long tails in the distributions, but they appear unimodal. Still, this might be masked by the one discussed next.

(3) The interval between P1 and P2 is_an "empty" interval. The interval between WS and P2 is a "divided" or "filled" interval: P1 is interpolated between these stimuli. The presence of an interpolated stimulus typically affects the judged duration of that interval. A divided interval is judged longer than an empty one of the same physical duration, the simplest example of the filled duration illusion (see Fraisse, 1963, for a review). A divided interval in which the stimulus occurs early is generally judged longer than an equally long one in which the interval occurs late (eg. Buffardi, 1971; Israeli, 1930). Suppose that the subject is timing a synchronization response from WS and that he is subject to this illusion, due to the presence of P1.7 The later that P1 occurs, relative to WS, the shorter the perceived interval between WS and P2. If the subject is attempting to produce an interval of 1100 msec, to coincide with P2, but his perception of the time elapsed to this point has been influenced by the time of presentation of P1, he may respond at the wrong time. If P1 occurs early, he may overestimate the time elapsed so far, responding early. If P1 occurs late, he may underestimate elapsed time and respond late. Brewster (1983) used a three stimulus synchronization task, with a P3 interpolated between P1 and P2 and did find that synchronization latency (responses were to be synchronous with P2) was positively correlated with the interval between P1 and P3. She did not interpret her result in terms of the filled duration illusion but consider ion of the various possible explanations would take us far

afield. If the subject is timing from WS and his latency varies as a function of the time of P1 (because of an illusion, because P1 interrupts processing of the interval, for whatever reason), then the response latency distribution is a mixture of many distributions (one for each WS-P1 interval) with many means, and even if they all had the same variance we would not obtain a measure of this common variance but of a larger one, the variance of the mixture.

(4) Kristofferson (1976) found serial dependencies in one of his subject's response distributions. Successive trials' latencies were positively correlated. Hopkins (1982) also found this but the correlations diminished to negligible values with practice. I have not looked for sequential dependence in the data -- this is a common source of noise in many paradigms (see, eg. Ward's 1979 review of the scaling literature), depending on the specifics of the procedures involved -but it would not be surprising if it was unusually (for synchronization) strong when P1-P2 intervals are almost impossibly short.

(5) The very short P1-P2 interval task was more stressful than those involving longer intervals. The subjects sometimes reported that they felt "typer" after a session. I informally noted unusual palmar sweating after sessions from one subject. The other sometimes jokingly noted that he performed best if he had a few beers first (presumably, to relax) and I think he was right. Startle responses, hand twitches and so forth are probably more likely to arise in stressful situations and these may have played a role in increasing the apparent variance of the subjects, though most of these were removed from the data via the

truncation algorithm.

It is hardly inconceivable that these sources of extra variance could jointly contribute 100 to 300 msec² to the measured latency variances. Accordingly, it seems quite plausible that the subjects were operating at a q level of 25 or 30 (timing variances of 100 to 150 msec²) despite the much higher latency variances.

The second hypothesis, that internal variances are higher than those at short P1-P2 intervals (150-550 msec) and smaller than at long ones, is certainly concordant with the results. The latency variances measured were intermediate so the underlying variances may well have been.

The third hypothesis is that the subjects, when timing from WS, operated at the same q level (50 or so) as they would operate at in the 1500 to 1750 msec P1-P2 region. Suppose that the subjects sometimes timed from WS, with a q of 50 (variance of about 417 msec2) and other times they reacted to P1. Suppose further that their fast responses to P1 were less variable than their timed responses to WS, and that their mean reaction latency was not much different from the mean latency (relative to P1) of responses from WS, so the variance is not inflated due to bimodality of the mixture. In this case "the variance of the mixed distribution would be between the variances of the two component distributions, the exact value depending on the relative number of responses of each type. (The problem of added foreperiod variance on some trials still remains, but this cannot add more than 90 msec² (the variance of the foreperiod) and it probably adds less since it is not a factor in all trials.)

I have some theoretical difficulties with this hypothesis. In particular, q is conceptualized in terms of waiting times in the nervous system. Information about a stimulus is delayed by up to q msec before it can be passed to another stage of processing. How can q be small when the subject reacts to Pi but large when he times from WS? Especially if we follow Kornblum's reasoning and suppose that the subject is timing from WS on all trials, reacting to Pi only when it arrives earlier than expected. This is a remarkably fast switch of q levels, within the same trial.

At the empirical level, Kristofferson (1977, 1983) found that something very much like this occured in speeded duration discrimination. When the subject responded "long" (the internal criterion interval, corresponding to the time since WS, ends before the stimulus marking the end of the interval), his latency variance depended on the stimulus interval, increasing in a stepwise manner, with obtained q values ranging from 25 to 200 msec. When the subject responded "short", as a reaction to the marker ending the stimulus interval, his variance was about 280 msec², and this was independent of the duration of the stimulus interval.

Given the empirical demonstration, the theoretical objection that the subject's variance in reacting to P1 should not be lower than the variance of his response from WS has no force.[®] Accordingly, while this imposes its own theoretical difficulties, it is not implausible to suppose that the subject timed from WS at a q level near 50, the lower latency variances being due to a mixture of responses determined this way with less variable responses triggered by P1.

I find it impossible to discriminate between the three alternative views of this experiment's data because of the complexity of the subjects' response strategies. They evidently neither timed exclusively from WS nor exclusively reacted to on timed from P1 nor in the way suggested by nor exclusively reacted to on timed from P1 nor in the way suggested by Kornblum. (The negative skew of the distributions from WS, noted above, seems just as difficult for this hypothesis as for the others.) I am impressed by how well the subjects performed in this task, but I am a complete loss as to how. exactly, they did it.

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5. Short Visual Intervals: Experiments 4 and 5

These two experiments used visual P1 and P2 stimuli. Experiment 4 was an extended study of performance across the range from 275 to 550 msec. Experiment 5 was a much briefer transfer experiment, examining the performance of one subject who had participated in Experiments 2 and 3.

There are fewer, long term and moderately long term visual timing experiments than auditory experiments. The only visual synchronization data come from Hopkins (1982). In that experiment Hopkins showed that the subjects' behavior was not controlled by the visual stimuli, so, in effect, there are no visual synchronization data. There are visual discrimination experiments, but none as extensive as Kristofferson's (1973, 1977, 1980, 1983) auditory experiments. None of the visual discrimination experiments involved speeded responding, which yields response latency distributions, and so there are no tests of the real time criterion theory for visually bounded intervals. There are suggestions of steps in variance with duration, but no orderly pattern has emerged from visual data. Again, the necessary proctracted studies of single subjects' data have not been conducted. 'The evidence for a continuously adjustable source of internal delays is also fairly weak for visual stimuli. From single stimulus studies, which involve few different test stimulus durations per base duration, one can roughly estimate the shape of the internal distribution, and the mean of the distribution. The distributions

appear symmetrical with means quite close to the midpoint of the single stimulus series (eg. Allan, et. al., 1971), which suggests that fine tuning of the type found in audition probably occurs in visual timing as well. The stronger tests of continuity, long term production tests across a number of different, but not <u>very</u> different, intervals, have not been conducted.

My goal in Experiment 4 was to demonstate that response distributions in the visual synchronization situation are the same as those obtained when auditory-stimuli are used. If enough similarities. (and few enough dissimilarities) between visual and auditory performance can be found, it would seem logical to suppose that the same mechanism underlies performance in both modalities.

A vast number of comparisons between visual and auditory r performance could probably be made. I chose to examine the following aspects of the data:

A. Reliance on P1: The experiment fails if subjects rely on the warning stimulus for timing rather than on P1. Hopkins' subjects did not use the visual P1 as the stimulus marking the start of their synchronization intervals. Instead, they used the tactile feedback available from their trial initiation responses. It hasn't yet been demonstrated that subjects can synchronize to visual stimuli. While we should not be surprised to find that subjects can time out intervals using visual P1 and P2 stimuli, given that Hopkins' subjects didn't do this, this aspect of performance must be examined. It is unfortunate that the present procedure appears to induce some reliance on the

warning stimulus, but this <u>was</u> found in both Experiments 1 and 2, so we should probably find a small correlation between response latency (from P1) and foreperiod duration in this experiment as well. The point is that this correlation should be small. If is isn't, we are studying timing from an auditory warning stimulus or from a tactile trial initiation response, which is not what we want here. I will predicate the rest of this discussion on the (in fact, correct) assumption that the correlations will be small.

B. Variance Levels: It is difficult to settle on precise criteria for absolute variance levels for a number of reasons. First, there are individual differences, as seen in the data of Experiments 1, 2 and 3 already. Brewster (1983), Hopkins (1982), Kristofferson (1976) and Michaels (1977) also obtained different results from different subjects. Further, the difference across their experiments, apparently due to differences in procedure, spans more than an order of magnitude. Variances in Experiment 2 at the most practiced intervals were generally between 100 and 200 msec². Most of Kristofferson's and Brewsters' subjects typically operated at 100 to 150 msec² levels. One of Brewster's subjects' variances were generally just above 200 msec².

<u>C. Estimates of g:</u> These should settle to levels close to 25, 50 or 100 msec after practice. Again, we should not be surprised by individual differences.

<u>D. Symmetry:</u> Distributions should appear symmetrical. From Hopkins' data and the results of Experiment 2 we should expect that if there is a slight degree of asymmetry, the skew should be positive.

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<u>E. Kurtosis:</u> Kurtosis coefficients should be between 2.4 and 4.2, though, given the variance and positive skew of Kurtosis statistics, occasional higher values are not unreasonable. From Experiments 1 and 2 we should expect that kurtosis coefficients will generally be slightly higher than 3.

F. Stability of Performance Across a Range: Ideally, the distributions will be the same in variance, symmetry, kurtosis, and correlation with the foreperiod, across the 275 to 500 msec range. We should not expect this to be true of the performance of all subjects. One of Kristofferson's (1976) subjects' variances declined as the P1-P2 interval increased from 150 to 300 msec, staying constant only from 300 to 550 msec. A different subject's variances did not increase at the shorter intervals. Individual differences are also evident in Hopkins' * and Brewster's data. In Experiment 1, DG's variances were roughly constant from 1500 to 2000 msec whereas GH's variances appeared constant only from 1500 to 1750 msec, increasing beyond this. In Experiment 2, variances at intervals longer than 200 msec were generally higher than at shorter intervals, again with differences between the subjects. There are also individual differences in visual discrimination data (Allan et. al., 1971; Allan and Kristofferson, 1974). (The comparable auditory research all involved a single very long term subject, so comparisons across subjects are not available.) The basic phenomenon demonstrated in all of these experiments is that there is constancy of performance across some range of intervals that is wide enough to be called a range. That much we should expect from all strects. We can hope, but cannot expect on the basis of previous

results, that all subjects will show constancy across the entire range.

<u>G. Continuous Adjustability</u>: The finding that response distributions are the same (except for mean latency) at widely separated intervals would not in itself imply that distributions would not vary with small changes in the interval. The simplest quantal counting models very definitely would not predict constancy across small changes (smaller than one quantal unit) in the interval. It was on this basis that Stroud's model was rejected in the introduction. Practiced performance was examined at a number of intervals differing by only 5 msec. We should expect mean latencies to track these changes, but variances and coefficients of skew, Kurtosis and correlation should be unaffected by them.

<u>H. Practice:</u> In auditory synchronization (eg. Experiments 1 and 2; Kristofferson, 1976) and in speeded discrimination (Kristofferson, 1983), variances at intervals at which there was extensive previous practice are generally lower after a scan across a range of other intervals than they were at the end of a long initial practice series. In Experiment 1, variances did not appear constant, or even roughly constant, until the final scan back to the initial interval. During the scan toward longer (GH) or shorter (DG) intervals than the original interval, variances increased with increasing stimulus duration. Kristofferson (1983) found that response latency variances in a speeded discrimination task initially increased with the separation of the stimuli, eventually settling at low and constant values only after two full scans (up and down twice) of the range. It is not clear why scanning should have such a beneficial result on

performance. I find it hard to believe that this is simply a general practice effect, resulting for example, in a decline of motor variability. After 50 sessions' initial practice at an interval the 'task is rather well learned already and motor variability should already have dropped to a rather low level. Whatever the basis of this phenomenon, it is quite noticeable in a fair bit of auditory data. It should be evident in the visual data as well.

These expectations for the visual results are not as precise as I would like them to be. It is fairly clear what "perfect" performance would entail: variances in the 100 to 150 msec² range, yielding q estimates close to 25, symmetry coefficients averaging about .05 or perhaps .10, average Kurtosis coefficients between about 2.9 and 3.4, -and average correlation coefficients smaller in absolute value than .15 or .10, with no change in any of these as the interval is varied in small steps across the range, once an initial scan of the range has been completed. Given the individual and procedurally based differences found in auditory work, however, I would not expect to find this perfect pattern, or, at least, not for all subjects. Despite the mass of data collected in Experiment 4, we will be confronted with a subjective)udgment at the end, about whether the results are in good enough general agreement with the prototypic pattern to warrant a conclusion that visual and auditory performance are the same. In the event, the data from the two subjects who completed both full scans of the range, IS and DK, were quite similar, and were similar enough to the prototype that this subjective decision should be a relatively easy one to make. That is a fortunate result, and the discussion below is

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fairly straightforward because of it, but the above notes outline what I thought (and think, if we consider replications in the future) should be obtained, not what was obtained. Given the variation in auditory results across subjects and experiments, a poorer match between expectations and data could well have been obtained, and could be obtained in the future, even if the underlying timing mechanism is the same across modalities.

There is one final point that I wish to make explicit about the list of expected outcomes above. The list of possible comparisons is probably endless. I have certainly not made all of the comparative analyses that were possible. When confronted with a selected series of comparisons from a much larger pool, a reader might wonder whether other comparisons were made, and if so, why they weren't reported. Sometimes this is justified: the practice of reporting only those "predictions" actually met by the data, leaving unmentioned the predictions not met or the results which might be considered unfavorable, is not unheard of. I consider such practices fundamentally dishonest and I do not engage in them. If there are unmentioned aspects of the data that in any way conflict with the conclusions drawn, they are unmentioned because I am unaware of them.

5.1 Method

5.1.1 Subjects

Five subjects started Experiment 4. DK, LL and IS participated throughout the experiment. AP, a female undergraduate, vanished after 39 sessions. JS, a male graduate student, started a few months later,

as a replacement for AP. By the time he had completed 75 sessions, the data from the other subjects indicated that this experiment would require more than twice as many sessions as I had originally anticipated. While JS very reliably completed the sessions for which he was scheduled each week, by that schedule he could not complete even half of the experiment. I gave him the choice of increasing his number of sessions per week or of leaving. Having other more pressing commitments, he left the experiment.

DK and LL are female, IS is male. DK and IS are senior graduate students. LL is a research technician. All had participated in other experiments involving duration perception.

5.1.1.1 Experiment 5

RW, the subject of Experiment 5, is a senior male graduate student, who had already participated in Experiments 2 and 3.

5.1.2 Procedure

5.1.2.1 Experiment 4

In the previous experiments, scanning across a range was done as quickly as possible, with extended practice only at a few intervals and very little practice at other intervals. In this experiment, the highly practiced intervals were 300, 400 and 500 msec. Extra practice, though not as much as at these three, was also given at 305, 310, 315, 320, 325, 475, 480, 485, 490 and 495 msec. My intention was to pay careful attention to performance at these intervals, in order to test for continuous adjustability.

Originally, I planned only one scan upward, to 500 msec, and a return to 300 msec, as in Experiment 1. The variances did not drop to a constant level during the return scan, but they did drop substantially, and the reduction was evident across the 6 very close intervals, 500 to 475 msec, at which extra training was being given. I increased the number of sessions per interval to 5, at each interval planned during the scan down for subjects DK and IS. These two subjects were willing to participate in an extraordinary number of sessions, allowing me to take a detailed look at this reduction in variance, which had been quite striking over the short range from 500 to 475 msec. The variances tended to level off eventually. In IS's case, they appeared constant from 400 to 275 msec during this scan down. It appeared that they might level off completely if the scan was repeated, and I had them conduct a full, slow, second scan back up to 500 msec. DK completed the scan. IS has progressed to 430 msec at this date. Due to illness and to other commitments, he has completed very few sessions in the past three months. I don't know at the time of this writing whether he will complete the full series or, for that matter, whether he will complete any further sessions. While I will still collect the data if he generates it, he participates so infrequently now that it seemed unwise in the extreme to delay describing this work until he does complete the series. If there are substantially more data from IS, I will discuss them during the oral examination and will append a summary of them to the dissertation at that time.

While DK and IS were willing to take part in more than twice as

many sessions as originally anticipated, LL could not do so. For her to complete a full scan up and down, it was necessary to abbreviate her schedule slightly, relative to the original plan, omitting the extra practice across the short range from 300 to 325 msec.

JS left the experiment during the first stages of his first scan up the range. AP left during stabilization training with her first interval.

5.1.2.2 Experiment 5

RW completed Experiments 2 and 3 far too late for extensive work in Experiment 5. I originally intended to examine his performance only at a single interval, 300 msec, which was the last interval he worked with in Experiment 2. His responses were correlated with the foreperiod at 300 msec, and his variances were high, so I conducted a very brief scan to 400 msec and back in the hope that this would yield better results at 300.

5.2 Results

Tables of the seasion by session data appear in Appendix 1. JS's results appear there only. I have not generated such a listing for AP's results, nor examined them in any detail, and there is thus no listing of her data in the appendix. She left the experiment after so few sessions that there isn't anything to learn from the data.

5.2.1 Latencies

Figures 12.1 (DK), 12.2 (IS), 12.3 (LL) and 12.4 (RW) show the

response latencies (upper panels) compared to the synchronization intervals (lower panels). All of the subjects' latencies tracked the intervals closely. The mean latencies were typically within 10 msec of perfect synchrony with the onset of P2.

5.2.2 Variance Levels and q

Figures 13.1 (DK), 13.2 (IS), 13.3 (LL) and 13.3 (RW) show each session's variance (upper panel) and synchronization interval (lower panel). In preceding figures of this type, for Experiments 1 to 3, 1 showed foreperiod durations in a third panel, at the bottom of the page.. There is little point to such a display here because almost all of the sessions were controlled by the Apple. Apple mean foreperiods were all quite close to 1000 msec. The only sessions in which the PDP-8 was used were IS's first 8 and DK's first 6. Accordingly, I have omitted this graph for these experiments.

Tables 7.1 (DK), 7.2 (IS), 7.3 (LL) and 7.4 (RW) give average values of variance, q, coefficients of symmetry $(\sqrt{b_1})$, kurtosis (b_2) , and correlation with the foreperiod, averaged across the final five sessions at each of the practiced intervals.⁴ All statistics are based on latencies as measured from P1. Session by session listings of statistics concerning latencies measured from WS are in Appendix 1 but these are of little or no interest because the correlations between response latency and foreperiod duration were generally quite small.

Figures 14.1 (DK), 14.2 (IS), 14.3 (LL) and 14.4 (RW) display the average variances listed in Tables 7.

FIGURE 12

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Upper panel: Mean synchronization latencies (from P1), in msec, of each session. Two sessions' data are missing and are indicated by gaps in the figure.

Lower panel: synchronization intervals, in milliseconds, of each session.

Figure 12.1: Subject DK. Figure 12.2: Subject IS. Figure 12.3: Subject LL. Figure 12.4: Subject RW.

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 $f_{i} = \frac{1}{2} \left[f_{i} \right]$



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FIGURE 13

Upper panel: Response latency variance (msec²) of each session. Two sessions' data are missing and are indicted by gaps in the figure. Variances shown as extending to the topmost line of the figure are larger than they appear. For exact values, see Appendix 1.

Lower panel: synchronization intervals, in milliseconds, of each session.

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Figure 13.1: Subject DK. Figure 13.2: Subject IS.





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FIGURE 14

Mean within-session response latency variances (msec²) as a ______ function of mean response latency (msec). Arrowheads show the first ascending series. Circles show data from the descending series. Squares show data from the second ascending series. The variances ______ plotted are listed in Tables 7.

Figure 14.1: Subject DK. Figure 14.2: Subject IS. Figure 14.3: Subject LL. Figure 14.4: Subject RW.



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TABLE 7

Mean within-session variances (msec²), q estimates, coefficients of , symmetry (ROOT B1), of Kurtosis (B2) and of correlation between response latency measured from P1 and the foreperiod duration. Only data from intervals at which the subject spent two or more consecutive sessions were averaged for this table. The averages are based on the last five sessions at this interval before the subject was transferred to the next, or, if there were fewer than five, on all sessions at the interval.4

Table 7.1: Subject DK. Table 7.2: Subject IS. Table 7.3: Subject LL. Table 7.4: Subject RW.

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SUMMARY STATISTICS FOR DK VISUAL RESPONSE LATENCY MEASURED FROM P1

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INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH FP
300	711	64.16	, 0.03	2.78	-0.04
305	535	55.33	0.02	3.20	-0.07
310	534	55.27	0.10	2.93	-0.04
315 🕈	474	51.91	-0.19	3.54	-0.03
320	337	43.26	-0.03	3.78	0.02
325	340	43.48	-0.10	3.58	-0.01
400	- 380	46.16	0.05	3.43	0.01
500	528	54.94	0.04	3.33	-0.04
495	468	51.53	-0.02.	2.92	-0.03
490	412	48_21	0.06	2.89 4	• -0.10
485	383	46.33	0.03	3.14	-0.02
480	399	47.37	0.05	2.92	-0.07
475	364	45.10	0.00	3.10	-0.03
470	349	44,12	0.06	3.15	-0.03
465	281	39.19	-0.11	3.11	-0.04
460	303	40.82	0.02	3.26	°-0.07
455	320	42.09	0.03	3.16	-0.05
450	316	41.81	0.00 .	3.25	-0.01
445	265	37.92	0.11	3.21	-0.05
440	279	39.02	0.17	3.42	-0.04
435	262	37.67	0.01	2.90	-0.03
430	278	38.97	, 0.05	3.19	-0.05
425	269	38.24	0.18	3.11	0.00
420	269	38.25	-0.02	3.26	-0.02
.415	- 249	36.63	0.12	3.11	-0.05
410	220 ·	34.24	0.0ð	3.28	-0.02
405	260	37.55	-0.01	3.21	-0.01
400	219	34.08	0.11	3.27	0.01
395.	212	33.49	0.12	3.47	0.01
390	226	34.73	0.09	3.41	0.01
385	. 176	30.10	0.02	3.12	-0.01
380	229	34.95	-0.09	3.04	0.05
375	227	34.84	0.17	3.93	0.02
370	191	31.55	-0.09	3.46	-0.02
365	210 -	33.28	-0.06	3.14	-0.04
360	196 .	31.99	-0.16	3.35	-0.04
355	200	32.36	-0.11	3.09	-0.06
350	203	32.69	0.09	3.08	-Ò.10
345	177	30.25	-0.23	3.36	-0:05
340	193	31.79	-0.01	2.97	-0 09

SUMMARY STATISTICS FOR DK VISUAL RESPONSE LATENCY MEASURED FROM P1

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INTERVAL	VARIANCE	Q	R00T 81	B2	CORR WITH FP
335	198	32.19	-0.20	3.64	-0.08
330	189	31.37	-0.12	3.04	-0.10
325	192	31.62	0.04	'3.13	-0.06
320	193	31.78	-0.01	3.43	-0.08
315	189	31.39	-0.04	2.87	-0.01
310	199	32.34	0.15	3.22	-0.09
305 _	1 <u>68</u> }	29.24	0.04	2.89	-0.15
300 (181	30.57	-0.08	3.26	-0.08
275 \	157	28.16	-0.20	3.33	-0.08
300	156	28.04	0.01	3.89	-0.06
305	171	29.62	-0.11	3.31	-0.14
310	145 -	26.84	, -0.03	2.86	-0.10
315	145	26.81	0,12	4.24	-0.03
320	179	30.41	-0,03	3.03	-0.06
325	195	31.94	-0.09	3.25	-0.09
330	192	31.65	0.01	3.16	-0.06
340	204	32.79	-0.01	3.30	-0.08 🖙
350	178	30.28	0.00	3.51	-0.08-
360 /	^{′′′} 163	28.76	0.19	3.20	-0.07
370 /	167	29.20	0.08	3.09	-0.08
380	173	29.78	0.03 /	3.30	-0.05
390	185 -	30.98	0.11	3.00	-0.07
400	178	30.25	0.04	2.78	-0.12
410	179	30.42	0.04	3.06	-0.11
420	187	31.22	0.03	2.95	-0.11
430 🕺	223	34.51	-0.02	3.10	-0.12
440	207	33.02	-0.02	3.03	-0.02
450	209	33.22	-0.04	2.96	-0.07
460	221	34.27	0.01	2.73	-0.13
470 -	213 د	33.58	0.01	3.09	-0.09 ·
480	200	32.44	-0.04	2.95	-0.10
490	215	33.79	-0.11	3.18	-0.11
500	156	27.99	-0.10	3.58	-0.09

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SUMMARY STATISTICS FOR IS VISUAL RESPONSE LATENCY MEASURED FROM P1

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INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH FP
300	345	43.82	0.05	3.72	0.02
305	346 _	43.89	0.16	3.61	-0.01
310	339	43.42	-0.13	3.81	-0.06
315	331	42.83	0.46	3.88	-0.02
320	333	43.01	0.25	3.46	-0.02
325	308	41.17	0.04	3.26	0.00
400	355	. 44.52	0.26	3.07	0.10
500	427	49.10	0.07	3.18	0.00
495 .	513	54.09	• 0.25	3.42	0.04
490	370	45.47	-0.04	2:97	-0.05
485	323	42.26	0.00	2.98	0.07
480	272	38.52	0.08	3.43	10.0
475	240	35.94	-0.01	2.98	-0.03
470	214	33.67	0.00	3.26	0.01)
465	236	35.60	-0.13	3.08	002
460	244	36.27	-0.21	3.15	-0.03
455	241	35.98	0.00	3.16	0.03
450	207	33.07	-0,03	3.36	0.03
445	216	33.81	0.]44	5.32	-0.01
440	237	35.66	0.00	3.03	-0.01
435	186	31.04/	0.01	2.82	-0.01
430	178	30.27	0.07	3.01	0.06
425	203	32.66	-0.15	2.95	-0.04
420	159	28.35	-0.08	3.15	-0.03
415	180 .	30.49	0.23	3.07	-0.03
410	151	27.52	0.06	3.60	0.01
405	155	27.89	-0.10	2.94	0.04
400	150	27.38	0.01	2.93	-0.02
395	149	27.28	0.12	3.26	0.02
390	153	27.76	-0.08	2.95	0.00
385	142	26.47	0.05	3.30	0.00
380	127	24.69	0.00	3.06	-0.02
375	159	28.38	-0.14	3.41	-0.02
370	162	28.66	-0.04	3.36	-0.01
360	170	29.45	-0.10	3.07	-0.08
360	132	25.28	0.04	3.15	-0.06
300	120	27.39	-0.05	3.32	0.01
300	179	30.40	-0.18	3.37	-0.06
345	194	31.84	-0.03	3.46	-0.11
34U	133	25.44	0.01	3.71	-0.02

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-SUMMARY STATISTICS FOR IS VISUAL RESPONSE LATENCY MEASURED FROM P1

INTERVAL	VARIANCE	Q	ROOT B1	82	CORR WITH FP
335 -	149	27.32	0.07	3.40	0.01
330	174	29.93	0.04	3.41	-0.10
325	177	30.20	0.08	3.53	-0.05
320	129	25.02	0.03	3.37	-0.09
315	156	28.03	-0.06	3.53	-0.03
310	158	28.21,	-0.04	4.55	0.06
305	141	26.34	0.08	2.93	0.05
300	139	26.13	-0.01	3.58	-0.06
275	132	25.31	-0.17	3.44	-0.03
300	95	20.52	-0.03	3.24	-0.03
305	109	22.51	0.08	3.08	0.02
310	113 🔍	22.92	0.19	3.40	0.02
315	129	25.02	0.13	3.61	0.02
320	116	23.37	0.11	3.76	0.01
325	117	23.51	-0.28	4.16	-0.04
330	122	24.17	0.05	3.39	0.02
340	142 /	26.44	-0.11	3.19	0.03
350	131	25.19	0.14	3.11	0.04
360	132	25.29	0.11	3.07	-0.01
370 .	159	28.34	0.10	3.30	0.05
380	130	25.11	0.22	3.44	-0.02
390	137	25.88	0.11	3.10	-0.04
400	118	23.68	0.17	3.40	-0.01
410	139	26.12	0.15	3.46	-0.02
420	. 148	27.18	0.20	3.76	-0.03
430	130	25.12	0.11	3.12	0.01

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SUMMARY STATISTICS FOR LL VISUAL RESPONSE LATENCY MEASURED FROM PI

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INTERVAL	VARIANCE	Q	ROOT BI	B2	CORR WITH FP
300	397	47.21	0.04	3.54	-0.02
. 305	493	52.99	0.31	3.58	-0.03
310	566	56.95	0.41	4.27	0.00
315	513	54.11	0.11	4.44	0.01
320	538	55.46	0.32	3.77	-0.01
325	416	48.41	0.10	3.06	-0.02
400	393	46.99	0.18	3.29	-0.02
465	428	49.19	0.30	3.32	-0.18
500	438	49.78	0.42	3.60	0.01
- 495	396	47.16	0.30	3.73	-0.01
490	386	46.56	0.30	4.31	-0.07
485	316	41.81	0.36	3.80	-0.01
480	321	42.14	0.08	3.15	°-0.03
475	280	39.11	0.24	3.20	0.01
450	341	43.55	0.28	3.53	-0.02
480	295	40.27	0.22	3.34	-0.06
350	233	35.29	0.17	3.42	-0.05
300	299	40.52	0.12	3.17	-0.02

SUMMARY STATISTICS FOR RW VISUAL RESPONSE LATENCY MEASURED FROM P1

INTERVAL	VARIANCE	Q	ROOT B1	B2	CORR WITH F	Ρ
ر 300 پ	281	39.22	0.12	3.24	-0.11	
350	310	41.37	-0.02	3.00	-0.06	
400	392	46,93	0.22	3.45	-0.09	
390 🅖	308	41.22	0.09	3.22	-0.07	۳
380	312	41.50	0.05	3.01	-0.09	
370	317	41.87	0.04	3.11	-0.07	
· 360	405	47.74	-0.08	3.55	-0.10	
350	269	38.30	0.03	2.99	-0,14	
340	244	36.28	0.07	3.13	÷0.16	
330	251	36.82	0.06	3.13	-0.12	
320	232	35.23	-0.03	3.13	-0.12	
310	238 -	35.71	0.04	3.25	-0.15	
300	287	39.64	0.22	3.77	-0.18	

The results from DK and IS are quite similar. DK's final variances (averaged across the last 5 sessions per interval) ranged from 156 to 223 msec². The variances at the intervals at which extensive stabilization training was done, 300, 400 and 500 msec, were 156, 178 and 156 msec² respectively. Average estimates of q during the final scan upward range from 28.0 to 34.5, hovering about 30. IS's variances were lower, ranging from 95 msec² to 148 msec². At the two intervals at which there was long practice, 300 and 400 msec, his variances were 95 and 118 msec² respectively. Estimates of q for IS range from 20.5 to 28.3, generally quite close to 25.

IS's variances were lower than any obtained in Experiment 2 and they compare well with the previous auditory results of Brewster (1983) and Kristofferson (1976). His q is right where it should be: close to 25. DK's variances were higher, but they were certainly as good as those obtained in Experiment 2. They are slightly better than those obtained from one of Brewster's subjects. DK's q.is not 25. I doubt that her variability would decline much further with more practice, so hould not expect her q estimates to drop to 25.

If we interpret the data in terms of Hopkins' model, q is the scale parameter of a triangularly distributed variable, which is convoluted with a logistic variable to yield the actual response distribution. I estimate q by subtracting 25 msec² from the response latency variance, assuming that this is the variance of the logistic motor component. The remaining variance is assumed to be that of the triangle, which is specified by the formula: Variance $= q^2/6$. This is a rough estimation procedure. One might

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whether DK's motor variance is not higher than IS's, in which case her \cdot q estimate would be higher than his simply because not enough motor variability was subtracted from her latency variance. This is not the explanation. DK's kurtosis coefficients are not higher than IS's -- &f they are different, DK's are lower -- and we should find kurtosis increasing with increasing motor variance, i.e. with increasing contribution of the higher Kurtosis logistic variable to the overall variance of the data. If DK's motor variance is so high that subtraction of all of it would result in a drop in the q estimate to 25, then IS's q_should_drop to 20, and we would face the same difficulty. Supposing that DK's q is 30, the question is, is this "close" to 25 or not? I don't know. The estimates of 12.5, 25, 50 and 100 for q levels are empirical. There is no theory that forces q to take on these particular levels. Values of 15, 30, 60 and 120 would be every bit as compatible with Kristofferson's (1980) quantal counting with steps model, for example. There is no quantitative theoretical expectation regarding variability of q across people, although given that guantal theory ties the glevel to a fundamental (if hypothetical) physiological process, we should not expect "too much" variance. Is a deviation, of 20% from the 25 msec level "too much"?

In terms of overall variance levels, I conclude from IS's and DK's data that visual and auditory synchronization variances are, or at least can be, the same. There is no evidence in these data that visual variances are higher than auditory variances. With respect to q, IS's results are unambiguous and, given these, I hesitantly interpret DK's q of 30 as passably close to 25.

LL's variances are higher than those of IS and DK but it must be remembered that she completed fewer than half as many sessions (261 compared to DK's 691 and IS's 540). Her variances during the initial scan upward, when she and the other subjects were on the same training schedule, are quite comparable to those of the other subjects, ranging from 393 to 566 msec², compared to DK's first scan range of 337 to 711 msec² and IS's 308 to 427 msec². LL's scan back to 300 msec was much faster than IS and DK's. Her variances on the way down the range were lower (280 to 396 msec², compared to 393 to 566 msec² during the scan up). Presumably they would drop further with further practice. Given that this is not asymptotic performance, I am not concerned that the q estimates are near 40 msec, with one value near 35. Values slightly higher than 40 might be acceptably close to 50 or they might not, but these variances have not settled down to a final level and the 25, 50 or 100 msec q values are predictions about final level performance.

LL's data highlight one problem with comparison of visual and auditory performance. The visual results may drop to the auditory levels, but it takes a long time to get them this low. It may take twice as many sessions, depending on the subject, before the low variance levels are reached. Perhaps with this much practice with auditory intervals, lower auditory levels would be reached.

I take it as granted that visual synchronization is more difficult than auditory synchronization. If there was no difference, Hopkins subjects would have used the visual stimuli, not the tactile feedback from the trial initiation response, and the delayed (auditory

and visual) feedback given after P2. I noted in the introduction a number of sources of difficulty, for the subject, in the visual situation. I don't think that these differences between the auditory and visual tasks can be eliminated, though perhaps a huge visual display (a wall, rather than a degree visual angle) would minimize them. It would be most pleasant, theoretically and practically, if visual subjects' variances declined at the same rate as auditory subjects' variances. But given the differences in physical variability (due to the changing posture of the subject) and of salience between the visual and auditory stimuli, the finding that subjects take longer to finally bring their visual synchronization standard deviations' down to 0 msec, a remarkable level of precision whatever the stimulus modally, cannot be taken as an important difficulty for theory.

Turning to Experiment 5, RW's data were more variable than his auditory data, and they remained more variable. His responses were correlated with foreperiod duration, negatively, as in his Experiment 3 data, rather than positively as they were in his comparably long Experiment 2 performance. I'll return to this later. For now I will note that the variances of the latencies measured from the warning stimulus are higher than the variances from P1. His variances are frankly disappointing, however they are calculated.

5.2.3 Variances Across the Range

The relation between duration and variance was not monotonic in LL's data. The variances differed so much from interval to interval that I am wary of calling them constant across the range. Figures 13.3

and 14.3 describe LL's results better than I can verbally and I will say no more about them.

DK_and IS showed an increasing relation between duration and variance during their initial scan up to 500 msec. This was also evident in the scan down, at least to 400 msec. From 400 to 275 msec the variances change little in either subject's data.

During the final scan upward, there was some tendency for the variances to increase again. As the intervals were lengthened, the variances of the first session or two were often higher than those obtained later, especially in DK's data. IS's variances were more stable and fluctuation in his variance at a given interval often appeared random.

The increase in DK's variances is most evident in her final five session averages from 400 to 500 msec. Variances rose from 178 ("msec² to values quite consistently, if sometimes only slightly, greater than 200 msec². The abrupt drop to 156 msec² at a P1-P2 interval of 500 msec from 215 msec² at 490 is not due to random fluctuation in the data. DK ended the experiment with 100 sessions at 500 msec. Her variances slowly declined from the low and mid 200's to values around 175 msec², and, finally, to values around 155 msec², the same as her performance at 300 msec. Transfer of training, and previous training at these intervals, did yield a substantial drop in variance relative to her earlier performance, during the first ascending series, but this transfer effect seemed to be timited. It appeared that to achieve levels below 200 msec², DK required extensive recent practice at the interval

under study. Kristofferson (1976) and Hopkins (1982) were more successful in transfer studies. Across the 150 to 550 msec auditory range, it appears that once a subject achieves a low level of variability, that level can be maintained across the range, even withchanges of 100 msec in the P1-P2 interval. That degree of success was not evident in Experiment 2 (or Experiment 1), but so little practice was given at intervals other than the highly practiced ones that we have little basis for comparison in those data. It is at least possible, given DK's results, that to achieve asymptotic visual performance levels one requires more training at the specific interval under study, as well as more training across the range, than is true for auditory synchronization. I am not certain that this is a "real" difference between auditory and visual results. This may merely reflect a difference between subjects. Supposing that the difference is replicable, and the slight increasing trend in IS's data suggests that it may well be, I am not at all sure that it should be considered an important difference.

I think that the key finding in these data is that after much training at specific intervals (300, 400, 500 msec), variances do not differ. Nor is there any trend in the coefficients of skewness (DK's were .01, .06, and -.10 at the three intervals; IS's were -.03 at 300 and .17 at 400, but they were generally positive from 300 to 430 msec) or in the coefficients of Kurtosis (DK: 3.09, 2.78, 3.58; IS: 3.24 and 3.40, but there is no hint of an increasing trend from 300 to 430) or in the correlation between latency and foreperiod duration (DK: -.08, -.12, -.09; IS: -.03, -.01). The distributions are very similar in

scale and shape at each of the practiced intervals. This is precisely what we should expect on the basis of the auditory results.

RW's transfer data again tell a different story. His variances are roughly increasing with duration, during the scan up and back down. His timing variances may have been constant from 350 to 300 msec, i.e. across half of the range studied, but I wouldn't argue forcefully that this was the case.

5.2.4 Scanning of the Range

All subjects' variances were lower during the descending series than they were during the initial ascending series. Variances tended to level off across a range over which they had previously been increasing in IS and DK's data (see Figures 14). LL's variances varied less during the descending series, and in this sense they approached constancy as well. During the final ascending series, DK and IS's variances declined further and they varied less over a wider range. This is all in accord with the expectations from the auditory data.

5.2.5 Continuous Adjustability

The question in this case is whether very small changes in the synchronization interval will have effects on the response distribution's scale and shape. If the subject can change her mean by a few msec without changing any other aspect of the distribution, we have evidence of a process that can generate delays on a continuous (rather than discrete, as in quantal) scale. If the subject can also produce widely different intervals without changing the scale

(variance) or shape of the distribution, which we have already seen can be done in visual synchronization, we have evidence for a deterministic source of internal delays, i.e. one which times intervals without variability. If the same process is involved in timing similar intervals and more widely separated intervals (and if the response distributions are all the same except for the means, why would we want to assume that different processes are involved?), then we have evidence for a source of continuously adjustable deterministic delays.

The relevant data come from DK and IS, especially at 300, 305, 310, 315, 320 and 325 msec. These subjects practiced for extra sessions at these intervals so if there are effects of shifting the P1-P2 interval slightly (by 20% or less of q), they should show up here more clearly.

Examining DK's data first, from 300 to 325 msec we find variances of 156 (181), 171 (168), 145 (199), 145 (189), 179 (193) and 195 (192) msec² respectively. The bracketed values are from the end of the descending series. The values not bracketed are from the start (final five sessions, but among the first intervals) of the final ascending series (which immediately followed the descending series). There is no greater variation across intervals than there is from replication (last 5 sessions, descending) to replication (last 5 sessions, ascending) at the same intervals. There is no trend in the symmetry coefficients either. The ascending and (bracketed) descending series values are .01 (-.08), -.11 (.04), -.03 (.15), .12 (-.04), -.03 (-.01) and -.09 (.04). The kurtosis values are 3.09 (3.26), 3.31(2.89), 2.86 (3.22), 4.24 and 2.87, 3.03 (3.43) and 3.25 (3.13). These

coefficients are highly variable (as they should be) but the variation does not look systematic. The correlations are all nearly the same.

I'll repeat this listing for IS in the next paragraph. There is no indication of a shift in scale, shape or correlation in his data either. The other way to look for trends, and probably the most powerful way, is to scrutinize Figures 13 (variance per session), 14 (average variances across the final five sessions at each interval), 15 (correlation), and 16 (symmetry and kurtosis) for periodic or monotonic changes in the distributions as the interval was changed during the final ascending series. Except for a slight increase in variance with duration, I see no trends in DK's or in IS's data in any of the graphs.

Returning to IS's data, from 300 to 325 msec, during the ascending and (bracketed) descending series the respective variances are 95 (139), 109 (141), 113 (158), 129 (156), 116 (129) and 117 (177) msec². The descending series variances are higher. That's the only trend that I see here. The symmetry coefficients are -.03 (-.01), .08 (.08), .18 (-.04), .13 (-.06), .11 (.03) and -.28 (.08). The kurtosis coefficients are 3.24 (3.58), 3.08 (2.93), 3.4 (4.55), 3.61 (3.53), 3.76 (3.37) and 4.16 (3.53).

If we accept the conclusion that there is a source of continuously adjustable internal delays for timing auditory intervals, we must also accept the conclusion that there is one for visually bounded intervals.

If we accept Kristofferson's (1976) Hypothesis that these internal delays are generated in the afferent component of the stimulus response chain, then either visual afferent latencie are not variable
or there is a temporally variable and a nonvariable subsystem(s) in the visual system, with recycling of the stimulus information (the basis of the delay) only being done in the nonvariable subsystem. To turn this around, Hopkins argued that visual latencies must be more variable than auditory latencies (which appear to be constant) because visual synchronization variances appeared greater than auditory variances. The poor visual performance in his task appears, from the results discussed so far, to have been due to the details of his procedure and stimuli, not to any problem intrinsic to the visual system. There is no reason to suppose, from these data, that visual afferent latencies are variable, or, if they are variable, that they are more variable than auditory latencies. Visual synchronization data, then, in no way conflict with Kristofferson's hypothesis that continuously adjustable internal delays (now equally strongly demonstrated in the visual and auditory modalities) are generated at the afferent level.

5.2.6 Correlations

There are indeed mild correlations between latency and foreperiod duration in the data of Experiment 4. Figures 15 display the correlations obtained during each session. Tables 7 list the averages at the practiced intervals. The correlations are no greater than those obtained in Experiment 1, when very long auditory intervals were studied. They are sufficiently small that we can consider the dominant controlling stimuli in the task the visual stimuli, i.e. Experiment 4 was in fact as well as in name a study of visual synchronization. There is no evident relation between correlation and

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FIGURE 15

Upper panel: Correlation between response latency (from P1) and foreperiod duration for each session. Some sessions' raw data were lost and correlations could not be calculated. These are indicated by gaps in the figure.

Lower panel: synchronization intervals, in milliseconds, of each session.

Figure 15.1: Subject DK. Figure 15.2: Subject IS. Figure 15.3: Subject LL. Figure 15.4: Subject RW.

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SESSION, 30 60 90 120 150 180 210 240 LL VISUAL

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FIGURE 16

Upper panel: Symmetry coefficients of the response latency (from P1) distributions of each session.

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Lower panel: Kurtosis coefficients of the response latency (from P1)

Some of the coefficients were too large in absolute magnitude to be represented accurately in these figures. Where there are two lines at the top or bottom of a figure, and a value is shown as extending to the outermost line, it is larger than it appears. For exact values, see Appendix 1.

Some sessions' raw data were lost and symmetry and Kurtosis coefficients could not be calculated. No values are shown for these sessions, which are indicated by gaps in the figures.

Figure 16.1: Subject DK. Figure 16.2: Subject I∰ ⇒ Figure 16.3: Subject LL. Figure 16.4: Subject RW.



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P1-P2 interval duration in Experiment 4.

RW's data are different. His correlations are substantial, comparable in sign and size to his performance at the very short intervals of Experiment'3. His variances are also at the Experiment 3 (eg. 100 msec P1-P2) levels. As in Experiment 3, RW is not relying exclusively on the warning stimulus. This can be checked by examining the variances of his latencies measured from WS, which bare higher than those obtained from P1 (see Appendix 1's tables). Further, and surprisingly given the correlatations, RW's distributions (latencies from Pi) are not negatively skewed, as they should be if he is timing from WS. They are positively skewed or symmetric. The kurtosis coefficients are not as large as they should be (or as they were in $^{
m N}$ Experiment 3). Whatever RW was doing in Experiment 3, he is doing something similar, but it is not the same thing, in this experiment. He is in some way mixing-a strategy of timing from P1 with one of timing from WS. This is evident in the correlations, such as -.18 at his most practiced interval (300 msec). I have no suggestions to offer about the details of this strategy.

RW was trained in Experiment 3 to use the warning stimulus instead of P1. It is unfortunate that RW's training with auditory stimuli transferred in this way, and it is somewhat surprising that it would transfer in this way since he was not timing from WS at the end of Experiment 2, at 300 to 350 msec intervals, with which he finished working only a few days before starting Experiment 5 at 300 msec. However, given that RW is relying more heavily on WS than the subjects of Experiment 4, it is not surprising that his pattern of results is

different from theirs.

5.2.7 Symmetry and Kurtosis

As in the previous experiments, there is a very slight positive skew to the response distributions. This can be seen in Figures 16 (upper panel) and in Tables 7. The skew is unusually large (relative to the other subjects: these are not very skewed distributions) and consistent in LL's case. DK's and IS's symmetry coefficients are no different from those of Experiments 1 and 2.

As in the previous experiments, the Kurtosis coefficients are usually between 2.4 and 4.2 (the logistic plus triangle limits) and the five session averages are usually between 2.9 and 3.4.

LL's kurtosis coefficients are large and they are larger than those obtained from IS and DK during their initial/ascending and descending series. This is not strictly idiosynchratic. JS's data look the same. His final averages at the 5 intervals (300 to 320 msec) at which his performance was studied ranged from .12 to .23 for the symmetry coefficients and from 3.23 to 3.75 for the kurtosis coefficients. However, these were not obtained after much practice (only 75 sessions total). IS's coefficients were also large during his first 75 sessions and his distributions eventually became symmetric and reasonably peaked, so perhaps JS's distributions would have shifted in shape as well.

On the basis of the data at hand, two different descriptions of visual synchronization distributions are defensible. One set of distributions is as symmetric and peaked as we find in auditory

synchronization. The other set are more positively skewed and they are less peaked. The distributions which look most like the auditory distributions come from the most practiced subjects, whose data look like auditory data in most other respects as well.

5.3 Summary

There were individual differences. What is true of the performance of IS and DK is often not true of the performance of LL, in Experiment 4. RW's performance was different again but the relatively high correlation between response latency and foreperiod duration in his case indicates that he was relying to a large degree (as much as when P1-P2 was 100 msec) on the auditory WS dr on tactile feedback from his trial initiation response as well as on the visual P1. There are curious aspects of his performance, such as the lack of negative skew in his response distributions, expected given the correlation, but conclusions about these data should be with reference to a multi-modality synchronization task, not to a unimodal visual synchronization task which was the desired object of study.

Considering only the data of Experiment 4, from IS and DK we see almost all of the same trends as are evident in auditory synchronization. The effects of practice are not quite the same. Scanning is effective; but there is less transfer of training from interval to interval, and more practice at any given interval seems necessary before asymptotic performance levels can be reached. Perhaps these differences would disappear if a bigger, brighter stimulus was used, assuming that that stimulus didn't have other deleterious effects

on performance. The main points of comparison, variance, symmetry, kurtosis, foreperiod correlation, variability and shape across narrow and wide ranges, scanning effects and absolute levels of q (with the possible reservation about DK's 30), all appeared to match the auditory results of Experiments 1 and 2 and of Kristofferson (1976) and Brewster (1983) acceptably. In terms of shape coefficients, the data are 'near Hopkins' values.

As noted in Experiment 1, if we accept Hopkins' model, the relatively high Kurtosis values (but not higher, or not much higher, than Hopkins') would suggest substantial motor variability, on the order of 50 to 100 msec². DK and IS participated in so many sessions that I find it extremely difficult to believe that their final motor variances were this high. However, despite the finding that their data do not fit well with the model constructed to describe Hopkins' (and Kristofferson's 1976) data, they are, except in variance level, very similar to Hopkins' data.

LL's data, and, with reservations due to the small number of sessions, JS's data, may indicate that there is more than one way to do the visual synchronization task. Their distributions were more skewed and less peaked than the others, auditory or visual. It is interesting that JS's and LL's variances were also relatively high. It is tempting to conjecture that IS's and DK's performance is better because they relied on the same timing mechanisms used by the auditory subjects, whereas JS and LL followed a suboptimal strategy.

However we interpret JS's and LL's performance, the results from DKmand IS make the point of the experiment. To the degree that we

are justified in concluding, on the basis of multiple similarities in performance across modalities that there is a common underlying mechanism across modalities, we are justified in concluding that the same mechanism is available to be used by auditory and visual synchronization subjects. Given the many dimensions of similarity, I would find it very difficult to believe that different mechanisms were responsible for auditory and visual synchronization performance, as obtained in the visual case from DK and IS.

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Conclusions

My conclusions are based mainly on the data from three subjects of the seven discussed in detail in the body of this work.

Two subjects failed to achieve stable performance levels, GH for reasons unKnown, and LL almost certainly because of insufficient training.

The two subjects of Experiments 2 and 3 were probably transferred too quickly to Experiment 3 (though "quickly" is a relative term and in practical terms, if they had spent more time in Experiment 2 pretraining, they would never bave made it through Experiment 3 and Experiment 2 retesting. Hundreds of sessions' data take years to collect. Both of these subjects would have preferred to have participated in fewer sessions than they did and they would not have participated in substantially many more.) In both cases, performance at intervals of 150 msec or longer was better after Experiment 3 than before, indicating that they were transferred to Experiment 3 before they had achieved final vaniance levels at the longer intervals. I hesitate to take their post-Experiment 3 data at face value. I don't know what stransfer effects from that extremely difficult task may be hidden in their performance. I compared Experiment 4's visual results to published auditory data as much as to those from Experiment 2 . because of my concern that the relatively high variances obtained in Experiment 2 may not only have been due to the specific differences between my version of synchronization and the others, but also to transfer effects.

This leaves three Key subjects, DG from Experiment 1, and IS and DK from Experiment 4, whose data have guided my conclusions. Their data strongly support the thesis. The data from the other subjects do not strongly conflict with it, but I would not accept it on the basis of those results alone.

There are three classes of findings that I consider firm in this work, one important hypothesis that is suggested by the results, and one finding that poses some difficulties. The three positive findings are:

(1) Equivalence of long and short interval synchronization: There is every indication in Experiment 1's data, especially in DG's performance, that auditory synchronization is based on the same mechanisms at 2000 msec as it is at 200 msec. Variances were constant from 1500 to 1750 msec, with a strong suggestion of constancy out to 2000 msec in DG's data. DG's q estimates of 51.3, 50.9 and 58.4, at 1500, 1750 and 2000 msec respectively, are near 50, one of the values we should expect q to hover about from other quantally-related data and theory. The distributions are symmetrical and the kurtosis values are reasonably low: 3.84, 2.98 and 2.99 at the three intervals. The average kurtosis of 3.84 over DG's last 5 sessions at 1500 msec is high, but from examination of the graph of kurtosis from each session I consider this random variation, undisturbing in a statistic known to be highly variable.

There is no support in these data for the hypothesis of a transition in timing mechanisms in the neighbourhood of 550 msec. That

idea has waxed and waned in popularity over the years, most recently being supported by scaling data from Michon (1967) and, I think more strongly, by Kristofferson's (1976) synchronization results.

I do not conclude from these data that there are never transitions at 550 msec. I believe that people can time intervals in many different ways, and some of those may not be possible at very short intervals. The point, though, is that a transition is not <u>necessary</u>. If there is a transition, it reflects a shift in strategy, not a boundary condition on ability. DG's performance at the long intervals is less variable than any reported previously in the temporal literature, and on good days GH's performance was not much worse. It takes work, or at least much practice, to use the precise timing mechanisms at the longer intervals that may be more readily available at the shorter ones, but they are available at the longer intervals nevertheless.

(2) Equivalance of visual and auditory synchronization: The variances of DK and IS were constant, or very nearly so, as were the symmetry and kurtosis coefficients. The variances were low and compare well with those reported by Kristofferson (1976) and Brewster (1983) for auditory synchronization in this range. The key difference is that DK and IS needed a great deal of practice before they showed the same trends visible in the data of excessively practiced, but not this excessively, auditory synchronization subjects.

Hopkins (1982; Hopkins and Kristofferson, 1980) showed that synchronization performance with tactile stimulation is also equivalent to auditory performance. This equivalence across three modalities

suggests to me that there is a common central mechanism that can be used for timing.

(3) <u>Continuously</u> <u>adjustable</u> <u>deterministic</u> <u>delays</u>: The finding of distributional (shape and variance) constancy actoss a relatively wide range of intervals suggests that the added timing delays at the long end of the range are produced by a deterministic mechanism. The finding of distributional constancy over very small changes in the . interval suggest a continuously adjustable mechanism. Both findings are well established with auditorily bounded intervals. Hopkins' finding of identical auditory and tactile synchronization variances suggests that the same will be true for intervals marked by tactile stimuli. The data of IS and DK establish continuous adjustability and deterministic timing in the visual modality as firmly as they are established in the auditory modality. (That there are convincing demonstrations in the auditory case in two further paradigms, speeded and very long term non-speeded discrimination, does not make the auditory case stronger but instead makes it more general.) Neither characteristic of timing had been strongly demonstrated for visually bounded intervals previously. Given the number of differences reported between visual and auditory timing and time perception, this result is neither obvious nor trivial.

Kristofferson (1976; Hopkins and Kristofferson, 1980) has argued that continuous adjustability is achieved via recycling of information at the afferent level. Auditory afferent latency variance is generally accepted to be negligible. On the grounds of equivalent variability, he and Hopkins have argued that tactile latency variance

must also be negligible. On the same grounds we could conclude that visual latency variances are negligible. Alternatively, and despite the difficulties this poses for quantal counting models, continuous adjustability may be a property of a central mechanism that is independent of input modality. I prefer the latter hypothesis, though I have no grounds for insisting upon it. With respect to quantal counting models, these have been most strongly supported by discrimination data which show stepped variances and which suggest, as the basis for each step, a maximum count of 16q. The hypothesis of a maximum of 16 does not fare well in synchronization, in the present data or in Hopkins' (1982). It is not at all clear why the maximum possible count should vary across tasks.

The hypothesis that I say is "suggested" by the present data is that we have seriously overestimated minimum reaction time. Minimum mean RT may be shorter than 100 or 105 msec. Further, with practice, moderately intense stimuli may be processed as quickly as extremely intense stimuli. There is little or no support for either suggestion in the RT literature, and there is much data to indicate that they are wrong.

I <u>think</u> that there were some very fast reactions in Experiment 3. There weren't any slow ones. If the subjects reacted to P1, their mean RT was short, perhaps as short as 85 msec, which is probably beginning to push the physical limits imposed by neural conduction times (I often hear 60 - 65 msec suggested as a value for the auditory system. Visual latencies are longer.)

I would be more inclined to press the argument that minimum RT -

has been overestimated if I had been able to develop a convincing model for the subjects' performance in Experiment 3. The negative skew of their latencies (from WS) fit with none of my notions and I eventually abandoned the attempt. Making the highly controversial assumption that mean RTs of 85 msec are possible bought little explanatory power. I see no alternative to this assumption (though 85 msec may be a bit low in any case), but my inability to arrive at an alternative does not imply that there is none.

The difficulty arising from the present data is that the kurtosis coefficients, from all of the experiments, are too high for Hopkins' (1982) model. I call the hypothesis of a triangle plus logistic distribution Hopkins' because he expressed it most formally and tested it. Kristofferson (1976) proposed much the same idea. If we accept that the logistic variance is motor variance, the kurtosis values suggest that motor variance may be as high as 250 mSec² in this task, after extensive practice. Previous results (such as Wing's, 1973) indicate that motor variances should be between 10 and 50 msec². The average of Hopkins' estimates for synchronization was close to 25 msec². A motor variance of 250 msec² is not plausible for this task.

It is unfortunate that the only other report of Kurtosis values comes from Hopkins (1982), whose subjects' latency variances were so low that almost any estimate of motor variance would be low. Kristofferson (1976) and Brewster (1983) described the shapes of their distributions with graphs and with tables that collapsed latencies into 5 msec bins. I have estimated the kurtosis coefficients from one table

of Kristofferson (1976), which provided four grouped latency distributions for one subject, GH, who later participated in Hopkins' research. (This is not Experiment 1's GH.) My estimates cannot be exact because of grouping, but they should not be too far off.

Table 8

Distributional Statistics for Subject GH

Interval 7		Variance (Grouped)	Variance (Reported)	Symmetry	Kurtosis	Logistic Variance	
A.	Krist	offerson ⟨i	, 976)				•
	· .						1-
175	msec	136	126	20	3.42	87	
310	msec	°106	112	05	2.97	42	
360	msec	108	103	.03 *	3.21	58	
405	msec	129	125	.06	2.90	45	
в.	НорК і	ns (1982) -	·		-		
460	msec.		41	80	2.87	10 .	

The values from Hopkins (1982) are from his Tables 12 and 13. An estimate of logistic variance based on the kurtosis would be slightly higher than his value of 10, based on minimum chi-square (as often as not my estimates from his data are lower than his), but the conclusion would not be changed by this, nor by selection of any other comparison data from Hopkins. As the variance of the same subject increases, so does the "motor" variance, all the way up to 87 msec². The same difficulty appears to be present in Kristofferson's (1976) data as is present in the data reported here.

I should also note that the autocorrelations of GH, not measured in the present data, were insignificant. This is not the reasop^{re}for the high kurtosis values in Kristofferson's (1976) data, and

it is probably not the reason for them in the present data. I doubt that the small foreperiod correlations added much to the present data's Kurtosis estimates either.

On the grounds of implausibly high estimates of motor variance, especially in Experiment 1, I think we should reject Hopkins' model.

I do not think that we should throw out quantal theory on the basis of the failure of the model, nor do I think that this failure is particularly damaging to the theory. The findings of distributional constancy across wide ranges of duration, and of consistent values of q (for spectacular examples, see Kristofferson (1983) for various data from the same subject) are the findings that I consider most supportative of the theory. The possibilities that the logistic (or the normal) is the wrong motor distribution, or that there is some further source of variability in the task, not a large one but with a thick tail, do not seem unreasonable for synchronization.

I do not have an alternative model to propose, nor was my intention in this research to search for one. My intention was to show that whatever model applies, it will apply equally to long and short interval timing, whether the intervals are bounded by visual or auditory markers.

Reference Notes

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Footnotes

1. One issue often discussed in the context of time perception is •whether there is such a thing as time perception. I don't think that this issue can be resolved with data and I have lost patience with it. It will not be discussed in the body of the thesis. However, the considerations component to this have had an impact upon the stated w positions of many authors in the field. To be fair to those positions, I should address a few points here.

An internal measure of the duration of an interval can be taken without being translated by the observer into a sensation or perception of that duration. This is especially evident in some of the cognitive positions. The measure of an interval is based on the amount of information that the observer remembers having processed during the interval. The judgment of the interval's length can be done solely on that basis, without at all involving a subjective experience of duration. Similarly, if the measure of an interval is the number of pulses counted within it, the count could be used without reference to a subjective impression of anything other than the count. There may or may not be a duration percept and if there is one it may or may not be epiphenomenal.

At the purely formal level, it doesn't matter in many theoretical positions whether or not duration perceptions are assumed. We could agree that a poisson process describes the characteristics of
a counting mechanism and that discrimination judgments are based on the count without deciding whether the count is translated into a duration percept or not. (The count may or may not be consciously experienced) and the conscious experience associated with it may or may not be a duration percept <u>per se</u>.) The mathematics will not change as a function of this. The same judgment theories (eg. Signal Detection) Theory) can be applied in either case. Experimenters can take a neutral position and avoid the perceptual issue when describing their formal models, or they can take either side, without limiting the utility of their ideas about mechanism for theorists holding opposite views on the perceptual issue.

An the body of the thesis, I will refer to internal clocks mechanisms which give rise to subjective experiences of duration, i.e. to duration percepts. On introspective grounds, I believe that I have duration percepts and that they are no Tess immediate or "perceptual" than percepts of size or color. More strictly, internal clocks are mechanisms which yield information that can be used as a basis for judgments about duration. The subjective experience associated with this information is a subject of controversy.

To keep the record straight on the positions of the main theorists cited, I should note that Creelman (1962) did not explicitly claim that his poisson count was translated into a duration percept. He merely claimed that the count was the basis of duration discrimination judgments. Stroud (1956) talked about subjective "dates" of events, but avoided the pssue of subjective duration

altogether. Kristofferson's (1977) real-time criterion theory explicitly denied the need for a duration perception as a basis for discrimination judgments in the single stimulus situation. 2. Latency variance could remain constant despite an increasing variance clock if the motor and timing variances were negatively correlated to a degree that changed in just the right way as clock variance changed. Kristofferson (1983) has wondered how such a hypothesis would address quantal steps in variance. I wonder why the response distributions don't change shape as the timing variance and timing - motor covariance change with duration. I don't consider this alternative to deterministic timing credible and to the best of my knowledge it has no advocates, but it is not logically impossible. This does not imply that there are no quantal delays in the system in naturalistic situations. Some components of even the most complex stimulus-response chains might still be timed internally using a quantal clock. My point is that there will be very few cases in which the main source of timing variance in the stimulus-response chain will be quantal.

4. The raw data from some sessions was lost due to unreadability of PDP-B paper tapes, and, less frequently, to disk problems of various types on the Apple. It is impossible to estimate symmetry, kurtosis, etc., from these data, so these values could not be included in the averages. Variances of the nontruncated data are available for these sessions, from the printouts obtained at the end of each session, but I excluded these from the averages as well, feeling that all averages reported together should be based on the same sessions' data.

Accordingly, the statistics are usually averaged across the last 5 sessions run at a given interval, but if some (rarely more than one) of these sessions were lost, the averages are based only on the remaining sessions. Similarly, if there were fewer than five sessions, the averages are across all consecutive sessions at this interval for which the raw data are still available. The tables in Appendix 1 list the intervals, means and variances for the lost sessions. It is evident in these tables which sessions' raw data were lost because no other statistics of the data are reported for them.

5. I have not performed this analysis for data from Experiment 3, which included P1-P2 intervals in the 85 - 150 msec range. The subjects often responded on the basis of time elapsed since the warning stimulus (WS), so the "real" latencies probably range from 1085 - 1150 msec. If they timed from WS onset rather than offset, these latencies would be 1185 - 1250 msec. The subjects did not always time from WS. Sometimes the actual latencies may be in the 85-150 msec range. In short, it is impossible to tell what the "real" mean latencies were, and, accordingly, it is impossible to express variance as a function of these latencies.

6. I do not imply that subjects adopt this strategy consciously (though some probably do). Hopkins (1982) noted that his subjects responded on the basis of time since their trial initiation response, ignoring visual P1 completely, but that they felt they were using P1, not the response. As a past subject in speeded tasks I will report that it can be very hard to tell what stimulus is controlling behavior. With practice the task become rather automatic and one doesn't think much

about how it's being done. One simply does what appears to work best, and faster is, in the reaction time case, better. Standard operant conditioning principles, assuming that shorter RTs are more reinforcing (they are, or were to me), could probably account for the development of such a strategy on a wholly unconscious level.

7. I say little in the text about duration illusions but I think that synchronization is a natural procedure to use to study them. Studies involving direct comparison of a filled and an empty interval are open to the critical suggestion that the filled interval is reported longer not because it was perceived as longer but because it is remembered as longer. The effect is on memory, not on perception. There may well be effects on memory, and many other biasing effects, but I doubt that these fully explain the illusion. Production tasks, such as synchronization, seem less vulnerable to such criticisms. Further, once a subject has stabilized in synchronization, examining the differences in response latencies with slightly more complex stimuli offers some hope of obtaining quantitative measures, in msec, of the magnitude of the illusions. If there are shifts in other aspects of the response distributions we may be able to develop better models of the temporal fariation associated with the processing of more complex stimuli. In the introduction I suggested that this should be a goal of the cognitive temporal theorist. Synchronization may provide a means.

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I find the present research exceedingly mechanistic and reductive. It is interesting to model the percepteion of identical stimuli in strictly mechanistic terms, and it is useful to know that the same mechanism is probably involve over a fairly wide range of

modalities and durations. But my biases are such that I regard this as a very useful baseline. Many different types of cues are involved in percepts of size and distance (see, eg., Sibson, 1950 and Gregory. 1966). I would be amazed if the same were not true for duration. I regard perception as an active, dynamic process, and percepts as the result of unconscious inferences drawn from as many sources of information as are available in the task.

Holway and Bering (1941) conducted a classic study of the perception of size in which they recorded observers' judgments under impoverished stimulus conditions (visual angle of the stimulus being the only available cue), and under various other more complex (and more naturalistic) conditions in which further information about stimulus size (information about its distance) was available. The research associated with quantal theory, including the present research, has provided a characterization of duration perception under impoverished stimulus conditions that Holway and Boring may not have imagined possible in the visual case. It is not a perfect characterization, as evidenced in the present data, for example, by the overly large Kurtosis values, but I find it remarKable nevertheless. It is hardly unreasonable to seek still better characterizations of the mechanismsinvolved in the well-practiced processing of strictly empty intervals, but if I were continuing in the study of temporal perception I would instead search for ways to use what we know now as a springboard for examining how people deal with more complex stimulus configurations, as they do every day.

8. The complexities of the speeded discrimination data (Brewster,

1983; Kristofferson, 1983) have very recently led Kristofferson (1983) to state that we may have to abandon the idea of a single quantal "clock", and to suggest that there may be more than one q level operative in the nervous system at the same time.

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APPENDIX 1

Summany statistics for each subject and each session.

The column headings are abbreviated. In case of confusion, they mean:

SESS: Session dumber

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NUMBER TRIALS: Number of trials included (excluded in brackets) in analysis for that session

MEAN FP: Mean foreperiod duration

MEAN RL: Mean response latency

VARIANCE OF RL: Response latency variance (from P1)

VARIANCE OF RL+FP: Response latency variance (latency measured from warning stimulus)

VARIANCE OF FP: Foreperiod duration variance.

CORRELATION OF RL, FP: Correlation of response latency (from P1) and foreperiod duration

CORRELATION OF RL+FP,FP: Correlation of response latency (from WS) and foreperiod duration

ROOT_B1 OF RC: Symmetry coefficient of response latency distribution (from P1)

KURTOSIS RL : Kurtosis of response latency (from P1) distribution . KURTOSIS RL+FP: Kurtosis of response latency (from WS) distribution.

For a discussion of timing from WS rather than from P1, see Chapter 4.

DG AUDITORY

SESS	NUMBER	M5an FP	Synch Int	MEAN RL	VARIANCE OF RL RL+FP FP	CORRELA RL,FP RI	TION OF L+FP,FP	ROOT B1 DF RL RL+FP	KURTOSIS (B2) RL RL+FP
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2.650 2.559 3.267 3.273 3.466 3.454 3.737 3.737 2.816 2.821 3.198 3.151 3.297 3.297 2.913 2.955 4.500 4.590 3.080 3.187 3.174 3.155 2.929 3.046 5.760 5.573 3.573 3.937 3.174 3.155 2.929 3.046 5.760 5.573 3.553 3.930 3.098 3.301 2.902 3.077 2.893 2.897 3.338 3.444 3.204 3.071 3.710 3.533 3.078 2.976 3.511 3.433 2.684 2.655 3.386 3.411 3.078 2.966 2.791 2.633 3.057 2.977 2.773 2.833 3.067 3.05 2.885 2.822 2.807 2.61</td></t<>	-0.072 0.037 -0.003 0.105 0.017 -0.020 0.021 0.028 -0.087 0.153 0.051 -0.030 0.059 -0.012 -0.020 -0.021 0.020 -0.021 0.029 -0.012 -0.020 -0.021 0.020 -0.021 0.020 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 -0.023 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0.177 \\ 0.062 & 0.200 \\ 9.120 & 0.123 \\ 0.140 & 0.064 \\ 0.302 & 0.301 \\ -0.177 & -0.113 \\ 0.113 & 0.177 \\ 0.056 & 0.081 \\ 0.165 & 0.162 \\ 0.109 & 0.086 \\ 0.263 & 0.219 \\ -0.039 & -0.096 \\ 0.263 & 0.219 \\ -0.039 & -0.096 \\ 0.102 & -0.131 \\ 0.102 & -0.131 \\ 0.102 & -0.131 \\ 0.102 & -0.131 \\ 0.97 & 0.294 \\ \end{array}$	2.401 2.426 2.597 2.605 2.611 2.416 2.402 2.403 2.625 2.578 3.239 3.203 3.202 3.178 2.650 2.559 3.267 3.273 3.466 3.454 3.737 3.737 2.816 2.821 3.198 3.151 3.297 3.297 2.913 2.955 4.500 4.590 3.080 3.187 3.174 3.155 2.929 3.046 5.760 5.573 3.573 3.937 3.174 3.155 2.929 3.046 5.760 5.573 3.553 3.930 3.098 3.301 2.902 3.077 2.893 2.897 3.338 3.444 3.204 3.071 3.710 3.533 3.078 2.976 3.511 3.433 2.684 2.655 3.386 3.411 3.078 2.966 2.791 2.633 3.057 2.977 2.773 2.833 3.067 3.05 2.885 2.822 2.807 2.61
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SESS	NUMBER TRIALS	Mean FP	Synch Int	M5an RL	VARIANCE OF RL RL+FP FP	CORRELAT RL,FP RL	ION OF +FP,FP	ROUT BI OF RL RL+FP	KURTOSIS RL	S (B2) RL+FP
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81 82 83 84 85 86	300 300 299 (1) 300 300	900 900 900 900	1650 1625 1600 1600 1600 1600	1651 1627 1599 1602 1607 1599	758 663 702 86 822 879 81 825 913 86 850 946 91 842	6 -0.100 1 -0.048 4 0.006 1 0.009	0.254 0.258 0.310 0.319	0.017 0.084 0.051 0.137 0.328 0.230 0.311 0.290	2.871 2.734 2.991 3.498	2.920 2.606 2.961 3.359
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SESS	NUMBER MEAN TRIALS FP	SYNCH MI INT I	ean var RL RL	RL+FP FP	CORRELATION OF RL,FP RL+FP,FF	ROOT B1 OF RL RL+FP	KURTDSIS (B2) RL RL+FP
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128 129 130 131 133 135 137 137 137 137 137 137 137 137 137 141 142 145 147 149 151 234 155 155 155 155 155 155	300 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900	1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 1500 1 <td< td=""><td>499 504 500 684 506 581 504 503 504 504 505 507 500 524 500 524 500 524 500 524 500 524 500 524 500 523 501 453 502 411 502 413 502 417 501 353 502 417 501 348 501 446 500 382 498 316 500 517 500 510 500 510 500 510 501 372 502 369 501 379 502 509 501 403 502 509 503</td><td>712 99 664 102 514 89 589 90 622 84 634 97 630 88 691 87 484 91 524 88 424 91 530 108 453 86 530 108 454 73 641 83 459 89 421 87 602 92 630 79 630 79 440 90 471 108 579 92 488 81</td><td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td><td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td><td>3.853 3.969 2.689 2.653 3.430 3.649 2.724 2.739 3.413 3.109 2.659 3.011 2.842 2.763 3.835 3.949 3.085 3.096 3.698 3.875 3.440 3.507 3.006 2.762 2.711 5.470 2.961 2.974 4.287 4.264 2.951 2.763 2.647 2.950 2.849 2.911 3.574 3.631 2.915 3.100 4.872 3.778 3.397 3.363 6.230 4.210 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690</td></td<>	499 504 500 684 506 581 504 503 504 504 505 507 500 524 500 524 500 524 500 524 500 524 500 524 500 523 501 453 502 411 502 413 502 417 501 353 502 417 501 348 501 446 500 382 498 316 500 517 500 510 500 510 500 510 501 372 502 369 501 379 502 509 501 403 502 509 503	712 99 664 102 514 89 589 90 622 84 634 97 630 88 691 87 484 91 524 88 424 91 530 108 453 86 530 108 454 73 641 83 459 89 421 87 602 92 630 79 630 79 440 90 471 108 579 92 488 81	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	3.853 3.969 2.689 2.653 3.430 3.649 2.724 2.739 3.413 3.109 2.659 3.011 2.842 2.763 3.835 3.949 3.085 3.096 3.698 3.875 3.440 3.507 3.006 2.762 2.711 5.470 2.961 2.974 4.287 4.264 2.951 2.763 2.647 2.950 2.849 2.911 3.574 3.631 2.915 3.100 4.872 3.778 3.397 3.363 6.230 4.210 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690 2.741 2.690
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SESS	NUMBER TRIALS	ngan FP	SYNCH INT	Mean Rl	VARIANCE OF RL RL+FP FP	CORRELATION C RL,FP RL+FP,F	F ROOT BI OF P RL RL+FP	KURTOSIS (82) RL RL+FP
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6H AUDITORY

SESS	NUMBER MEAN TRIALS FP	Synch Mean Int Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTDSIS (B2) RL RL+FP
121 122 123 124 125 126 127 128 129 131 132 133 134 135 136 137 138 139 141 142 143 145 151 152 153 155 156 789 161 152 155 156 161 162 163 165	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2000 1982 2025 1995 2050 2030 2075 2048 2100 2063 2100 20643 2100 2076 2005 2048 2100 2063 2100 2076 2005 2045 2050 2029 2050 2029 2050 2029 2000 1964 2000 1971 2000 1977 2000 1977 2000 1977 2000 1977 2000 1977 2000 1979 2000 1979 2000 1977 2000 1981 2000 1981 2000 1983 2000 1984 2000 1984 2000 1984 2000 1986 2000 1986 2000 </td <td>3672 3801 101 2437 2563 103 3471 3661 90 3111 3289 -96 3072 3307 89 2839 2995 87 3525 3708 81 9497 3769 95 5369 5534 82 2550 2699 89 2551 2703 88 3809 4041 99 2976 3286 112 2359 2472 81 2623 2765 98 2304 2447 76 2305 3094 91 2208 2270 69 1855 2020 75 1890 2056 100 1754 1868 74 3163 3314 83 1676 1742 90 1301 1503 83 1266 1362 91 1964 2013 83 1472<td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td><td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td><td>3.050 2.996 3.634 3.351 3.137 3.100 2.575 2.571 2.893 2.921 3.238 3.202 3.568 3.686 2.941 2.937 3.008 3.030 3.990 3.809 3.103 2.963 3.426 3.535 2.755 2.797 4.122 4.203 2.822 2.808 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.885 2.947 2.741 2.766 2.810 2.778 3.085 3.159 3.013 2.937 3.935 3.890 2.778 2.755 3.594 3.338 3.300 \$.494 3.311 3.249 3.495 3.662 3.514 3.318 3.267 2.890 2.818 2.677 2.875 2.712 2.727 2.709 2.686 4.123 3.703 3.031 3.109 3.122 3.417 3.198 2.891 3.117 3.431 3.310 3.109 3.046</td></td>	3672 3801 101 2437 2563 103 3471 3661 90 3111 3289 -96 3072 3307 89 2839 2995 87 3525 3708 81 9497 3769 95 5369 5534 82 2550 2699 89 2551 2703 88 3809 4041 99 2976 3286 112 2359 2472 81 2623 2765 98 2304 2447 76 2305 3094 91 2208 2270 69 1855 2020 75 1890 2056 100 1754 1868 74 3163 3314 83 1676 1742 90 1301 1503 83 1266 1362 91 1964 2013 83 1472 <td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td> <td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td> <td>3.050 2.996 3.634 3.351 3.137 3.100 2.575 2.571 2.893 2.921 3.238 3.202 3.568 3.686 2.941 2.937 3.008 3.030 3.990 3.809 3.103 2.963 3.426 3.535 2.755 2.797 4.122 4.203 2.822 2.808 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.885 2.947 2.741 2.766 2.810 2.778 3.085 3.159 3.013 2.937 3.935 3.890 2.778 2.755 3.594 3.338 3.300 \$.494 3.311 3.249 3.495 3.662 3.514 3.318 3.267 2.890 2.818 2.677 2.875 2.712 2.727 2.709 2.686 4.123 3.703 3.031 3.109 3.122 3.417 3.198 2.891 3.117 3.431 3.310 3.109 3.046</td>	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	3.050 2.996 3.634 3.351 3.137 3.100 2.575 2.571 2.893 2.921 3.238 3.202 3.568 3.686 2.941 2.937 3.008 3.030 3.990 3.809 3.103 2.963 3.426 3.535 2.755 2.797 4.122 4.203 2.822 2.808 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.888 2.864 3.041 3.179 2.885 2.947 2.741 2.766 2.810 2.778 3.085 3.159 3.013 2.937 3.935 3.890 2.778 2.755 3.594 3.338 3.300 \$.494 3.311 3.249 3.495 3.662 3.514 3.318 3.267 2.890 2.818 2.677 2.875 2.712 2.727 2.709 2.686 4.123 3.703 3.031 3.109 3.122 3.417 3.198 2.891 3.117 3.431 3.310 3.109 3.046
166 167 168 169 170 171 172 173 174 175 176 177 178 179 180	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2006 1978 2000 1979 2000 1982 2000 1982 2000 1977 2000 1975 2000 1975 2000 1975 2000 1975 2000 1975 2000 1975 2000 1975 2000 1975 2000 1975 2000 1977 2000 1976 2000 1973 2000 1985 2000 1981	1169 1300 107 1516 1608 79 1442 1491 86 1460 1627 92 1442 1523 82 1877 2078 93 1575 1740 95 2787 2926 75 2458 2789 114 1991 2085 59 1769 1804 70 1922 2014 63 2235 2414 94 1493 1643 105 1612 1695 107	$\begin{array}{ccccc} 0.035 & 0.319 \\ 0.019 & 0.240 \\ -0.053 & 0.188 \\ 0.103 & 0.335 \\ -0.002 & 0.230 \\ 0.129 & 0.344 \\ 0.089 & 0.319 \\ 0.070 & 0.228 \\ 0.016 & 0.217 \\ 0.051 & 0.218 \\ -0.049 & 0.148 \\ 0.043 & 0.218 \\ 0.093 & 0.287 \\ 0.057 & 0.307 \\ -0.029 & 0.223 \\ \end{array}$	0.217 0.294 0.090 0.079 -0.051 -0.117 0.094 0.167 0.201 0.238 0.288 0.255 0.246 0.187 -0.034 0.041 0.111 0.147 0.153 0.186 -0.010 -0.018 -0.339 -0.232 -0.078 -0.106 0.297 0.317 0.176 0.184	3.765 3.626 3.336 3.318 2.838 2.910 2.966 3.243 3.200 3.040 3.267 3.276 3.036 2.789 3.102 3.258 2.663 2.575 3.418 3.547 2.942 2.959 4.482 4.246 3.137 3.007 2.623 2.648 2.966 2.984

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GH AUDITORY

SESS	NUMBER	mean	Synch	nean	VARIANCE OF	CORRELATION OF	ROOT B1 OF	KURTOSIS (B2)
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SESS	NUMBER TRIALS	mean FP	SYNCH I INT	'IEAN Rl	VARIANCE OF RL RL+FP FI	CORRELATIO	EN OF EP, FP	ROOT B1 OF RL RL+FP	KURTOSI -RL	S (82) RL+FP
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3.614 2.955 3.419 3.223 3.614 2.955 3.419 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.223 3.257 3.223 3.257 3.223 3.257 3.223 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.255 3.257 3.257 3.257 3.255 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 3.257 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				- -	GH AUD	ITORY			
SESS	NUMBER TRIALS	nean Fp	Synch Int	nean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS (B2) RL RL+FP	~
301 302	300 300	1000 1000	1500 1500	1491 1491	691 780 77 673 707 69	0.026 0.339 -0.082 0.233	0.204 0.262 -0.070 0.008	3.185 2.959 3.118 2.873	

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RW AUDITORY

SESS	NUMBER TRIALS	nean Fp	SYNCH INT	nean Rl	var Rl	IANCE OF RL+FP FP	CORRELA RL,FP R	TION OF L+FP,FP	ROOT BI OF RL RL+FP	KURTOSIS (82) -RL RL+FP
1 2 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 4 5 6 7 8 9 0 11 12 3 14 5 16 7 8 9 0 11 12 3 14 5 16 7 8 9 0 11 12 3 14 5 16 7 8 9 0 11 12 3 14 5 16 7 8 9 0 11 12 3 11 2 11 2 11 2 11 2 11 2 11 2	300 (3) 299 (1) 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	150 150 150 150 150 150 150 150 150 150	159 162 156 156 157 152 155 155 155 155 155 155 155 155 155	1004 315 246 629 1047 225 218 202 183 195 258 177 173 179 285 247 191 162 171 185 171 185 171 185 170 152 170 152	1084 90 393 109 319 81 715 86 102 84 287 87 260 81 255 84 230 87 272 79 290 87 272 79 290 87 272 79 298 106 245 87 248 99 331 76 339 88 280 84 259 84 259 84 242 84 303 94 243 89 369 98 278 89 225 84 243 89 369 98 278 89 225 84	-0.017 -0.082 -0.029 -0.001 -0.049 -0.146 -0.121 -0.159 -0.059 -0.055 0.072 -0.082 -0.074 -0.102 0.015 0.015 0.021 0.056 0.095 -0.070 -0.146 -0.097 -0.056 -0.070 -0.141 -0.069 -0.141 0.011	Q⇒272 0.452 0.479 0.346 0.228 0.471 0.425 0.466 0.474 0.533 0.390 0.525 0.569 0.525 0.569 0.525 0.569 0.525 0.569 0.525 0.569 0.525 0.565 0.564 0.543 0.574 0.574 0.574 0.574 0.548 0.574 0.548 0.574 0.548 0.574 0.548 0.548 0.548 0.548 0.574 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.548 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.554 0.555 0.554 0.554 0.554 0.554 0.554 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.55500000000	-0.734 -0.584 0.108 0.501 0.006 0.156 -1.635 -1.491 0.410 0.319 -0.397 0.136 -0.163 0.144 -0.249 0.115 -0.429 -0.116 -0.031 0.198 -0.256 -0.079 -0.321 0.333 0.861 2.350 -0.327 -0.123 0.423 0.912 -0.025 0.147 -1.136 -0.109 -0.077 0.252 -0.432 0.1092 -0.432 0.1092 -0.432 0.1092 -0.432 0.1092 -0.432 0.207 -0.573 -0.176 -0.230 0.276 -0.573 -0.056 -0.486 -0.081 -1.494 -0.399	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
29 30 312 33 34 35 36 37 38 40 41 42 43 44	300 300 (1) 300 (1) 300 (1) 300 (1) 300 300 300 (1) 300 (1) 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	150 150 150 150 150 150 150 150 150 150	155 155 155 150 155 157 150 157 156 157 156 151 153 151 158 152 152	171 141 182 187 155 152 277 180 152 155 142 149 222 149 222 149 222 149	248 91 241 91 279 101 243 91 275 105 370 73 281 101 242 89 228 88 259 86 231 98	0.072 -0.126 -0.034 -0.015 0.069 0.066 0.002 0.004 -0.066 0.141 -0.069	0.659 0.504 0.574 0.601 0.670 0.601 0.610 0.502 0.601 0.610 0.598	-0.099 0.860 -0.157 0.111 D.071 0.748 -0.158 0.307 -0.251 0.427 -1.004 -0.614 -0.184 0.486 -0.056 0.380 -0.278 0.478 -0.072 0.596 -0.042 0.356	3.183 6.057 3.049 3.208 3.726 4.516 3.804 3.609 3.899 4.294 7.305 5.486 4.128 4.594 3.190 3.469 3.058 4.251 3.934 4.220 3.463 3.748
45 46 47 48 50 51 52 53 55 55 55 56 58 50 58 50	300 301 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	150 150 150 150 150 150 150 150 150 150	152 155 162 159 147 152 151 145 159 149 154 155 155 155 153	168 296 183 175 172 145 143 220 160 139 141 276 185 171 137 185	392 95 283 101 260 81 276 91 242 99 249 83 303 101 264 86 227 79 224 86 375 112 301 83 305 106 257 95 265 90	0.004 -0.003 0.019 0.055 -0.004 0.103 -0.060 0.079 0.044 -0.016 -0.036 0.135 0.106 0.107 -0.040	0.496 0.596 0.573 0.617 0.635 0.526 0.526 0.623 0.623 0.607 0.516 0.631 0.668 0.686 0.550	-2.210 -0.648 -0.335 0.364 -0.471 0.116 -0.051 0.734 -0.089 0.645 -0.447 -0.046 -0.400 0.205 -0.418 0.291 0.228 0.432 -0.030 0.395 -0.938 -0.302 -0.317 0.294 -0.797 0.566 0.086 0.588 -0.343 0.144	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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RU AUDITORY

SESS	NUMBER TRIALS	mean FP	, synch , int	mean Rl	VARIANCE OF RL RL+FP FI	CORRELAT RL,FP RL	ION OF	ROOT B1 OF RL RL+FP	KURTOSIS- RL R	(B2) (L+FP
61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 879	300 (1) 300 300 300 (2) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (300 300 300 300	900 900 900 900 900 900 900 900 900 900	155 160 165 175 175 175 175 175 175 180 185 195 200 200 200 200 200	156 163 167 175 175 177 171 176 182 185 184 194 196 201 203 202 198 199 199	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.086 0.095 0.114 0.085 0.113 0.176 0.073 0.176 0.078 0.104 0.104 0.104 0.116 0.118 0.1173 0.118 0.087 0.118 0.087 0.183	0.686 0.637 0.666 0.655 0.655 0.655 0.627 0.701 0.704 0.632 0.637 0.631 0.668 0.533 0.667 0.616 0.528	-0.207 0.620 -0.200 0.543 0.063 0.538 0.189 0.561 0.091 0.671 0.065 0.561 -0.127 0.946 0.409 1.583 0.292 0.491 0.069 0.552 0.187 0.836 0.511 0.761 0.76 0.598 1.073 0.889 0.003 0.462 0.218 0.683 0.214 0.676 0.292 0.325	2.876 4 3.168 4 3.046 3 2.807 3 3.510 3 2.977 3 2.880 4 5.207 11 4.719 3 3.107 3 3.275 4 4.281 3 7.755 4 2.811 3 2.935 3 3.518 3 3.314 3	.504 .036 .477 .993 .856 .440 .887 .824 .645 .645 .965 .314 .426 .735 .264 .735 .264
70 81 82 83 84 85 88 89 91 92 92 92 92 92 92 92 92 92 92 92 92 92	300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 301 301 300 300 301 301 300 300 300 300 300 300 300 300 300 300 300 300 300 300 3	900 900 900 900 900 900 900 900 900 900	2100 2200 2200 2200 2200 200 200 200 200	210 2103 2006 2011 192 1916 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 180 1772 1772 1772 1772 1772 1772 1772 177	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{c} 0.087\\ 0.101\\ 0.100\\ 0.200\\ 0.028\\ 0.128\\ 0.008\\ 0.028\\ 0.028\\ 0.028\\ 0.028\\ 0.028\\ 0.008\\ 0.008\\ 0.008\\ 0.008\\ 0.007\\ 0.140\\ 0.037\\ 0.106\\ 0.037\\ 0.000\\ -0.028\\ 0.000\\ -0.028\\ 0.000\\ -0.028\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.258\\ 0.000\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.235\\ -0.164\\ 0.081\\ 0.000\\ -0.092\\ 0.0056\\ -0.004\\ -0.092\\ 0.0056\\ -0.004\\ -0.092\\ 0.0056\\ -0.004\\ -0.092\\ 0.005\\ -0.000\\ -0.092\\ 0.0056\\ -0.000\\ -0.092\\ 0.0056\\ -0.000\\ -0.000\\ -0.000\\ -0.000\\ -0.000\\ -0.000\\ -0.000\\ 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0.5276 0.5277 0.5277	0.207 $0.5970.542$ $0.5610.323$ $0.6650.294$ $0.5550.288$ $0.6611.083$ $1.1120.209$ $0.6420.149$ $0.6180.038$ $0.349-0.111$ $0.735-0.057$ $0.489-0.319$ $0.361-0.102$ $0.583-0.445$ $0.321-0.445$ $0.321-0.445$ $0.321-0.505$ $0.279-0.321$ $0.353-0.505$ $0.279-0.321$ $0.353-0.396$ $0.051-1.182$ $-1.037-1.694$ $-0.469-1.036$ $-0.627-1.846$ $-1.096-0.613$ $-0.365-0.974$ $-0.761-1.523$ $-0.751-0.845$ $-0.215-0.845$ $-0.215-0.845$ $-0.215-0.869$ $-0.668-0.398$ $-0.117-0.356$ $0.459-0.206$ $1.138-0.365$ $0.047-0.345$ $0.047-0.345$ $0.047-0.345$ $0.047-0.345$ $0.047-0.345$ $0.047-0.408$ $0.141-0.473$ -0.200	2.803 3.507 3.816 4.230 4.230 4.230 3.093 4.230 3.124 3.306 3.329 2.986 3.329 2.986 3.329 2.986 3.329 2.986 3.329 2.986 3.329 2.986 3.329 2.986 3.329 2.986 3.124 3.306 5.856 4.20 3.014 3.583 5.856 4.925 7.11.015 4.893 4.925 7.11.015 4.893 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 4.293 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RU AUDITORY

SESS	NUMBER	Mban	SYNCH	nean	var	IANCE OF	CORRELA	TION OF	ROOT B1 OF	KURTOS	IS (B2)
	TRIALS	FP	INT	Rl	Rl	RL+FP FP	RL,FP R	L+FP,FP	RL RL+FP	·RL	RL+FP
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4.583 3.364 4.774 4.650 4.5123 5.478 5.474 4.5123 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.514 5.474 4.5773 5.478 5.474 4.5773 5.4748 5.4748 5.6774 4.099	3.763 3.441 3.087 3.885 4.302 4.181 4.680 3.914 6.947 5.503 4.172 4.176 3.371 4.176 3.493 3.353 4.172 4.373 3.704 6.839 4.680 4.181 3.704 4.502 3.704 4.502 3.714 4.502 3.572 3.321 4.502 3.572 3.321 4.502 3.572 3.321 4.502 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 3.572 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RW AUDITORY

SESS	NUMBER	nean	synch mean	VARIANCE OF	CORRELATION OF	RCOT B1 OF	KURTOSIS (B2)
	TRIALS	FP	Int rl	RL RL+FP FP	RL,FP RL+FP,FP	RL - RL+FP	-RL RL+FP
241 242 243 244 245 246 247 248 247 253 2554 2552 2554 2557 2557 2557 2557 2557	300 300 300 300 300 300 300 300 300 300	1001 1000 1000 1001 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 10000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000	85 82 85 86 85 86 85 87 85 85 85 87 85 85 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 87 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85 85	430431 78 425 379 89 483 417 109 381 374 86 371 355 89 531 514 129 377 322 101 387 363 97 433 373 92 509 371 112 446 456 66 461 386 92 488 455 75 410 377 97 394 350 84 404 369 78 349 350 80 246 247 71 174 171 89 349 353 70 327 303 80 266 247 71 174 171 89 127 160 114 200 242 84 148 242 92 126 242 104 127 260 112 123 242 94 144 267 94 142 261 94 131 236 94 157 259 97 188 299 95 154 254 80 209 353 102 173 299 76 218 343 107 270 367 83 381 482 94 315 482 94 <t< th=""><th>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</th><th>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</th><th>3.488 2.975 3.407 4.349 3.127 2.875 3.746 3.784 3.239 3.413 4.858 6.161 3.235 3.243 3.178 3.108 3.752 3.460 3.376 3.076 3.376 3.078 3.752 3.460 3.376 3.078 3.767 3.377 3.793 3.749 10.301 9.927 3.562 3.908 3.162 2.907 4.606 3.864 5.456 4.813 3.966 4.821 3.566 4.813 3.966 4.821 3.525 5.251 4.323 4.651 5.002 3.526 5.251 4.323 4.651 5.002 3.526 4.202 5.251 4.323 4.651 5.002 3.526 5.280 3.028 4.390 3.028 4.390 3.029 7.3182 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.549 2.742 3.927 3.004 3.544 2.754 4.471 2.779 3.004 3.113 3.023 2.907 3.036 2.992 3.549 2.811 2.775 3.352 3.037 2.782 2.869 2.872 3.549 2.811 2.775 3.352 3.037 2.782 2.869 3.892 3.819</th></t<>	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	3.488 2.975 3.407 4.349 3.127 2.875 3.746 3.784 3.239 3.413 4.858 6.161 3.235 3.243 3.178 3.108 3.752 3.460 3.376 3.076 3.376 3.078 3.752 3.460 3.376 3.078 3.767 3.377 3.793 3.749 10.301 9.927 3.562 3.908 3.162 2.907 4.606 3.864 5.456 4.813 3.966 4.821 3.566 4.813 3.966 4.821 3.525 5.251 4.323 4.651 5.002 3.526 5.251 4.323 4.651 5.002 3.526 4.202 5.251 4.323 4.651 5.002 3.526 5.280 3.028 4.390 3.028 4.390 3.029 7.3182 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.276 4.044 3.251 3.282 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.182 3.229 3.549 2.742 3.927 3.004 3.544 2.754 4.471 2.779 3.004 3.113 3.023 2.907 3.036 2.992 3.549 2.811 2.775 3.352 3.037 2.782 2.869 2.872 3.549 2.811 2.775 3.352 3.037 2.782 2.869 3.892 3.819
297	300	1001	300 295 300 290 300 292 300 296	416 526 97	0.030 0.457	0.299 0.396	3.459 3.128
298	300	1000		530 692 95	0.149 0.501	-0.198 0.222	3.511 3.855
299	300	999		583 672 79	0.022 0.364	0.443 0.409	3.635 3.382
300	300	1000		481 629 92	0.133 0.499	0.261 0.451	3.472 3.756

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SESS	NUMBER - TRIALS	Mean FP	Synch Int	nean Rl	VARIANCE OF RL RL+FP FP	CORRELATION RL,FP RL+FP,	DF ROOT B1 OF FP RL RL+FP	KURTOSIS (B2) RL RL+FP
301 302 303 304 305 306 307 308 309 310	300 300 300 300 300 300 300 300 300 300	999 1000 999 1001 999 1001 1000 999 1000	300 300 300 300 300 300 300 300 300	301 296 295 287 290 282 297 297 297 297 297	473 639 80 427 574 96 465 558 67 499 623 100 511 582 64 912 1083 135 559 747 88 495 590 79 408 532 88	0.223 0.5 0.125 0.5 0.075 0.4 0.053 0.4 0.019 0.3 0.052 0.4 0.224 0.5 0.042 0.4 0.096 0.4	15 0.134 0.243 17 0.038 0.248 14 0.126 0.338 15 -0.047 0.365 16 0.443 0.302 17 0.138 0.443 18 0.179 0.256 10 0.178 0.464 10 0.138 0.464 10 0.061 0.056 10 0.015 0.199	3.089 3.203 2.869 2.947 3.547 3.657 3.070 4.115 3.653 3.191 3.131 3.053 2.843 3.342 3.582 3.246 2.705 3.110
309 310 311 312 313 314 315 314 315 314 315 314 315 314 315 314 315 314 315 322 322 322 322 322 322 322 32	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000	300 300 300 300 300 300 300 300 300 300	- 294 300 303 297 298 300 301 298 298 300 298 298 298 298 298 298 298 298 298 298	408 532 88 595 476 609 88 388 547 90 386 500 633 456 604 88 339 481 933 389 550 114 341 483 87 416 548 87 476 608 99 297 387 80 411 524 94 270 389 92 351 448 94 357 466 89 290 406 92 281 388 933 276 462 119 371 473 80 287 366 92 290 406 81 317 484 99 330 496 104 267 396 80 245 316	0.096 0.4 0.109 0.4 0.185 0.5 0.164 0.5 0.148 0.5 0.138 0.5 0.138 0.5 0.110 0.5 0.141 0.5 0.111 0.5 0.075 0.4 0.058 0.5 0.047 0.4 0.058 0.5 0.047 0.4 0.055 0.4 0.045 0.5 0.45 0.5 0.45 0.5 0.45 0.5 0.45 0.5 0.045 0.4 0.055 0.4 0.00	0.015 0.199 0.015 0.199 0.001 0.549 0.058 0.092 0.058 0.092 0.076 0.629 0.076 0.424 0.748 48 0.076 0.429 0.076 0.429 0.189 57 0.424 0.748 48 0.076 0.429 60 -0.116 0.296 0.192 0.044 0.388 0.720 1.046 0.198 0.413 66 -0.140 -0.023 57 -0.085 0.192 63 -0.062 0.862 0.031 0.248 0.052 0.077 0.067 0.092 68 -0.275 0.068 0.073 0.2240 0.573 0.179 0.510 0.439 0.021 0.067 0.924	2.705 3.110 2.681 3.456 3.278 3.895 3.003 3.074 4.285 4.905 2.948 3.804 2.765 4.038 3.159 3.646 2.950 3.300 2.916 3.698 5.371 6.129 3.476 3.938 2.584 2.819 2.815 3.233 3.021 2.921 2.606 2.849 3.763 3.573 4.045 3.920 3.099 4.454 3.203 3.072 3.499 3.252 2.858 2.861 2.972 3.251 3.205 3.180 2.972 3.251 3.205 3.773 3.160 3.965 3.344 3.455 3.618 3.624 3.072 5.043 3.703 4.017 3.583 3.773 4.045 3.920 2.972 3.251 3.205 3.773 3.160 3.965 3.344 3.455 3.618 3.624 3.072 5.043 3.703 4.017 3.583 3.773 4.045 3.920 4.528 3.205 3.773 3.160 3.965 3.344 3.455 3.310 4.696 3.038 3.018 3.151 4.190 6.373 8.812 2.770 3.325 3.907 4.411 3.240 6.839 2.972 2.855
358 359 360	300 300 300	1000 1000 999	300 300 300	302 295 295	339 455 81 253 325 79 275 402 86	0.106 0. -0.023 0. 0.131 0.	13 0.068 0.457 73 0.427 0.447 72 0.140 0.490	3.069 4.306 3.483 3.236 3.142 3.804

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•	SESS	TRIALS	FP	INT	RL /	RL I	RL+FP FP	RL,FP F	ALION OF RL+FP,FP	ROOT RL	B1 OF RL+FP	RURTOS:	IS (82) RL+FP		
	361 362 363 364 365 366 367 368 370 371 372 373 374 375 377 378 377 378 379 380 381 382 383 384 385 388 389	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 1000 1000 1000 1000 100	300 300 300 300 300 300 300 300 300 350 35	295 299 294 295 305 305 305 316 329 316 329 346 350 347 351 348 337 326 307 295 302 297 302 297 303 301 297 303 301	254 286 239 224 254 252 253 254 254 255 255 255 255 255 255 255 255	364 97 408 108 313 76 368 96 371 103 358 99 377 100 326 78 358 100 326 74 388 86 437 79 373 95 379 107 366 73 97 107 366 73 97 107 366 73 97 107 341 105 270 85 400 98 804 89 327 73 267 86 274 71 348 81 329 100	0.038 0.039 -0.009 0.166 0.046 0.074 0.056 0.137 -0.010 0.093 0.052 0.052 0.036 0.075 0.062 -0.078 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 0.025 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0.025	$\begin{array}{c} \textbf{0.549}\\ \textbf{0.548}\\ \textbf{0.639}\\ \textbf{0.586}\\ \textbf{0.586}\\ \textbf{0.586}\\ \textbf{0.536}\\ \textbf{0.536}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.547}\\ \textbf{0.586}\\ \textbf{0.515}\\ \textbf{0.515}\\ \textbf{0.573}\\ \textbf{0.585}\\ \textbf{0.5515}\\ \textbf{0.5738}\\ \textbf{0.5515}\\ \textbf{0.5738}\\ \textbf{0.558}\\ \textbf{0.558}\\ \textbf{0.558}\\ \textbf{0.521}\\ \textbf{0.528}\\ \textbf{0.521}\\ \textbf{0.5221}\\ \textbf{0.521}\\ \textbf{0.5221}\\ $	0.382 0.279 0.283 0.168 0.101 -0.012 0.298 0.044 0.101 -0.012 0.298 0.154 -0.277 0.148 0.092 -0.185 0.152 -0.061 -0.231 0.242 0.353 0.170 0.133 0.204 0.133 0.204 0.133 0.204 0.148 -0.622 0.148 -0.062	0.577 0.527 0.163 0.588 0.706 0.507 0.507 0.507 0.497 0.497 0.497 0.498 0.396 -0.082 0.474 0.396 0.298 0.254 0.298 0.254 0.254 0.254 0.5536 0.273 0.536 0.253 0.365 0.253 0.515	2.859 2.604 3.011 2.969 3.262 2.914 2.598 3.336 3.246 2.794 3.238 3.376 3.733 2.608 2.830 2.697 3.396 2.819 3.518 3.420 2.953 3.200 2.682 3.408 2.646 3.368 3.133 2.961 2.897	3.408 3.498 3.345 3.345 3.345 3.345 3.355 3.351 3.355 3.351 4.559 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 3.359 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JB AUDITORY

SESS	NUMBER	mban FP	Synch Int	MEAN RL	var RL	IANCE OF RL+FP FP	CORRELA RL,FP R	TION OF RL+FP,FP	ROOT BI RL RI	OF +FP	KURTOSI -RL	S (82) RL+FP
1234567890112345678901123456789021222245678902122222222222222222222222222222222222	300 () 300 () 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1) 300 (1)	900 900 900 900 900 900 900 900 900 900	200 200 200 200 200 200 200 200 200 200	206 201 205 209 195 206 192 203 196 202 207 208 209 203 200 200 200 200 200 200 200 205 205 205	1401 1177 849 874 810 618 641 494 680 424 545 390 418 554 466 476 326 572 518 603 326 572 518 466 379 481 421	1454 80 1300 86 975 105 970 84 908 88 695 97 732 94 590 91 763 73 468 85 635 94 471 80 516 95 532 107 606 84 633 91 549 95 566 86 677 74 448 98 718 98 605 81 529 74 464 95 511 100 519 98	$\begin{array}{c} -0.040\\ 0.058\\ 0.035\\ 0.018\\ -0.040\\ -0.005\\ 0.011\\ 0.022\\ -0.107\\ -0.008\\ 0.014\\ 0.030\\ 0.014\\ -0.075\\ 0.087\\ -0.029\\ 0.008\\ -0.058\\ -0.058\\ -0.058\\ -0.058\\ -0.058\\ -0.058\\ -0.058\\ -0.029\\ 0.008\\ -0.031\\ -0.024\\ -0.160\\ -0.008\end{array}$	0.195 0.312 0.361 0.329 0.354 0.329 0.354 0.323 0.374 0.404 0.323 0.376 0.447 0.4455 0.463 0.302 0.458 0.390 0.329 0.329 0.425 0.329 0.324 0.323 0.324 0.323 0.324 0.323 0.324 0.323 0.325 0.463 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 0.329 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5.004 4.078 8.455 4.916 2.950 4.693 2.933 3.884 3.514 3.805 4.310 3.837 3.393 3.539 2.981 3.887 3.305 4.310 3.814 4.568 3.814 4.568
29 30 31 32 33 34 35 36 37 38 37	300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900	200 200 200 200 200 200 200 200 200 200	201 202 193 203 205 194 210 205 209 195	419 451 523 380 446 379 354 439 552 513 345	551 100 590 97 577 79 482 95 566 103 469 83 441 98	0.077 0.101 -0.061 0.020 0.038 0.018 -0.029	0.495 0.494 0.312 0.462 0.461 0.438 0.438 0.444	0.374 0 0.588 0 -0.164 0 0.374 0 0.232 0 0.129 0 0.313 0	.530 .703 .019 .629 .343 .311 .096	3.720 4.845 3.450 4.359 3.813 2.772 3.294	3.693 4.484 3.768 4.368 3.514 3.137 2.960
441 444 445 447 447 447 555 555 555 555 555	300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	200 200 200 200 200 200 200 200 200 200	201 204 195 200 192 201 199 200 201 199 200 201 198 198 198 188 188 185 175 175	371 377 364 499 523 449 758 379 297 469 379 398 399 469 398 469 398 442 545 545 545 301 371	461 90 418 97 593 88 588 89 616 97 529 77 804 91 396 85 460 84 368 62 561 110 398 89 550 97 473 88 490 84 620 91 594 97 416 75 342 83 423 78	0.001 -0.118 -0.093 0.001 -0.014 0.007 -0.087 -0.131 -0.008 0.032 -0.040 -0.086 142 -0.036 -0.094 -0.035 0.042 -0.083 -0.134 -0.079	0.444 0.377 0.295 0.390 0.389 0.252 0.339 0.420 0.438 0.407 0.395 0.540 0.397 0.326 0.351 0.443 0.345 0.368 0.356	$\begin{array}{cccccccccccccccccccccccccccccccccccc$.307 .098 .141 .153 .252 .042 .119 .228 .425 .265 .089 .282 .242 .053 .289 .282 .242 .053 .718	3.688 3.446 4.012 4.924 4.925 3.990 4.341 2.792 3.356 3.530 3.404 3.489 4.076 2.836 3.958 13.050 3.456 3.928 4.497 7.380	3.580 3.528 3.556 4.601 3.790 3.471 4.072 2.810 3.373 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.673 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.675 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.655 3.6555 3.6555 3.6555 3.65555555555

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SESS ,	NUMBER TRIALS	Mean FP	Synch JNT	mean Rl	VARI RL R	ANCE OF L+FP FP	CORREL4 RL,FP F	ATION OF RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOS	IS (82) RL+FP
61 62 63 64 65 66 67 68 70 71 72 73 74 75 77 78 79 80	300 279 (1) 300 300 300 300 299 (1) 300 300 300 300 300 300 300 300 300 30	900 900 900 900 900 900 900 900 900 900	175 175 175 175 175 175 175 175 160 155 150 150 150 150 150 150 150 150	168 180 174 175 169 165 157 155 157 152 153 154 148 150 166 152 152 152 152 152	344 378 318 296 375 374 359 431 431 431 551 431 551 431 551 373 373 282 389 315	415 82 436 93 384 104 386 89 413 88 467 76 476 102 476 102 461 113 457 92 383 78 669 88 465 101 565 101 421 95 385 92 295 84 438 84 360 100 425 84	-0.032 -0.094 -0.104 0.004 -0.138 0.049 0.047 -0.137 -0.211 -0.181 0.015 0.066 -0.162 -0.138 -0.216 -0.229 -0.098 -0.152 -0.174	0.415 0.375 0.425 0.484 0.331 0.448 0.489 0.333 0.289 0.272 0.464 0.424 0.311 0.250 0.344 0.276 0.308 0.347 0.383 0.274	-0.176 0.346 -0.458 -0.017 0.177 0.425 0.507 0.640 -1.006 -0.694 -0.751 -0.405 0.190 0.831 -0.724 -0.067 -0.355 -0.086 -0.722 -0.540 -0.211 0.318 -0.310 -0.087 -0.588 -0.409 -0.821 -0.106 -0.457 -0.033 -0.135 0.039 -0.457 -0.078 -1.762 -0.979 -0.147 0.192 -0.603 -0.109	5.683 5.226 4.279 6.627 6.227 4.223 3.661 4.733 3.839 4.204 4.513 3.091 4.235 7.089 3.909 3.528 3.700 11.709 2.843 7.032	6.032 4.097 4.761 6.077 4.810 4.124 5.213 3.586 3.761 5.584 3.717 6.798 5.621 3.276 3.346 8.369 3.056 5.606
81 82 83 85 85 85 87 89 89 89 90 91	300 300 300 300 300 300 300 300 300 302 302	900 900 900 900 900 900	135 130 125 120 115 110 105 100 95 90 90	135 137 129 124 122 118 112 108 99 99 103	348 315 448 347 408 384 564 486 732 651 •778	346 96 476 84 448 91 747 79 634 90	-0.188 -0.146 -0.306 -0.132 -0.222	0.348 0.279 0.132 0.194 0.153	-0.367 0.018 -0.068 0.094 -0.782 -0.686 -0.819 -0.590 -0.975 -1.028	3.524 4.006 4.712 4.416 4.901	3.436 3.860 4.580 4.047 5.536
92 93 94 95 97 98 97 100 101 102 103 104 107 108 109 111 112 113 114 115 116 117 118 119 120	300 301 301 301 300 301 300 299 301 300 301 300 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 301 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 3	900 901 900 900 900 1000 1000 1000 1000	95 100 100 100 100 100 100 100 100 100 10	105 104 109 106 100 104 103 101 108 95 107 101 108 95 107 101 109 103 107 101 104 106 107 105 111 109 105 111	559 44 536 534 563 563 563 563 563 563 563 563	630 74 537 100 634 98 620 84 611 89 510 92 510 92 476 91 856 84 948 67 641 64 467 91 467 87 737 85 488 88 476 97 737 85 476 97 378 90 380 76 394 112 392 77 281 83	$\begin{array}{c} -0.135\\ -0.213\\ 0.004\\ -0.268\\ -0.175\\ -0.316\\ -0.316\\ -0.316\\ -0.182\\ -0.164\\ -0.181\\ -0.184\\ -0.181\\ -0.164\\ -0.195\\ -0.196\\ -0.196\\ -0.235\\ -0.196\\ -0.235\\ -0.173\\ -0.069\\ -0.173\\ -0.062\\ -0.177\\ -0.289\\ -0.162\\ -0.153\\ -0.176\\ -0.155\\ -0.120\\ -0.132\\ -0.155\end{array}$	0.208 0.219 0.397 0.090 0.208 0.208 0.204 0.260 0.150 0.082 0.152 0.152 0.181 0.327 0.233 0.188 0.228 0.324 0.353 0.271 0.355 0.318 0.358 0.358 0.358 0.358 0.358 0.358 0.358 0.358	$\begin{array}{c} -0.499 & -0.446 \\ -1.038 & -0.759 \\ -0.428 & -0.198 \\ -0.865 & -0.834 \\ -0.421 & -0.435 \\ -0.321 & 0.106 \\ -0.512 & -0.394 \\ -0.904 & -0.360 \\ -0.990 & -0.746 \\ -0.967 & -0.867 \\ -0.528 & -0.526 \\ -0.917 & -0.608 \\ -1.096 & -0.920 \\ -0.979 & -0.538 \\ -1.096 & -0.920 \\ -0.563 & 0.285 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.823 \\ -1.384 & -0.633 \\ -0.563 & 0.225 \\ -1.033 & -0.226 \\ -1.033 & -0.226 \\ -1.052 & -0.334 \\ -1.058 & -0.334 \\ -1.633 & -0.954 \\ -0.959 & -0.347 \\ \end{array}$	4.376 6.390 2.915 4.302 3.188 6.655 3.210 5.838 8.321 4.151 4.417 5.111 6.493 7.420 8.594 5.289 5.654 7.232 8.749 5.289 5.654 7.050 5.738 5.834 6.518 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074 5.074	4.377 5.897 3.045 4.680 3.342 4.949 3.340 5.374 4.053 4.053 4.053 4.053 4.053 3.367 5.311 8.287 3.367 5.311 8.287 5.344 5.745 6.806 3.956 4.569 5.038 4.702 5.263 3.707 6.653 4.927

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JB AUDITORY

SESS	NUMBER TRIALS	mean FP	SYNCH INT	Mean Rl	Vari RL F	IANCE OF RL+FP FP	CORRELA RL,FP R	TION OF L+FP,FP	RDOT RL	B1 OF RL+FP	KURTOSI -RL	(B2) RL+FP
$\begin{array}{c} 121\\ 122\\ 123\\ 124\\ 125\\ 126\\ 127\\ 128\\ 127\\ 128\\ 129\\ 130\\ 131\\ 132\\ 133\\ 135\\ 136\\ 137\\ 138\\ 139\\ 141\\ 142\\ 143\\ 144\\ 145\\ 151\\ 152\\ 153\\ 156\\ 156\\ 156\\ 156\\ 166\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 167\\ 168\\ 168\\ 168\\ 168\\ 168\\ 168\\ 168\\ 168$	300 301 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 3	1000 1000 1000 1001 1000 1000 1000 100	100 100 100 100 100 100 100 100 100 100	110 107 108 103 107 112 104 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422 71 381 99 370 88 357 75 295 78 320 96 295 70 426 89 328 97 543 120 429 102 432 71 368 72 432 71 368 72 409 106 332 74 425 78 365 101 419 78 339 89 233 91 272 100 212 97 275 97 226 88 222 74 181 67 226 85 251 84 194 83 226 85 251 84 194 83 241 128 243 90 241 114 289 89 308 86	$\begin{array}{c} -0.085\\ -0.153\\ -0.125\\ -0.205\\ -0.146\\ -0.032\\ -0.082\\ -0.082\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.190\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.121\\ -0.124\\ -0.218\\ -0.195\\ -0.214\\ -0.218\\ -0.195\\ -0.214\\ -0.218\\ -0.195\\ -0.214\\ -0.218\\ -0.195\\ -0.214\\ -0.218\\ -0.195\\ -0.045\\ -0.140\\ -0.040\\ -0.040\\ -0.040\\ -0.040\\ -0.040\\ -0.040\\ -0.045\\ -0.124\\ 0.077\\ -0.028\\ 0.045\\ -0.124\\ 0.077\\ -0.028\\ 0.045\\ -0.160\\ 0.017\\ -0.028\\ 0.017\\ -0.028\\ 0.0165\\ 0.017\\ -0.053\\ 0.037\\ 0.037\\ 0.039\\ 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4.955 3.759 4.095 3.755 4.085 3.667 3.155 4.985 3.497 3.360 5.197 3.667 3.360 5.197 3.360 5.197 3.667 3.360 5.197 3.360 5.197 3.667 3.360 5.197 3.667 3.360 5.197 3.667 3.360 5.197 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 3.667 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170 171 172	300 300 300	1000	· 250 · 250 250	257 251 257	289 239 248	314 87	-0.041	0.490	0,428	0.460	3.841	4.061
173 174	300 300	1000	250 250	253 251	290 194	.382 84	0.028	0.492	0.533	0.670	4.206	3.780
175 176 177 178 179 180	300 300 300 300 300 300	1001 1000 1000 1000 1000 1000	260 270 280 290 300 250	268 272 282 292 300 254	216 254 310 290 317 207	286 101 299 92 349 85 390 93 429 99 264 75	-0.104 -0.154 -0.140 0.020 0.037 -0.074	0.504 0.414 0.360 0.505 0.513 0.467	-0.107 -0.117 0.002 -0.005 0.085 0.265	0.109 0.186 0.082 0.376 0.169 0.192	2.923 2.870 2.628 2.747 2.945 3.375	3.371 2.905 2.594 3.344 2.975 2.920

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JB AUDITORY

SESS	NUMBER TRIALS	nean FP	SYNCH	nean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS (82) RL RL+FP
181 182 183 184 185 186 187 188 189 199 199 199 200 202 203 204 207 209 211 212 213 215 215 220 222 222 222 222 222 222 222 222 22		1001 999 1000 1001 1000 1001 1000 1001 1000 1001 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 10000 100000 10000 10000 1000000		3074 30077 30233073075 30233023014 301122305589 30233023009 302330229020 3009330229020 3009330229020 3009330229020 3009330229020 3009330229020 3009330223023302230 300555555555555555555555	365 475 119 373 458 81 287 340 81 290 329 69 295 349 101 236 320 92 328 399 81 237 299 79 296 381 94 250 334 104 229 287 80 285 397 135 258 312 81 205 296 108 278 333 68 194 277 113 337 409 101 332 429 97 217 285 71 216 299 84 216 314 103 230 302 104 214 287 79 214 287 79 214 287 79 214 287 79 214 287 79 214 287 73 185 277 80 247 317 69 191 260 97 214 297 73 185 277 80 247 317 69 298 267 81 200 313 95 208 321 107 162 247 92 208 296 113 162 247 92 208 296	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	4.084 4.555 4.585 4.370 2.748 3.357 3.215 3.244 3.119 3.276 2.865 3.395 3.733 3.632 2.746 3.092 2.746 3.032 2.746 3.032 2.746 3.952 2.746 2.981 3.229 3.746 2.981 3.332 2.942 3.652 2.746 2.981 3.229 3.746 2.981 3.332 2.942 3.652 2.748 2.801 3.229 3.746 2.788 2.801 3.057 4.446 2.771 2.633 2.949 3.151 3.255 3.711 3.801 3.251 3.255 3.726 3.255 3.726 3.277 2.854 4.068 4.452 3.973 3.282 2.920 2.877 3.623 3.844 4.068 4.452 3.953 3.282 2.854 4.074 3.2953 3.282 2.854 4.074 3.2854 4.074 3.645 3.771 2.854 4.074 3.122 5.62 2.894 2.621 3.645 3.792 3.645 3.792 3.646 3.792 3.646 3.792 3.642 3.622 3.643 $3.$

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JE AUDITORY

SESS	NUMBER TRIALS	mean FP	SYNCH INT	nean Rl	VAR) RL	ANCE OF RL+FP FP	CORRELA RL,FP F	TION OF L+FP,FP	ROOT RL	B1 OF RL+FP	KURTOSI 'RL	S' (B2) RL+FP
241 242 243 244 245 246 247 248 250 251 252 253 254 255 256 257 258 259	300 300 300 300 300 300 300 300 300 300	1000 1001 1000 1000 1000 1000 1000 100	300 300 310 320 330 350 350 350 350 350 350 350 350 35	303 301 300 318 323 335 349 357 357 357 357 355 348 329 329 329 329 329 329 329 329 329 329	181 180 167 192 203 175 203 175 203 175 203 217 203 211 180 210 212 222 239 239 217 226 7 221 184 195	252 71 266 117 248 -72 243 88 259 84 274 101 250 80 354 92 280 93 261 91 283 83 317 85 296 83 352 97 284 106 345 135 269 90 246 83 231 81	0.002 -0.108 0.045 -0.106 -0.079 -0.105 -0.023 0.081 -0.084 -0.039 -0.041 0.052 -0.002 -0.002 -0.045 -0.002 -0.149 -0.087 -0.177	0.533 0.573 0.574 0.509 0.502 0.517 0.547 0.502 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.509 0.502 0.502 0.509 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.502 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508 0.508	-0.008 -0.033 -0.031 -0.207 -0.274 -0.211 -0.188 -0.324 -0.104 0.105 -0.198 0.103 -0.198 0.103 -0.198 0.345 -0.107 -0.121 -0.432 0.186 -0.083	0.044 0.402 0.394 -0.031 -0.143 -0.119 0.191 0.090 0.159 0.524 0.186 0.702 0.091 0.833 0.913 -0.311 0.216 0.594	3.460 2.992 2.843 3.114 4.385 3.723 2.949 3.616 2.443 3.386 2.959 3.508 2.813 3.416 3.941 2.976 3.948 2.929 5.286	3.293 3.822 3.912 3.195 3.649 3.250 2.625 3.287 2.868 3.963 2.877 4.979 2.827 4.979 2.827 4.925 4.821 3.777 3.777 3.24(7.297)

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KC AUDITORY .

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SESS	NUMBER TRIALS	mean FP	SYNCH INT	nean Rl	VARIANCE OF RL RL+FP F	CORRELA P RL,FP I	ATION DF RL+FP,FP	* ROOT B1 DF RL RL+FP	KURTOS	IS (B2) RL+FP
t 1 2 3 4 5 6 7 8 9 10 112 3 4 5 6 7 8 9 10 112 3 14 5 6 7 8 9 10 112 3 14 15 16 7 8 9 21 22 3 4 5 6 7 8 9 10 112 3 14 15 16 7 8 9 20 21 22 3 4 5 6 7 8 9 20 20 20 20 20 20 20 20 20 20 20 20 20	305 (2) 315 (2) 302 (1) 307 304 300 (2) 307 303 208 (2) 301 (2) 303 (2) 300 (2) 301 (2) 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	150 150 150 150 150 150 150 200 200 200 200 200 200 200 200 200 2	160 142 186 155 160 147 178 140 142 201 198 195 189 202 199 203 207 204 197 197 190 195 192 203 196 197 190 195 192 202	3408 3402 7 4837 4930 10 4537 4532 9 3127 3144 6 2593 2610 8 2258 2196 8 2323 2295 6 2924 2965 6 2945 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 2045 1994 8 1048 115 8 1075 1195 11 1289 1382 10 1437 1499 9 1203 1316 10 1120 1185 6 771 835 9 1061 1110 9	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.967 0.138 0.059 0.097 0.108 0.035 0.063 0.126 0.037 0.135 0.285 0.278 0.247 0.316 0.260 0.208 0.278 0.246 0.208 0.247 0.216 0.228 0.244 0.228 0.244 0.2281 0.225 0.321 0.285 0.349 0.294 0.297 0.294 0.297 0.2980	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	9.313 5.716 4.354 4.172 4.746 4.867 4.055 3.924 4.004 9.419 5.052 4.126 4.933 3.950 3.188 4.015 4.336 3.379 5.704 4.762 8.561 3.337 3.764 4.409 5.853 7.089 3.875 4.582 3.765 4.582 3.765	9.134 5.588 4.229 4.171 4.459 4.657 3.735 3.949 3.400 9.384 4.899 4.198 4.921 3.655 3.635 3.635 3.635 3.635 3.635 3.635 3.635 3.635 3.635 3.64 5.478 8.304 5.478 8.304 5.168 3.655 4.201 5.012 6.843 5.168 4.039 4.353
31 323 334 35 36 37 38 39 40 42 43 44 45 44 45 47 89 51	300 301 299 (1) 300 300 300 300 300 (1) 300 300 300 300 300 300 300 30	900 900 900 900 900 900 900 900 900 900	200 200 200 200 200 200 200 200 200 200	196 198 189 196 196 196 197 207 193 209 204 200 197 198 189 189 189 183 201 200 194 189	732 1144 937 1052 9 1055 1056 7 1055 1069 9 729 788 9 706 777 9 867 907 9 915 1018 8 958 950 9 602 705 11 529 609 9 374 434 8 523 553 8 539 610 7 563 672 9 791 868 11 499 891 616 676 8 733 759 9	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.333 0.133 0.169 0.282 0.305 0.233 0.318 0.341 0.365 0.365 0.365 0.265 0.340 0.403 0.303 0.303	-0.547 -0.266 0.180 0.121 -0.126 -0.033 -0.288 0.111 0.581 0.486 0.313 0.270 -1.335 -0.946 1.143 1.201 0.228 0.321 0.155 0.318 -0.022 -0.033 -0.103 -0.071 -0.562 -0.460 0.286 0.158 -0.361 -0.119 0.082 0.115 -0.678 -0.420	6.918 5.052 2.754 6.850 3.745 4.461 11.614 8.517 4.268 3.398 3.316 3.029 7.908 4.964 3.772 4.428 7.162	6.494 5.120 2.766 4.615 3.484 3.951 9.534 8.299 3.589 3.589 3.589 3.589 3.589 3.775 6.898 3.738 3.938 3.779 4.252
52 53 54 55	300 (1) 300 300 (1) 300 (1) 300	901 900 900	200 200 200 200	195 184 187 192	562 623 10 609 687 10 535 578 10 591	-0.127 -0.087 0 -0.044 -0.128	0.323 0.339 0.300	-0.878 -0.420 0.571 0.619 1.058 1.080 -0.334 -0.262	7.162 3.837 8.146 7.922	6.252 4.441 7.013 7.077
57 58 59 60	300 300 299 (1) 300	900 900 900 900 900	200 200 200 200 200	196 196 199 183 186	536 1011 9 560 628 10 -734 799 9 1115 1154 8 722 835 8	B -0.0/1 1 -0.069 1 -0.052 3 -0.073 6 0.056	0.243 0.335 0.288 0.197 0.372	0.715 0.708 0.132 0.025 -1.072 -0.750 -0.305 -0.194 0.386 0.461	5.617 4.391 11.108 5.274 3.725	5.329 4.179 9.956 5.558 4.143

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SESS	NUMBER TRIALS	nean Fp	SYNCH INT	RL	var Rl	IANCE RL+FP	of FP	CORRELA RL,FP R	TION OF L+FP,FP	ROOT B1 OF RL RL+FP	KURTOS: -RL	IS (B2) RL+FP
61 62 63 64 65 66 67 68 69 70 71 72	300 300 (1) 301 300 299 (2) 300 300 300 300 300 300	900 900 900 900 900 900	200 200 200 200 200 200 200 200 200 200	196 200 188 186 200 210 236 200 205 195 195 187	509 500 877 1105 663 877 911 702 613 1204 683 849	593 585 943 1199 751 909	103 102 99 91 98 100	-0.040 -0.036 -0.057 0.004 -0.020 -0.115	0.379 0.383 0.269 0.280 0.342 0.219	0.429 0.638 0.353 0.487 -0.107 0.100 -1.164 -1.108 0.175 0.421 -1.677 -0.963	4.247 4.663 8.953 8.331 4.319 12.935	4.694 4.309 7.143 8.272 4.789 7.775
73 74 75	300 300 300	900	200 200 200	218 232 211	800 576 1073	625	87	-0.084	0.292	0.355 0.512	3.293	3.869
76	298 (3)	900	200	220	1029	1062-	84	-0.087	0.195	-0.488 -0.542	9.616	9.501

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SESS	NUMBER	nean FP	SYNCH INT	nean Rl	VARIANCE OF RL RL+FP FP	CORRELA RL,FP R	TION OF L+FP,FP	ROOT B1 O RL RL+FI	F KURTOSIS (B2) RL RL+FP
1234567890112345678901123456789012322222222222233333333333333344424444	298 (2) 297 (3) 300 300 300 300 300 300 300 300 300 3	$\begin{array}{c} 900\\ 900\\ 900\\ 900\\ 900\\ 900\\ 900\\ 900$	200 200 200 200 200 200 200 200 200 200	198 188 194 197 201 195 201 195 197 204 197 200 201 194 203 198 204 198 195 200 198 204 198 203 198 204 197 200 198 204 197 200 197 200 197 201 202 199 204 203 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 197 200 207 200 197 200 207 200 197 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 207 200 200		$\begin{array}{c} -0.039\\ 0.022\\ 0.022\\ 0.027\\ -0.077\\ -0.026\\ 0.007\\ 0.044\\ 0.929\\ -0.990\\ -0.090\\ -0.090\\ -0.090\\ -0.090\\ -0.090\\ -0.090\\ -0.0016\\ -0.022\\ -0.065\\ -0.033\\ -0.022\\ -0.065\\ -0.033\\ -0.028\\ -0.078\\ 0.003\\ -0.028\\ -0.078\\ -0.028\\ -0.078\\ -0.029\\ -0.078\\ -0.029\\ -0.078\\ -0.029\\ -0.078\\ -0.029\\ -0.078\\ -0.029\\ -0.078\\ -0.029\\ -0.001\\ -0.008\\ -0.029\\ -0.001\\ -0.004\\ -0.004\\ -0.004\\ -0.006\\ -0.043\\ -0.033\\ -0.064\\ -0.033\\ -0.064\\ -0.033\\ -0.064\\ -0.033\\ -0.064\\ -0.038\\ -0.015\\ -0.041\\ -0.038\\ -0.063\\ 0.101\\ 0.044\\ \end{array}$	0.181 0.252 0.2253 0.2253 0.2253 0.4431 0.4424 0.4424 0.4424 0.4424 0.4424 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4631 0.4532 0.4313 0.4451 0.4532 0.4313 0.4451 0.4532 0.4451 0.5515 0.4880 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605 0.4605	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	5 7.024 7.050 3 10.080 8.902 4 .372 4.062 5 9.076 8.058 3 4.579 4.256 6 8.625 6.345 7 3.441 3.196 3 3.407 4.142 5 6.641 5.698 3 3.203 3.284 7 3.233 3.284 7 3.642 3.182 7 5.403 4.166 3 3.197 3.287 3 6.139 5.560 4 .807 4.058 1 7.559 6.745 3 3.430 3.016 3 3.115 3 3.430 3.056 4 .807 4.058 1 7.559 6.745 3 3.233 3.115 3 3.430 3.056 4 .807 4.058 1 7.559 6.745 3 3.233 3.115 3 3.430 3.056 4 .807 4.058 1 2.249 8.518 2 4.257 3.943 7 3.539 2.909 1 3.046 2.892 5 4.168 4.564 7 4.871 4.708 5 5.428 4.543 3 6.134 3.514 4 3.826 3.569 2 2.905 3.394 5 3.175 4.103 3 3.705 3.706 3 3.741 4.219 3 3.580 3.256 3 3.741 4.219 3 3.799 4.001 4 1.51 5.456
447 48 50 51 52 53 55 55 56 57 58 50 50	300 300 300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	200 200 200 200 195 195 180 175 175 175 175 175	199 201 203 203 205 197 189 186 179 174 174 179 178 179 172	512 608 81 261 388 114 315 392 103 311 348 84 275 363 106 168 277 95 193 280 89 152 247 91 226 335 92 211 275 375 83 251 330 81 187 299 93 124 206 96 249 377 93 377 93	0.038 0.037 -0.070 -0.145 -0.052 0.057 -0.008 8.014 0.057 -0.103 0.057 -0.005 0.074 -0.005 0.074 0.045 0.114	0.400 0.572 0.449 0.354 0.494 0.494 0.557 0.620 0.620 0.482 0.519 0.482 0.519 0.492 0.615 0.632 0.589	-0.207 0.097 -0.290 0.377 -0.372 0.034 -0.792 -0.420 0.407 0.450 -0.210 0.366 -0.040 0.370 -0.096 0.370 -0.434 0.942 -0.096 0.141 -0.400 0.180 0.126 0.197 -0.005 0.364 0.060 0.512 -0.155 0.292	3.200 3.530 3.714 5.284 3.500 3.828 4.829 4.241 4.035 3.702 4.157 4.388 2.823 3.643 3.260 3.523 2.6214 6.066 3.012 3.019 4.270 3.872 2.865 3.022 3.094 3.567 3.691 4.208 4.018 3.820

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SESS	NUMBER TRIALS	mean FP	SYNCH INT	MEAN RL	VARIA RL RL	NCE OF +FP FP ,	CORRELA RL,FP R	TION OF L+FP,FP	RDOT RL	B1 OF RL+FP	KURTOSI -RL	IS (B2) RL+FP
61 62 63 64 65 66 67 70 71 72 73 74 75 76 77 78 80 81 82	300 300 300 300 300 300 300 300	900 900 900 900 900 900 900 900 900 900	165 160 155 150 150 150 150 150 150 150 150 15	167 162 158 157 153 153 153 153 152 152 152 152 154 154 152 147 147 145 134 132 128 130 116 117	286 170 166 194 228 170 187 203 199 201 158 259 194 300 139 160 133 251 303 380 1319 1386	341 81 248 83 257 89 264 103 276 100 247 96 247 96 247 96 247 96 247 96 247 96 247 96 2180 98 346 87 221 96 291 80 334 81 430 108 280 90 314 89	$\begin{array}{c} -0.087\\ -0.042\\ 0.008\\ -0.116\\ -0.072\\ -0.045\\ -0.050\\ -0.044\\ -0.102\\ -0.084\\ -0.102\\ -0.084\\ -0.128\\ -0.0537\\ -0.044\\ -0.128\\ -0.143\\ -0.143\\ -0.143\\ -0.144\\ -0.144\\ -0.168\\ -0.164\end{array}$	0.408 0.560 0.525 0.488 0.564 0.537 0.518 0.537 0.518 0.537 0.511 0.538 0.489 0.556 0.582 0.502 0.502 0.631 0.393 0.341 0.367 0.304	-0.120 -0.128 -0.185 -0.455 -0.324 -0.325 -0.206 -0.614 -1.028 1.580 -0.332 -0.332 -0.332 -0.339 -0.799 0.020 0.008 -0.429 -0.175 -0.323 -1.351 0.147 0.644 -0.732	$\begin{array}{c} -0.013\\ 0.270\\ 0.461\\ 0.130\\ 0.033\\ -0.046\\ -0.120\\ 1.171\\ 0.242\\ 0.110\\ 0.298\\ 0.264\\ 0.001\\ 0.695\\ 0.653\\ -0.083\\ -0.929\\ 0.421\\ 0.643\\ -0.421\\ \end{array}$	3.051 3.024 3.114 4.740 3.070 3.586 3.701 4.535 6.968 14.763 4.467 4.467 4.4675 4.421 4.753 3.172 3.204 10.683 9.592 9.983 3.756	2.986 2.819 4.066 4.279 3.285 3.721 3.457 3.111 8.107 8.635 4.091 3.872 4.020 3.888 3.563 7.523 5.921 2.811 9.869 7.428 9.129 3.705

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SESS	NUMBER MEAN TRIALS FP	synch mean Int Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	RDOT BI OF RL RL+FP	KURTOSIS (B2) RL RL+FP
SESS 1 2 3 4 5 6 7 8 9 10 11 12 3 4 15 6 7 8 9 10 11 12 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 6 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 22 3 4 15 7 8 9 20 12 20 12 20 12 10 10 10 10 10 10 10 10 10 10 10 10 10	NUMBER TRIALS MEAN FP 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 900 300 999 300 999 300 999 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 <	SYNCH INT MEAN RL 300 248 300 301 300 295 300 295 300 295 300 297 300 297 300 287 300 287 300 287 300 287 300 287 300 291 300 297 300 297 300 297 300 297 300 297 300 297 300 297 300 297 300 297 300 297 300 302 300 301 300 301 300 304 300 304 300 304 300 304 300 304	VARIANCE OF RL RL+FP FP 4990 3215 112 3458 3599 82 2113 2235 76 5008 2155 73 2655 2259 87 2029 2086 89 1550 1547 71 1303 1445 99 1905 1968 65 1270 1345 103 1542 1704 104 1436 1546 79 1291 1382 90 1032 1119 94 890 929 90 939 1054 90 846 965 90 749 831 110 786 857 112 713 793 121 713 793 144	CORRELATION OF RL,FP RL+FP,FP -0.072 0.114 0.055 0.205 0.057 0.239 -0.098 0.099 -0.001 0.183 -0.022 0.221 -0.038 0.170 -0.111 0.102 0.059 0.319 -0.003 0.179 -0.038 0.239 0.073 0.316 0.046 0.270 0.004 0.279 -0.091 0.222 0.043 0.333 0.051 0.354 -0.048 0.318 -0.069 0.325 -0.069 0.325	RDOT B1 OF RL RL+FP 1.666 1.548 1.660 1.641 0.589 0.669 0.640 0.726 0.028 0.003 0.232 0.201 0.056 0.211 -0.435 -0.387 0.191 0.240 0.196 0.201 0.530 0.515 0.075 0.109 0.229 0.244 0.383 0.480 0.431 0.279 0.323 0.295 0.136 0.276 0.522 0.700 1.562 1.320 0.277 0.347 0.282 0.295	KURTOSIS (B2) RL RL+FP 10.738 9.872 9.310 9.633 5.652 5.944 6.143 6.383 6.157 6.011 3.940 3.822 5.600 4.713 4.284 4.268 3.881 3.670 4.989 4.924 4.822 4.621 3.627 3.267 3.199 3.102 3.142 3.315 3.317 3.167 3.272 2.907 3.117 2.891 3.469 3.180 5.152 4.908 11.447 8.771 3.162 3.273 3.944 3.440
2267890123345678901234567890123345678901233456789012334567890123345678901233456789012334567890	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	300 294 300 298 300 299 300 304 300 299 300 304 300 302 300 302 300 302 300 302 300 302 300 302 300 304 300 293 300 293 300 293 300 293 300 294 300 294 300 305 300 303 300 304 300 303 300 304 300 304 300 304 300 304 305 308 305 308 305 304 305 304 305 304 305 304 305	822 878 81 822 927 105 747 836 110 814 905 94 631 685 94 774 823 85 735 864 101 813 840 66 1056 1104 79 644 696 73 516 604 115 761 840 99 744 818 83 616 626 103 554 614 80 530 631 92 549 614 81 732 791 75 896 1057 102 907 1049 99 547 679 70 777 839 74 672 717 106 646 718 75 781 878 151 815 910 97 1149 1258 96 547 507 111 478 509 911 599 711 110 575 601 57 510 563 74	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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1000 1000 1000 1000 1000 1001 1001 100	nean Fp
310 310 310 310 315 315 315 320 320 320 320 320 320 320 320 320 320	SYNCH INT
311 310 312 315 316 317 316 317 316 317 316 317 316 317 317 316 317 317 317 317 317 317 317 317 317 317	nean Rl
573 498 518 4229 980 445 4404 4224 4109 5287 879 405 1333 307 284 3485 274 273 314 2729 404 429 275 407 438 287 405 1333 2728 43485 274 22331 42557 874 4227 2543 2777 4358 2770 4358 2770 4444 2773 2777 4051 2770 2770 4371 2773 2777 4051 2770 2770 4457 4051 2770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 4770 2770 2	VARI RL R
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	ANCE OF L+FP FP
$\begin{array}{c} -0.045\\ -0.051\\ -0.070\\ 0.012\\ -0.029\\ -0.024\\ -0.156\\ -0.073\\ -0.073\\ -0.073\\ -0.073\\ -0.093\\ -0.089\\ -0.044\\ -0.094\\ -0.089\\ -0.044\\ -0.094\\ -0.0113\\ -0.0113\\ -0.014\\ -0.014\\ -0.013\\ -0.014\\ -0.014\\ -0.013\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.014\\ -0.025\\ -0.044\\ -0.025\\ -0.048\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038\\ -0.038$	CORRELA RL,FP R
$\begin{array}{c} 0.337\\ 0.344\\ 0.314\\ 0.353\\ 0.430\\ 0.387\\ 0.435\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.320\\ 0.423\\ 0.420\\ 0.420\\ 0.425\\ 0.440\\ 0.325\\ 0.440\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.443\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.325\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.445\\ 0.455\\ 0.455\\ 0.455\\ 0.455\\ 0.455\\ 0.455\\ 0.455\\ 0.$	TION OF L+FP,FP
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	ROOT BI DF RL RL+FP
3.144 2.746 2.808 3.148 2.787 3.305 3.474 3.456 4.671 2.973 3.355 3.355 2.973 3.355 2.973 3.440 4.206 2.973 3.440 4.206 2.973 3.440 4.206 2.973 3.273 3.440 4.206 3.440 4.206 3.440 4.206 3.440 4.206 3.440 4.206 3.440 2.979 3.130 3.627 3.306 3.306 3.307 3.519 3.306 3.2973 3.2973 3.2973 3.2975 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2975 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2973 3.2974 2.726 3.2274 3.2273 3.2274 2.726 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2274 3.2273 3.2374 3.2726 3.2274 3.2273 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2726 3.2374 3.2936 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937 3.2937	KURTOSI RL
4.002 3.133 3.157 3.050 3.954 3.050 3.953 3.953 3.953 3.953 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.927 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273 3.9273	S (B2) RL+FP

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DK VISUAL

SESS	NUMBER	nean	SYNCH	nean	VARIANCE OF	CORRELA	TION OF	ROOT B1 OF	KURTOSIS (82)
	TRIALS	FP	INT	Rl	RL RL+FP F	P RL,FP F	L+FP,FP	RL RL+FP	'RL RL	+FP
$\begin{array}{c} 121\\ 122\\ 123\\ 124\\ 125\\ 126\\ 127\\ 128\\ 129\\ 130\\ 131\\ 132\\ 133\\ 134\\ 135\\ 136\\ 137\\ 138\\ 139\\ 140\\ 141\\ 142\\ 143\\ 144\\ 145\\ 153\\ 155\\ 157\\ 158\\ 157\\ 158\\ 157\\ 158\\ 166\\ 163\\ 164\\ 165\\ 166\\ 167\\ 172\\ 173\\ 177\\ 178\\ 180\\ 180\\ 180\\ 180\\ 180\\ 180\\ 180\\ 18$	300 300 300 300 300 300 300 300	1001 1001 1000 1000 1000 1000 1000 100	$\begin{array}{c} 400\\ 400\\ 400\\ 400\\ 410\\ 420\\ 425\\ 425\\ 425\\ 425\\ 425\\ 425\\ 425\\ 425$	$\begin{array}{c} 399\\ 397\\ 399\\ 407\\ 422\\ 877\\ 420\\ 228\\ 437\\ 450\\ 450\\ 450\\ 450\\ 450\\ 450\\ 450\\ 450$	381 478 1478 427 506 376 500 352 426 473 532 448 477 514 639 536 617 577 680 442 552 511 583 446 488 690 718 442 552 511 583 442 552 517 576 478 558 579 649 629 745 748 909 528 581 517 576 629 745 748 909 528 581 517 576 629 745 549 638 511 598 638 714 767 804 520 527 <th>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</th> <th>$\begin{array}{c} 0.451\\ 0.396\\ 0.457\\ 0.416\\ 0.339\\ 0.294\\ 0.342\\ 0.349\\ 0.442\\ 0.356\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.321\\ 0.249\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.3229\\ 0.395\\ 0.324\\ 0.329\\ 0.327\\ 0.442\\ 0.3219\\ 0.325\\ 0.324\\ 0.3249\\ 0.327\\ 0.304\\ 0.328\\ 0.365\\ 0.325\\ 0.304\\ 0.328\\ 0.365\\ 0.325\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.353\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 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0.457\\ 0.416\\ 0.339\\ 0.294\\ 0.342\\ 0.349\\ 0.442\\ 0.356\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.325\\ 0.321\\ 0.249\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.395\\ 0.3219\\ 0.3229\\ 0.395\\ 0.324\\ 0.329\\ 0.327\\ 0.442\\ 0.3219\\ 0.325\\ 0.324\\ 0.3249\\ 0.327\\ 0.304\\ 0.328\\ 0.365\\ 0.325\\ 0.304\\ 0.328\\ 0.365\\ 0.325\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.365\\ 0.353\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.353\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 0.355\\ 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DK VISUAL

SESS	NUMBER MEAN TRIALS FP	synch mean Int RL	VARIANCE OF . RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT 81 OF RL RL+FP	KURTOSIS (B2) -RL RL+FP
181 182 183 184 185 185 185 187 188 187 191 192 193 194 195 195 195 197 198 199 200 201 202	300 979 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1001 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 999 300 999 300 999 300 999	490 487 490 491 485 481 485 484 485 483 485 482 485 487 485 487 485 482 485 487 480 482 480 483 480 481 480 482 475 475 475 475 475 479 475 479 475 479 475 470 470 470 470 470 470 471 470 471	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2.995 2.888 2.768 2.753 2.681 2.829 3.175 3.251 2.785 3.233 3.224 15.528 3.369 3.123 3.127 4.363 2.800 3.283 2.976 4.013 2.871 3.062 3.029 2.916 3.015 2.919 3.435 2.819 3.260 3.100 2.962 3.036 3.009 3.151 2.820 3.848 3.205 3.797 2.785 4.897 3.639 3.009 3.022 3.168
203 204 205 206 207 210 211 212 213 214 215 216 217 218 219 221 212 213 214 215 216 227 228 229 221 2223 2224 225 2230 2331 2334 2354 2357 238 239 2312 2334 2356 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 238 237 237 238 237 237 237 237 237 237 237 237 237 237	0 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 300 1000 3	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{c} 0\\ 315 & 393 & 89\\ 273 & 350 & 93\\ 316 & 394 & 98\\ 282 & 365 & 89\\ 285 & 340 & 88\\ 249 & 353 & 89\\ 310 & 398 & 97\\ 283 & 342 & 88\\ 349 & 420 & 89\\ 293 & 374 & 89\\ 293 & 374 & 89\\ 293 & 374 & 89\\ 293 & 374 & 89\\ 274 & 338 & 88\\ 240 & 298 & 83\\ 315 & 415 & 95\\ 224 & 299 & 114\\ 430 & 550 & 87\\ 281 & 366 & 96\\ 365 & 395 & 90\\ 301 & 374 & 81\\ 320 & 426 & 89\\ 382 & 440 & 98\\ 260 & 373 & 89\\ 279 & 350 & 88\\ 342 & 422 & 83\\ 342 & 422 & 83\\ 342 & 422 & 83\\ 263 & 343 & 89\\ 279 & 350 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 308 & 367 & 88\\ 289 & 390 & 95\\ 241 & 289 & 97\\ 290 & 381 & 114\\ 322 & 378 & 102\\ 257 & 352 & 92\\ 266 & 369 & 93\\ 275 & 327 & 96\\ 256 & 368 & 94\\ 191 & 278 & 89\\ \end{array}$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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DK VISUAL

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SESS	NUMBER TRIALS	nean Fp	SYNCH INT	mean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS (B2) RL RL+FP
241 242 243	300 300 300	1000 999	435 435 435	437 433 436	269 338 92 316 406 92 263	-0.074 0.456 -0.006 0.472	0.038 0.212 -0.082 0.212	2.819 3.217 2.796 3.101
22222222222222222222222222222222222222	1) 1) 1) 1) 1) 1) 1) 1) 1) 1)	1000 1000 1000 1000 1000 1000 1000 100	4430004425555000000055555500000000555555555	7331 428 4322 424 424 425 537 428 425 537 428 417 418 410 410 410 400 400 400 400 229 60 399 399 395 395 395 388 388 388 388 388 388 388 388 388 38	253 318 92 303 374 80 288 346 79 268 366 102 298 366 89 293 369 84 220 305 93 282 383 86 250 345 94 270 384 89 246 307 84 250 345 94 270 384 89 246 307 84 250 345 94 270 384 89 246 307 84 250 345 94 271 362 94 2721 294 97 217 312 92 230 320 119 193 277 79 219 358 135 235 329 84 287 344 90 287 344 79 <	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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DK VISUAL

SESS	NUMBER TRIALS	mean Fr	SYNCH INT	MEAN RL	VARI. RL R	ANCE OF L+FP FP	CORRELA RL,FP R	TION OF L+FP,FP	ROOT B1 OF RL RL+FP	KURTOSI -RL	S (B2) RL+FP
301 302 303 304 305 306 307 308 307 310 312 313 314 315 317 319 321 322 321 322 3224 325 326 327	300 300 300 300 300 300 300 300 300 300	1000 999 1000 1000 1000 1000 1000 1000	375 375 370 370 370 365 365 365 360 360 360 360 360 355 355 355 355 355 355 355 355 355 35	372 374 373 370 370 370 363 363 363 364 365 366 362 364 362 364 362 355 355 355 355 355 355 355 355 352 352	451 168 164 193 208 168 205 180 232 161 209 216 209 216 209 216 209 216 209 216 209 216 201 195 188 211 206 203 193 226 164 229	558 88 249 79 223 90 243 87 306 86 266 111 331 113 269 87 323 86 249 84 298 108 286 95 277 69 317 103 271 79 268 108 279 90 268 108 277 80	$\begin{array}{c} 0.048\\ 0.010\\ -0.129\\ -0.146\\ 0.041\\ -0.050\\ 0.041\\ 0.007\\ 0.015\\ -0.138\\ -0.015\\ -0.138\\ -0.015\\ -0.004\\ -0.005\\ -0.041\\ 0.027\\ -0.041\\ -0.045\\ -0.045\\ -0.044\\ -0.045\\ -0.035\\ -0.079\\ -0.044\\ -0.045\\ -0.079\\ -0.044\\ -0.045\\ -0.079\\ -0.041\\ -0.045\\ -0.079\\ -0.079\\ -0.041\\ -0.079\\ -0.076\\ -0.111\\ \end{array}$	0.440 0.572 0.523 0.469 0.545 0.545 0.575 0.575 0.575 0.575 0.529 0.480 0.515 0.495 0.505 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.545 0.555 0.545 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.555 0.550 0.555 0.555 0.555 0.550 0.555 0.555 0.555 0.550 0.555 0.550 0.555 0.555 0.555 0.550 0.555 0.555 0.555 0.550 0.555 0.550 0.555 0.550 0.550 0.555 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.550 0.5500 0.5500 0.5500 0.5500000000	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	8.045 3.188 2.456 3.220 4.145 3.127 3.626 3.168 3.270 2.956 3.854 2.916 2.956 3.854 2.816 2.956 3.479 4.173 3.309 2.803 3.502 3.442 2.979 2.803 3.502 3.442 2.979 2.845 3.255 3.133	5.455 3.318 3.318 4.638 4.638 4.638 4.638 4.638 4.638 4.638 4.638 4.638 4.638 3.012 3.012 3.642 3.257 3.228 3.859 2.830 3.925 3.925 3.925 3.925 3.759 3.759 3.757 3.135
328 329 331 332 3334 335 334 335 335 335 335 335 335 341 342 343 345 345 345 345 355 355 355 355 355	300 300 300 300 300 300 300 300 300 300	1001 1000 1000 1000 1000 1000 1000 100	350 345 345 345 345 340 340 340 340 340 335 335 335 335 330 330 330 330 330 325 325 325 320 320 320 320 320 320 320 320 320 320	346 348 348 346 348 346 339 339 349 339 349 339 349 339 349 339 33	153 180 215 166 152 176 175 211 252 154 235 210 191 210 142 179 192 220 173 181 187 185 184 173 185 184 173 185 183	243 111 328 104 203 64 270 113 271 113 272 84 277 111 273 103 302 79 212 64 334 113 249 80 276 103 311 107 203 81 209 71 236 74 279 86 246 90 266 92 261 89 273 113 239 96 247 84 341 103 234 79 242 81 264 103 327 97 248 103 327 97 281 84 253 73	$\begin{array}{c} -0.170\\ 0.027\\ -0.128\\ 0.013\\ -0.003\\ -0.138\\ -0.138\\ -0.104\\ -0.028\\ -0.159\\ -0.159\\ -0.066\\ -0.022\\ -0.091\\ -0.185\\ -0.129\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.083\\ -0.083\\ -0.083\\ -0.083\\ -0.0156\\ -0.043\\ -0.015\\ -0.088\\ -0.017\\ -0.013\end{array}$	0.530 0.587 0.445 0.660 0.597 0.478 0.407 0.492 0.478 0.525 0.545 0.555 0.554 0.554 0.554 0.534 0.534 0.573 0.573 0.573 0.573 0.573 0.573 0.573 0.573 0.573 0.573 0.573 0.575 0.534 0.575 0.534 0.575 0.534 0.575 0.534 0.575 0.575 0.534 0.575 0.553 0.546 0.575 0.553 0.5534 0.5575 0.534 0.575 0.575 0.575 0.534 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.575 0.558 0.558 0.5528 0.5528	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2.789 3.474 2.925 4.260 3.328 2.853 3.305 2.938 2.761 2.980 2.833 3.334 5.832 2.806 3.402 3.322 2.538 3.402 3.322 2.538 3.402 3.322 3.418 3.402 3.322 3.418 3.160 3.047 2.567 3.864 3.135 3.028 3.264 3.264 3.2714 2.688	3.421 3.131 3.388 3.3757 3.388 3.3757 3.388 3.3757 3.388 3.3757 3.388 3.3757 3.399 3.34.544 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.327 3.3277 3.327 3.3277 3.3277 3.3277 3.3277 3.327

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DK VISUAL

SESS	NUMBER TRIALS	mean FP	SYNCH INT	Mean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS (B2) RL RL+FP
361 362 363 364 365 366 367 373 374 375 377 377 377 377 377 377 377 377 378 377 378 377 378 381 382 383 381 382 383 385 385 386 387 392 392 392 392 392 392	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 1000 1000 1000 1000 100	33333333333333333333333333333333333333	319 318 311 311 302 303 305 305 305 305 305 305 305 305 305	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.024 0.537 -0.060 0.540 -0.093 0.481 -0.097 0.463 -0.158 0.464 -0.088 0.551 -0.158 0.464 -0.088 0.551 -0.143 0.492 -0.080 0.510 -0.228 0.457 -0.186 0.552 -0.184 0.552 -0.184 0.552 -0.184 0.575 -0.118 0.524 -0.027 0.512 -0.057 0.530 -0.069 0.536 -0.140 0.546 -0.204 0.509 -0.082 0.514 -0.092 0.526 -0.047 0.514 -0.092 0.526 -0.047 0.511 -0.123 0.531 -0.224 0.474 -0.082 0.552 -0.137 0.505 -0.137 0.505 -0.126 0.533 -0.208 0.551	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
375 397 397 398 398 400 401 402 403 404 405 407 408 407 408 407 408 407 408 407 411 412 414 415 416 419 420	0 300 300 300 300 300 300 300 300 300 3	1000 1000 1000 1000 1000 1001 1001 100	275505020000000555555555555555555555555	0 279 280 284 290 296 304 301 304 300 303 306 300 300 300 306 306 306 306	U 138 208 83 145 211 88 191 246 84 174 273 103 150 200 100 168 213 97 133 211 109 132 197 106 137 219 94 144 232 86 134 177 89 132 180 83 165 228 88 148 228 85 158 280 103 177 241 80 164 220 73 178 227 71 169 251 49 177 226 80 205 230 84 197 276 85 160 186 86 158 200 71	$\begin{array}{ccccccc} -0.058 & 0.583 \\ -0.172 & 0.492 \\ -0.111 & 0.485 \\ -0.013 & 0.603 \\ -0.206 & 0.531 \\ -0.211 & 0.495 \\ -0.125 & 0.617 \\ -0.175 & 0.591 \\ -0.050 & 0.615 \\ 0.011 & 0.618 \\ -0.213 & 0.524 \\ -0.167 & 0.535 \\ -0.103 & 0.532 \\ -0.025 & 0.592 \\ 0.075 & 0.662 \\ -0.066 & 0.521 \\ -0.074 & 0.511 \\ -0.098 & 0.472 \\ 0.061 & 0.575 \\ 0.022 & 0.582 \\ -0.131 & 0.480 \\ -0.224 & 0.391 \\ -0.25 & 0.535 \\ -0.254 & 0.443 \\ -0.134 & 0.477 \end{array}$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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DK VISUAL

SE5S	NUMBER	mean	SYNCH	nean	varjanci	OF	CORRELA	TION OF	root	B1 OF	KURTOSI	S (B2)
	TRIALS	FP	INT	Rl	RL' rl+fi	FP	RL,FP RI	L+FP,FP	RL	RL+FP	RL	RL+FP
421 422 423 424 425 427 425 427 428 427 428 427 430 431 432 433 435 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 437 438 438 437 438 437 438 438 437 438 437 438 437 438 437 438 438 437 438 438 437 438 438 437 438 438 438 438 438 438 438 438 438 438	300 300 300 300 300 300 300 300	1000 1000 1000 1001 1001 1000 1001 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 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\\ 130 \\ 177 \\ 244 \\ 132 \\ 227 \\ 130 \\ 177 \\ 244 \\ 132 \\ 227 \\ 130 \\ 177 \\ 244 \\ 132 \\ 227 \\ 130 \\ 177 \\ 244 \\ 132 \\ 227 \\ 130 \\ 177 \\ 244 \\ 132 \\ 227 \\ 130 \\ 177 \\ 235 \\ 235 \\ 255 \\ 297 \\ 160 \\ 244 \\ 146 \\ 205 \\ 255 \\ 297 \\ 160 \\ 244 \\ 146 \\ 205 \\ 297 \\ 154 \\ 244 \\ 146 \\ 205 \\ 297 \\ 154 \\ 244 \\ 146 \\ 205 \\ 297 \\ 154 \\ 244 \\ 146 \\ 205 \\ 297 \\ 154 \\ 244 \\ 146 \\ 205 \\ 297 \\ 163 \\ 211 \\ 213 \\ 213 \\ 213 \\ 214 \\ 213 \\ 214 \\ 213 \\ 214 \\ 213 \\ 214 \\ 214 \\ 213 \\ 214 \\ 214 \\ 213 \\ 214 \\ 214 \\ 213 \\ 214 \\ 215 \\ 215 \\ 250 \\ 33 \\ 161 \\ 257 \\ 174 \\ 215 \\ 250 \\ 33 \\ 161 \\ 257 \\ 174 \\ 215 \\ 250 \\ 33 \\ 161 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 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257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 257 \\ 25$	7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 8 9 9 7 1 1 7 7 7 7 8 9 9 7 1 1 1 9 9 7 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	-0.040 -0.108 -0.022 -0.048 -0.022 -0.048 -0.113 -0.053 -0.196 -0.160 -0.160 -0.183 -0.017 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 -0.024 -0.028 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480	300	999	360	360	154 24	17 108	-0.056	0.617	0.166	1.012	3.104	7.365

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SESS	NUMBER	Mean	SYNCH	nean	VARI	ANCE OF	CORRELA	TION OF	ROOT B1 OF	KURTOSI	\$ (82)
	TRIALS	FP	INT	Rl	RL R	L+FP FP	RL,FP R	L+FP,FP	RL RL+FF	RL	RL+FP
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SESS	NUMBER TRIALS	Mean FP	Synch Int	nean Rl	VARIANCE OF RL RL+FP F	F CORRELA	ATION OF RL+FP,FP	ROOT 81 OF RL RL+FP	KURTOSIS (B 'RL RL+	12) FP
121 122 123 124 125 126 127 128 129 130 131 132 133 134 135 136 139 141	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 1001 1001 1000 1000 100	400 400 400 400 400 400 400 400 400 400	397 392 381 392 384 387 391 402 397 390 393 393 393 399 410 393 395 408 407 407	483 557 6 393 492 5 771 811 5 359 418 7 543 648 5 489 569 7 429 485 5 374 437 8 384 477 10 466 608 5 305 390 6 450 521 5 352 438 12 352 438 12 329 488 12 342 455 10 418 525 8 331 484 11 358 539 10 315 457 11	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$	0.365 0.449 0.241 0.377 0.403 0.375 0.403 0.375 0.403 0.375 0.453 0.4491 0.382 0.442 0.442 0.442 0.442 0.442 0.442 0.458 0.576 0.500 0.458 0.568 0.561 0.561	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2.590 2.9 2.973 3.6 5.342 4.5 4.126 4.2 7.582 5.4 3.479 3.3 4.074 4.8 3.226 3.4 3.866 3.7 3.935 3.4 2.963 3.4 3.134 2.7 3.149 3.5 3.343 2.8 3.331 4.4 2.995 3.7 3.388 3.7 2.807 3.1 2.841 3.6 2.881 3.2	751 124 1932 1813 1813 1813 1813 1813 1813 1813 18
141 142 143 1445 1447 148 149 150 152 153 155 157 158 161 163 164 1689 171 1773 1756 1789 180	300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 3	1000 1000 1000 1000 1000 1000 1000 100	410 412 425 425 425 425 425 425 425 425 425 42	410 419 421 421 429 434 452 444 452 444 452 444 452 450 450 450 450 450 450 450 450 450 450	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.372 0.404 0.429 0.407 0.444 0.354 0.447 0.442 0.376 0.376 0.442 0.442 0.446 0.448 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.446 0.386 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.376 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375 0.375	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	5.623 4.5 3.367 3.2 4.519 4.1 3.535 3.4 3.512 3.1 4.293 3.6 5.604 4.6 3.013 2.9 3.317 3.0 3.145 3.0 4.719 4.4 3.675 3.2 2.935 3.1 2.773 2.9 2.912 3.0 3.237 3.2 4.364 4.1 3.982 3.6 3.321 2.9 2.912 3.0 3.321 2.9 2.906 3.0 3.462 3.6 3.271 3.1 5.843 5.6 2.882 3.6 3.271 3.1 5.843 5.6 2.882 3.6 3.219 3.2 4.018 4.9 3.001 3.1 2.793 2.7 3.170 3.8 2.924 3.0 3.219 3.2 4.2918 3.0 3.231 3.2 4.2918 3.0 3.2219 3.2 4.2918 3.0 3.2219 3.2 3.231 3.7 3.123 3.7 3.231 3.7	7581 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970 1970

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SESS	NUMBER TRIALS	nean Fp	STNCH INT	Mean Rl	Vari. RL R	ANCE OF L+FP FP	CORRELA RL,FP R	TI ON OF L+FP,FP	ROOT B1 DF RL RL+FP	KURTOSI 'RL	S (82) RL+FP
181 182 183 184 185	300 300 300 300 300 300	999 1000 1000 999	490 490 490 485 485	490 491 482 481 487	262 476 364 288 399	354 80 553 104 400 91 357 79	0.042 -0.060 -0.152 -0.034	0.511 0.377 0.333 0.440	-0.059 0.345 -0.105 -0.172 -0.018 0.060 0.448 0.553	2.707 3.251 2.798 3.355	3.363 3.063 2.797 3.551
186 187 188 190 191 192 193 194 195 196 197 198 199 200	300 300 300 300 300 300 300 300 300 300	1001 1000 999 1000 999 1001 1000 1000 1	485 485 485 485 485 485 485 485 485 485	472 485 476 486 485 488 488 488 488 488 488 488 481 480 480 480 480	520 430 584 465 419 277 309 328 257 461 441 297 303 353 324	542 98 701 95 578 80 489 91 353 67 409 100 403 87 354 95 532 79 516 64 370 89 474 89 445 85	0.034 0.046 0.086 -0.054 0.035 -0.001 -0.035 0.007 -0.020 0.026 -0.050 0.117 0.088 0.109	0.455 0.410 0.448 0.382 0.466 0.494 0.433 0.524 0.367 0.382 0.445 0.566 0.510 0.530	0.409 0.474 0.909 0.765 -0.021 0.108 -0.114 0.121 -0.028 0.357 0.058 0.328 0.068 0.088 0.223 0.354 -0.186 -0.009 -1.186 -0.987 0.177 0.521 0.144 0.349 0.003 0.294 0.055 0.262	4.288 7.702 3.081 3.160 3.212 3.077 2.827 2.843 3.206 9.945 3.358 2.816 2.592 2.992	4.303 5.925 3.203 3.640 2.842 2.874 3.115 3.097 8.476 3.461 2.968 3.043
201 202 203	300 300 300	1000 1000	485 485 480	482 482 476	335 279 224	435 80 343 102	0.062 0.055	0.482 0.590	-0.390 -0.025	3.152 3.015	3.176 3.771
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IS VISUAL

SESS	NUMBER	mean	synch mean	VARIANCE OF -	CORRELATION OF	ROOT B1 OF	KURTOSIS (82)
	TRIALS	FP	Int Rl	RL RL+FP FP	RL,FP RL+FP,FP	RL RL+FP	RL RL+FP
4223 4223 4223 4223 4223 4224 4225 4228 4220 4228 4220 4228 4220 4220 4220	300 300 300 300 300 300 300 300 300 300	999 1001 999 1000 1000 1000 1000 1000 1	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	153238801472371061352309410017091124203791332321191051998712219675118209841091707694181841072051179017087861578099174761462519614725291117266128101205938916397931899514123984103205911072048911226313810318297138235891593008917628187921477312324710689168879214773123247106891688792147731232471068916413910319172151247761382339012824613910319772151247781011018110315776126 <td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td> <td>$\begin{array}{c} -0.204 & 0.346 \\ -0.175 & 0.225 \\ -0.135 & 0.635 \\ -0.402 & 0.807 \\ -0.223 & 0.514 \\ -0.173 & 0.744 \\ -0.029 & 0.671 \\ 0.042 & 0.817 \\ 0.041 & 0.373 \\ 0.192 & 0.421 \\ -0.085 & 0.812 \\ 0.192 & 0.421 \\ -0.085 & 0.812 \\ 0.115 & 1.292 \\ -0.037 & 0.339 \\ 0.169 & 0.774 \\ -0.322 & 0.593 \\ -0.262 & 0.527 \\ 0.157 & 0.608 \\ 0.102 & 0.862 \\ 0.030 & 0.691 \\ 0.120 & 0.710 \\ -0.010 & 0.587 \\ 0.347 & 0.654 \\ 0.235 & 0.897 \\ 0.648 & 1.031 \\ -0.014 & 2.597 \\ 0.017 & 1.275 \\ 0.028 & 0.488 \\ 0.253 & 0.881 \\ 0.283 & 0.487 \\ -0.021 & 0.527 \\ 0.062 & 0.559 \\ 0.315 & 0.940 \\ 0.017 & 0.861 \\ 0.012 & 0.343 \\ -0.021 & 0.527 \\ 0.062 & 0.559 \\ 0.315 & 0.940 \\ 0.017 & 0.861 \\ 0.012 & 0.343 \\ -0.343 & 0.704 \\ 0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ 0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ 0.059 & 0.776 \\ 0.003 & 0.776 \\ 0.023 & 0.572 \\ -0.118 & 0.678 \\ -0.257 & 0.579 \\ 0.133 & 0.563 \\ 0.053 & 0.974 \\ 0.003 & 0.776 \\ 0.283 & 0.562 \\ -0.202 - 0.166 & 0.182 \\ -0.202 - 0.166 & 0.182 \\ -0.203 & 0.776 \\ 0.203 & 0.776 \\ 0.203 & 0.776 \\ 0.023 & 0.776 \\ 0.021 & 0.431 \\ 0.271 & 0.859 \\ 0.037 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.859 \\ 0.035 & 0.037 \\ 0.048 & 0.846 \\ -0.0$</td> <td>$\begin{array}{cccccccccccccccccccccccccccccccccccc$</td>	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{c} -0.204 & 0.346 \\ -0.175 & 0.225 \\ -0.135 & 0.635 \\ -0.402 & 0.807 \\ -0.223 & 0.514 \\ -0.173 & 0.744 \\ -0.029 & 0.671 \\ 0.042 & 0.817 \\ 0.041 & 0.373 \\ 0.192 & 0.421 \\ -0.085 & 0.812 \\ 0.192 & 0.421 \\ -0.085 & 0.812 \\ 0.115 & 1.292 \\ -0.037 & 0.339 \\ 0.169 & 0.774 \\ -0.322 & 0.593 \\ -0.262 & 0.527 \\ 0.157 & 0.608 \\ 0.102 & 0.862 \\ 0.030 & 0.691 \\ 0.120 & 0.710 \\ -0.010 & 0.587 \\ 0.347 & 0.654 \\ 0.235 & 0.897 \\ 0.648 & 1.031 \\ -0.014 & 2.597 \\ 0.017 & 1.275 \\ 0.028 & 0.488 \\ 0.253 & 0.881 \\ 0.283 & 0.487 \\ -0.021 & 0.527 \\ 0.062 & 0.559 \\ 0.315 & 0.940 \\ 0.017 & 0.861 \\ 0.012 & 0.343 \\ -0.021 & 0.527 \\ 0.062 & 0.559 \\ 0.315 & 0.940 \\ 0.017 & 0.861 \\ 0.012 & 0.343 \\ -0.343 & 0.704 \\ 0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ 0.059 & 0.776 \\ 0.002 & 0.614 \\ -0.059 & 0.776 \\ 0.002 & 0.614 \\ 0.059 & 0.776 \\ 0.003 & 0.776 \\ 0.023 & 0.572 \\ -0.118 & 0.678 \\ -0.257 & 0.579 \\ 0.133 & 0.563 \\ 0.053 & 0.974 \\ 0.003 & 0.776 \\ 0.283 & 0.562 \\ -0.202 - 0.166 & 0.182 \\ -0.202 - 0.166 & 0.182 \\ -0.203 & 0.776 \\ 0.203 & 0.776 \\ 0.203 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.023 & 0.776 \\ 0.021 & 0.431 \\ 0.271 & 0.859 \\ 0.037 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.846 \\ -0.071 & 0.431 \\ 0.271 & 0.859 \\ 0.035 & 0.037 \\ 0.068 & 0.859 \\ 0.035 & 0.037 \\ 0.048 & 0.846 \\ -0.0$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

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SESS	NUMBER TRIALS	mean FP	SYNCH INT	Mean Rl	VA RL	RIANCE RL+FP	OF FP	CORREL RL,FP	ATION OF RL+FP,FP	ROOT RL	B1 OF RL+FP	KURTOSI 'RL	S AB2) RL+FP
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3500	$\begin{array}{c} 347\\ 351\\ 345\\ 345\\ 352\\ 362\\ 372\\ 362\\ 372\\ 375\\ 375\\ 375\\ 375\\ 375\\ 375\\ 375\\ 375$	1473 105 1204 1294 1294 1294 1294 1294 1294 1294 129	$\begin{array}{c} 217\\ 250\\ 196\\ 194\\ 186\\ 225\\ 204\\ 213\\ 237\\ 282\\ 204\\ 213\\ 237\\ 282\\ 205\\ 205\\ 205\\ 205\\ 205\\ 205\\ 205\\ 20$	781 871 667678989500326768998107267689999999999999999999999999999999999	$\begin{array}{c} -0.039\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0788\\ -0.0128\\ -0.0128\\ -0.0128\\ -0.0128\\ -0.0128\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0156\\ -0.0258\\ -0.0258\\ -0.026\\ -0.026\\ -0.026\\ -0.026\\ -0.026\\ -0.026\\ -0.026\\ -0.026\\ 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SESS	NUMBER	nean	Synch	Nean	VARIANCE OF	CORRELATION OF	ROOT BI OF	KURTOSIS (B2)
	TRIALS	FP	Int	, Rl	RL RL+FP FP	RL,FP RL+FP,FP	RL RL+FP	RL RL+FP
123456789011234567890122222222222222333333333444444444444455555555	300 300 300 300 300 300 300 300 300 300	1001 1000 1000 1000 1000 1000 1000 100	300 300 300 300 300 300 300 300 300 300	303 303 305 305 305 305 305 305 305 307 305 307 307 303 307 307 307 307 307 307 307	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	4.778 4.813 3.295 3.196 3.537 3.154 4.730 3.643 3.475 4.340 3.644 3.428 2.841 2.553 3.223 3.765 2.912 3.212 3.813 3.665 3.193 3.169 3.911 3.038 3.055 3.068 3.134 3.435 3.196 3.194 2.813 3.699 3.353 6.176 3.384 3.134 3.374 3.179 3.243 3.439 3.143 3.289 2.669 2.833 3.143 3.289 2.669 2.631 3.705 3.247 3.183 3.261 4.184 4.503 2.890 3.031 3.508 4.496 2.696 2.643 3.144 3.438 3.195 3.051 3.508 4.496 2.696 3.540 3.144 3.438 3.195 3.362 3.143 3.289 3.363 3.572 2.909 3.560 3.405 3.224 3.607 3.284 3.292 3.518 3.405 3.292 3.174 2.977 3.274 3.297 3.274 3.297 3.274 3.297 3.274 3.297 3.765 3.974 3.176 3.9

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SESS	NUMBER TRIALS	nean FP	SYNCH MEAN INT RL	var RL	RIANCE OF RL+FP FP	CORREL RL,FP	ATION OF RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSI	IS (B2) RL+FP
6123345667890123777777778901234566788901233456678890112344567889011111111111111111111111111111111111	300 300 300 300 300 300 300 300 300 300	1000 1001 1000 1000 1001 1001 1001 100	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	502 576 549 708 578 578 578 578 578 578 578 57	544 81 674 120 612 89 802 71 646 90 831 86 703 113 618 85 479 93 656 85 476 88 440 113 506 87 482 98 592 84 653 117 581 96 597 107 750 100 539 73 497 70 474 78 499 99 411 82 1276 119 370 67 599 103 448 93 1165 755 976 474 487 88 540 89 487 88 560 89 914 413 935 562 914 413 938 84 441 85 599 94 418 84 599 94 418 84 592 87 300 82 344 81 592 87	-0.098 -0.043 -0.059 0.053 -0.048 -0.081 0.025 0.051 -0.005 0.020 -0.113 -0.142 -0.122 0.011 -0.026 0.042 -0.032 -0.042 -0.031 0.016 -0.042 -0.031 0.016 -0.028 -0.029 -0.038 -0.017 0.031 0.016 -0.029 -0.029 -0.085 -0.015 0.015 0.015 0.015 -0.045 -0.028 -0.015 0.015 0.015 -0.045 -0.028 -0.015 0.015 -0.045 -0.028 -0.015 0.015 -0.028 -0.015 -0.028 -0.015 -0.028 -0.015 -0.028 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141 142 143 144 145 146 147 148 149 151 152 153 155 157 158 160 161 163 165 168 170 172 173 175	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 1001 1001 1001 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000 1000	480 485 500 500 500 500 500 500 500 500 500 5	487 487 493 497 506 500 500 500 500 500 500 500 500 500	11 33 1373 54 435 54 4360 52 329 52 329 52 520 52 520 52 520 52 520 52 548 52 550 548 520 52 548 52 550 548 520 52 548 52 550 548 520 52 548 52 520 52 548 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52 520 52	4 76 77 103 89 59 10 11 11 10 10 10 10 10 10 10 10 10 10	-0.030 0.032 0.020 0.001 0.028 -0.055 0.106 -0.053 -0.105 0.057 0.007 -0.054 -0.054 0.002 -0.164 0.002 -0.161 0.004 0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.015 -0.024 -0.015 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- LL VISUAL

SESS	NUMBER	mean	SYNCH	Mean	vari	IANCE OF	CORRELA	TION OF	RDOT	B1 OF	KURTOSI	S (B2)
	TRIALS	FP	INT	Rl	RL F	RL+FP FP	RL,FP R	L+FP,FP	RL	RL+FP	•RL	RL+FP
241 242 243 244 245 246 247 248 247 248 247 250 251 252 253 254 255 255 255 255 255 255 255 257 258 257 258 257 258 257 258 257 264	300 300 300 300 300 300 300 300 300 300	1000 1001 1000 1000 1000 999 1001 1001	350 350 349 335 325 325 325 310 300 300 300 300 300 300 300 300 300	360 352 344 337 330 322 315 314 302 305 310 304 305 300 303 300 305 300 305 300	213 189 254 293 229 316 255 205 273 281 309 378 303 337 440 272 332 376 262 251	255 83 304 105 341 98 397 93 315 80 363 78 390 81 430 120 311 121 339 92 394 76 426 105 465 76 406 118 500 128 570 105 397 85 400 75 435 90 319 92 343 92	-0.155 0.038 -0.031 0.021 0.021 -0.014 -0.021 0.159 -0.049 -0.083 0.127 0.031 0.034 -0.039 0.057 0.057 0.057 0.057 0.057 0.057 0.057 0.057 0.024 -0.024 -0.024 -0.024 -0.03	0.428 0.616 0.508 0.512 0.522 0.452 0.436 0.522 0.436 0.584 0.584 0.584 0.584 0.584 0.584 0.525 0.585 0.505 0.435 0.480 0.413 0.376 0.413 0.376	$\begin{array}{c} 0.069\\ -0.077\\ -0.018\\ 0.531\\ -0.058\\ -0.204\\ 0.062\\ -0.019\\ -0.154\\ 0.137\\ 0.413\\ 0.087\\ 0.153\\ 0.494\\ 0.564\\ 0.564\\ 0.195\\ -0.186\\ 0.450\\ -0.003\\ 0.166\end{array}$	0.299 0.608 0.294 0.551 0.163 0.267 0.467 0.467 0.084 0.381 0.381 0.311 0.311 0.127 0.649 0.641 0.108 0.583 0.213 0.122	3.380 3.324 2.913 4.626 3.202 3.575 3.421 3.148 3.059 3.042 3.219 3.182 3.365 3.522 4.984 5.000 3.652 2.548 3.547 2.575 3.510	3.039 5.157 3.769 3.731 3.141 3.075 3.669 3.669 3.669 3.669 3.652 3.857 2.757 2.824 3.652 3.485 4.226 4.372 4.372 3.749 2.963 3.664 3.209 3.142

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JS VISUAL

SESS	NUMBER	mean FP	SYNCH INT	nean Rl	VARIANCE OF RL RL+FP FP	CORRELAT: RL,FP RL	ION OF FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS -RL	(B2) RL+FP
1234567890112345678901122222222222222222222222222222222222	300 298 (2) 300 300 300 299 (1) 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 300 3	999 1000 1000 1000 1000 1000 1000 1000	 300 300	308 314 315 314 285 287 285 287 285 287 285 287 285 287 285 287 285 285 287 285 285 285 285 285 285 285 285 285 285	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{c} -0.068\\ -0.030\\ 0.006\\ -0.005\\ 0.050\\ -0.027\\ -0.012\\ 0.090\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.088\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.097\\ -0.095\\ -0.095\\ -0.095\\ -0.095\\ -0.095\\ -0.011\\ -0.055\\ -0.095\\ -0.011\\ -0.055\\ -0.055\\ -0.011\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.003\\ -0.055\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ -0.005\\ 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0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2292\\ 0.2392\\ 0.2392\\ 0.2392\\ 0.2392\\ 0.2392\\ 0.2392\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2292\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 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0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.2332\\ 0.23$	0.419 0.402 0.092 0.128 0.012 -0.092 0.173 0.604 0.537 0.648 0.414 0.481 0.301 0.258 0.468 0.811 1.421 1.367 0.205 0.226 0.179 0.242 0.215 0.117 0.597 0.571 0.676 0.667 -0.546 -0.518 0.086 0.211 0.408 0.619 0.533 0.479 0.091 0.260 0.235 0.469 0.143 0.036 0.046 0.200 0.222 0.257 -0.046 -0.091 0.121 0.195 0.037 0.455 0.102 0.260 0.227 0.349 0.112 0.313 0.142 0.436 0.453 0.527 0.258 0.120 0.260 0.276 0.289 0.394 0.227 0.349 0.112 0.313 0.142 0.436 0.453 0.527 0.258 0.120 0.445 0.459 -0.107 -0.207 -0.051 -0.032 -0.130 -0.071 0.161 0.220 0.238 0.325 0.511 0.538 0.325 0.445 0.445 0.459 -0.130 -0.071 0.161 0.220 0.238 0.325 0.511 0.538 0.329 0.445 0.445 0.459 -0.130 -0.071 0.145 0.429 0.332 0.324 0.238 0.325 0.511 0.538 0.329 0.445 0.445 0.459 -0.134 0.272 0.344 0.279 0.332 0.344 0.533 0.469 0.102 0.252 -0.168 0.0998 0.272 0.207 0.209 0.125 0.194 0.304 0.304 0.299 0.010 0.076 -0.070 -0.141	$\begin{array}{c} 4.644\\ 3.250\\ 4.622\\ 4.046\\ 5.626\\ 4.622\\ 4.046\\ 4.803\\ 3.283\\ 3.097\\ 3.908\\ 5.612\\ 5.626\\ 4.622\\ 4.046\\ 4.702\\ 8.033\\ 3.097\\ 3.908\\ 5.612\\ 5.612\\ 5.612\\ 3.097\\ 3.908\\ 3.221\\ 3.546\\ 4.432\\ 3.546\\ 4.432\\ 3.546\\ 3.3546\\ 3.227\\ 3.502\\ 3.709\\ 3.221\\ 3.502\\ 3.702\\ 3.703\\ 3.662\\ 3.101\\ 3.662\\ 3.101\\ 3.662\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.101\\ 3.662\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3.005\\ 3$	4.491 4.315 5.713 5.286 5.713 5.062 3.703 6.354 3.289 4.325 5.062 3.703 6.354 3.304 3.324 4.267 3.316 4.327 3.324 4.325 4.327 3.324 4.325 4.327 3.324 4.325 5.022 3.324 4.327 3.324 4.325 5.022 3.324 4.325 5.022 4.327 3.324 4.325 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 5.022 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JS VISUAL

SESS	NUMBER	nean	SYNCH	nean	VARI	IANCE OF	CORRELA	TION OF	root	B1 OF	KURTOSI	(S (B2)
	TRIALS	Fp	INT	Rl	RL I	RL+FP FP	RL,FP R	L+FP,FP	Rl	RL+FP	•RL	RL+FP
61 62 63 64 65 66 67 68 69 70 71 72 73 74 75	300 300 300 300 300 300 300 300 300 300	1000 1000 999 1000 1000 1000 1000 1000	310 310 310 315 315 315 315 315 320 320 320 320 320 320	310 307 310 309 316 324 317 312 304 317 314 317 320 324 318	489 570 456 360 557 524 522 563 491 562 572 509 608 663 468	555 94 624 81 493. 72 416 95 630 74 577 88 581 105 614 88 558 85 605 82 656 63 569 105 679 91 747 104 508 83	-0.065 -0.062 -0.096 -0.104 -0.002 -0.082 -0.089 -0.082 -0.045 -0.093 0.054 -0.097 -0.042 -0.039 -0.110	0.350 0.300 0.289 0.381 0.340 0.312 0.331 0.300 0.349 0.349 0.349 0.349 0.349 0.349 0.349 0.349 0.349 0.337 0.337 0.337	0.298 -0.141 0.287 -0.052 0.299 0.307 0.290 0.172 0.499 -0.114 0.095 0.277 0.522 0.116 -0.018	0.298 0.091 0.254 0.211 0.416 0.285 0.374 0.374 0.442 0.104 0.201 0.525 0.201 0.138	2.892 4.442 3.299 2.910 3.326 3.444 3.059 3.605 3.931 4.691 3.485 2.843 3.475 3.327 3.923	2.820 4.014 3.078 2.964 3.864 3.316 3.072 3.316 3.756 3.831 2.790 3.450 3.501 3.125

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RU, VISUAL

SESS	NUMBER	nean	SYNCH	nean	VARIANCE OF	CORRELATION OF	ROOT B1 OF	KURTOSIS (B2)
	TRIALS	FP	INT	Rl	RL RL+FP FP	RL,FP RL+FP,FP	RL RL+FP	RL RL+FP
391 393 394 395 397 3990 401 234 402 402 402 402 402 402 402 402 402 40	33000000000000000000000000000000000000	$\begin{array}{c} 1001\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 1000\\ 10000\\ 10000\\ 10000\\ 10000\\ 10000\\ 10000\\ 10000\\ 10000\\ 0000000\\ 0$	300 300 300 300 300 300 300 300 300 300	300 302 302 302 302 302 302 302 302 302	782 844 116 538 570 89 463 460 80 463 460 80 308 391 85 555 553 76 354 377 84 537 562 85 505 529 86 472 468 81 419 439 90 440 476 83 419 454 81 360 409 73 371 415 102 483 532 81 371 382 90 511 485 106 404 460 94 396 436 91 295 349 108 319 388 69 400 415 87 386 429 72 290 355 83 272 343 82 290 355 83 272 343 82 386 417 109 340 396 106 357 386 71 327 388 76 419 467 71 328 343 106 276 342 109 301 308 74 259 312 91 288 358 156 327 453 83 239 316 81 314 337 <	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	5.578 5.367 3.248 2.884 3.422 4.411 3.177 3.194 3.024 3.045 4.093 3.973 2.798 2.919 3.274 3.115 3.819 4.199 2.898 3.204 3.274 3.115 3.819 4.199 2.898 3.206 3.46 3.148 3.159 2.702 2.695 2.704 3.058 2.890 2.498 2.748 2.885 2.702 2.695 2.704 3.056 2.852 3.773 3.361 3.036 2.852 3.179 2.961 3.179 2.961 3.179 2.961 3.179 2.961 3.179 2.961 3.179 3.489 3.227 3.652 3.179 2.961

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RU VISUAL

SESS	NUMBER TRIALS	mean FP	synch Int	nean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL,FP RL+FP,FP	ROOT B1 OF RL RL+FP	KURTOSIS (B2) -RL RL+FP
451 452 453 454 455 456 457 458 460 461 462 463 464 465 466 465 4667	300 300 300 300 300 300 300 300 300 300	1000 1000 1000 999 979 1000 1000 999 1000 1000	350 350 350 350 350 360 365 370 375 380 390 395 400 400 400	353 355 344 348 347 356 370 366 379 372 386 379 372 386 379 382 395 395 400 397 399 407	216 291 103 335 371 90 382 433 94 299 407 87 230 274 73 306 404 97 328 381 90 394 484 94 348 399 72 372 420 76 428 519 85 330 395 78 326 397 90 544 660 103 494 533 87 476 515 67 485 595 89	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	0.216 0.381 0.064 0.229 -0.188 -0.061 0.074 0.623 0.111 0.287 -0.168 0.262 0.638 0.974 0.281 0.333 0.131 0.254 0.450 0.399 0.506 0.636 0.071 0.084 0.265 0.507 0.312 0.405 1.246 0.958 0.181 0.139 0.085 0.111	3.846 4.458 2.787 2.832 3.350 3.632 3.291 4.765 2.775 3.358 2.806 4.020 5.952 7.424 3.453 3.294 3.273 3.299 3.739 3.211 4.341 3.748 3.546 3.350 3.432 4.191 3.598 3.875 10.094 8.049 3.639 3.534 3.162 2.625
469 4711 4773 4774 4775 4778 4778 4777 4778 4812 4885 4887 4887 49901 4975 5012 505 505 505 505 505 505 505 505 505 50	300 300 300 300 300 300 300 300 300 300	1000 1000 1001 1000 1000 1000 1000 100	$\begin{array}{c} 400\\ 400\\ 400\\ 400\\ 400\\ 400\\ 400\\ 400$	404 404 400 400 400 400 27 33 355 33 355 33 355 33 355 355 355 35	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{c} -0.075 \\ -0.072 \\ -0.072 \\ -0.324 \\ -0.070 \\ 0.540 \\ -0.145 \\ 0.330 \\ -0.093 \\ 0.392 \\ -0.050 \\ 0.431 \\ -0.203 \\ 0.254 \\ -0.032 \\ 0.366 \\ -0.145 \\ 0.371 \\ -0.057 \\ 0.411 \\ 0.02 \\ 0.401 \\ -0.142 \\ 0.322 \\ -0.112 \\ 0.424 \\ -0.108 \\ 0.316 \\ -0.132 \\ 0.359 \\ 0.056 \\ 0.474 \\ 0.394 \\ 0.470 \\ -0.67 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.067 \\ 0.461 \\ -0.124 \\ 0.382 \\ -0.049 \\ 0.401 \\ -0.140 \\ 0.331 \\ -0.199 \\ 0.304 \\ -0.103 \\ 0.439 \\ -0.029 \\ 0.430 \\ 0.014 \\ 0.459 \\ -0.029 \\ 0.438 \\ 0.170 \\ -0.073 \\ 0.447 \\ -0.131 \\ 0.138 \\ 0.170 \\ -0.073 \\ 0.445 \\ -0.132 \\ 0.384 \\ \end{array}$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	3.642 $3.9963.247$ $4.0183.940$ $8.7722.776$ $3.4194.242$ $3.3723.582$ $4.9423.327$ $3.1733.705$ $3.3544.010$ $3.5973.884$ $3.9354.078$ $3.8003.056$ $3.5723.070$ $3.3663.861$ $3.6694.275$ $4.1642.986$ $3.4333.103$ $2.8692.746$ $2.7263.737$ $3.6392.655$ $2.7903.861$ $4.2193.246$ $3.6322.578$ $2.9193.165$ $2.7213.602$ $3.7602.865$ $3.6973.417$ $3.6262.903$ $4.2122.742$ $2.9273.429$ $3.9382.749$ $3.0442.707$ $2.6725.217$ $3.2693.136$ $2.7342.943$ $2.2693.136$ $2.7342.943$ $2.2693.2683.274$

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RU VISUAL

SESS	NUMBER TRIALS	Mean FP	SYNCH INT	nean Rl	VARIANCE OF RL RL+FP FP	CORRELATION OF RL, FP RL+FP, FP	ROOT B1 OF RL RL+FP	KURTOSIS (B2) -RL RL+FP
511 512 513 515 516 517 518 521 522 522 522 522 522 522 522 522 522	300 300 300 300 300 300 300 300 300 300	1000 999 1000 1000 1000 1000 1000 1000	340 340 340 340 330 330 330 330 320 320 320 320 320 32	346 343 344 342 331 329 330 328 329 330 328 329 330 328 329 330 328 329 329 329 329 329 329 329 329 329 329	256 309 87 221 296 81 240 276 91 238 282 92 291 338 74 198 293 91 261 287 95 213 255 84 291 323 78 184 284 120 312 370 91 213 249 73 238 281 75 212 252 76 253 296 85 183 257 81 264 317 119 234 279 91 215 295 96 236 265 76 218 271 81 344 371 76 310 373 91 245 284 74 260 288 80	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
542 543 544 545 546 547 548 549	300 300 300 300 300 300 300 300	1001 1000 1001 1000 1000 1000 1000	300 300 300 300 300 300 300	300 295 297 300 302 295 303 298	258 308 106 322 373 96 239 253 73 313 377 87 255 272 69 276 290 91 308 332 87 283 297 80	-0.170 0.432 -0.129 0.387 -0.223 0.319 -0.071 0.415 -0.195 0.315 -0.243 0.323 -0.193 0.327 -0.219 0.305	-0.034 0.112 -0.505 -0.117 -0.289 -0.127 0.178 0.237 0.194 0.050 0.319 0.334 0.142 0.035 0.260 0.236	3.201 3.081 4.408 3.671 3.938 3.589 3.138 3.234 3.891 3.776 4.437 4.377 3.409 3.088 3.992 3.012

Appendix 2

Distributional Information

(a) Notation and Basic Theorems

Let f(x) be the density function of a random variable X and denote the expected value of a function, g(), of X by E(g(X)). By definition (eq. Freund, 1971)

 $E(g(X)) = \Sigma g(x) * f(x) \quad \text{if } X \text{ is discrete}$ $= \int g(x) * f(x) dx \quad \text{if } X \text{ is continuous.}$

Let μ be the population mean. For discrete distributions, the second, third and fourth central moments about the mean are defined as:

 $\mu_{2} = \Sigma(x-\mu)^{3} + f(x) = V(X) \quad (Variance)$ $\mu_{3} = \Sigma(x-\mu)^{3} + f(x)$ $\mu_{4} = \Sigma(x-\mu)^{4} + f(x).$

The definitions for continuous distributions are the same except for replacement of discrete summation by integration.

The population coefficients of symmetry, $\sqrt{\beta_{N_{1}}}$ and kurtosis, β_{2} are defined as:

 $\sqrt{\beta_1} = \mu_3 / V(X)^{3/2}$ $\beta_2 = \mu_4 / V(X)^2.$ Symmetrically distributed variables have $\int \beta_1 = 0$. Positively skewed distributions (such as the geometric, exponential and chi-square) have positive symmetry coefficients.

Distributions with thin tails (data points many standard deviations from the mean are extremely unlikely) have low kurtosis coefficients. As extreme values become more likely, the kurtosis increases.

From Johnson and Kotz (1970) and Hastings and Peacock (1974)we obtain the following information:

Distribution	Variance	Coefficient of Symmetry	Coefficient of Kurtosis	4th Central Moment	
Triangular	,q2/6 A	0	2.4	q*/15	
Normal	ر 2	0	3.0	30"	
Logistic	P5	0	4.2	4.25*	-
Geometric	r/p2	(2-p)/(r	9+p²/r՝	9r²/p+ + c/p²	(r≢1-p)

Let X and Y be two random variables and let x and y be observed values of X and Y respectively. The variable Z is called the sum, or convolution, of X and Y when observed values, z, are formed by taking z = x+y.

Most of the work below will involve convolutions of two or more of the four distributions above. The following two theorems will be useful:

Expectation of a Product (Freund, 1971, p. 179)

If X and Y are statistically independent then E(X * Y) = E(X) * E(Y).

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Covariance of a Linear Combination (Freund, 1971, Theorem 6.4, p. 196)

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If X_1, X_2, \ldots, X_n are random variables and c_1, c_2, \ldots, c_n and d_1, d_2, \ldots, d_n are constants and $Y_1 = \Sigma c_1 X_1$, and $Y_2 = \Sigma d_1 X_1$, then the covariance of Y_1 and Y_2 , COV(Y_1, Y_2), is given by:

 $COV(Y_1,Y_2) = \Sigma c_1 d_1 V(X_1) + \Sigma \Sigma (c_1 d_1 + c_1 d_1) * COV(X_1,X_1).$

Finally, the geometric variable of interest in the present work is the foreperiod duration. On the PDP-8 geometric variates were generated with a mean of 10 msec (plum 890), in units of 1 msec. On the Apple, geometric variates were generated in 200 microsecond units, with p = .0208608. Converting to milliseconds, the foreperiod distributions of the present research had the following statistics:

	Mean	Variance	β 1	β2	۴з	μu
Apple	9.5874	90	2.00011	9.00044	1707.78	72903.6
PDP-8	10.0000	90	2.00278	9.01111	1710-00	72990.0

I will use the Apple values-below.

(b) Symmetry and Kurtosis of the Sum of Independent Random Variables Let X and Y be independent random variables and let Z be their convolution, Z = X+Y. Let the means of X, Y and Z be denoted μ_x , μ_y and μ_Z respectively. Since X and Y are independent, the variance V(Z) = V(X) + V(Y). By definition, the symmetry coefficient is $\overline{\beta_1} = \mu_3/V(Z)^{3/2} = EI(Z-\mu_Z)^{3}/V(Z)^{3/2}$. Focusing on the numerator, we have $E[(Z-\mu_Z)^3] = E[(X+Y-\mu_x-\mu_y)^3]$

 $= E[(X-\mu_{x})^{3}] + 3E[(X-\mu_{x})(Y-\mu_{y})^{2}] + 3E[(X-\mu_{x})^{2}(Y-\mu_{y})] + E[(Y-\mu_{y})^{2}]$

Since X and Y are independent, so are $(X-\mu_x)$ and $(Y-\mu_y)^2$, as are $(X-\mu_x)^2$ and $(Y-\mu_y)$. And since $E[X-\mu_x] = E[Y-\mu_y] = 0$, we have $E[(X-\mu_x)(Y-\mu_y)^2] = E[X-\mu_x] * E[(Y-\mu_y)^2] = 0$, and $E[(X-\mu_x)^2(Y-\mu_y)] = E[(X-\mu_x)^2] * E[Y-\mu_y] = 0$. Thus, $E[(2-\mu_z)^3] = E[(X-\mu_x)^3] + E[(Y-\mu_y)^3]$, and $\sqrt{\beta_1} = (E[(X-\mu_x)^3] + E[Y-\mu_y)^3])/(V(X)+V(Y))^{3/2}$.

Substituting X+Y for Z and dropping terms that reduce to zero, for <u>Kurtosis</u> we have:

 $\beta_{2} = E[((X - \mu_{x}) + (Y - \mu_{y})) +]/(V(X) + V(Y))^{2}]$

= $(E[(X-\mu_x)^{*}] + E[(Y-\mu_y)^{*}] + 6E[(X-\mu_x)^2(Y-\mu_y)^2]]/(V(X)+V(Y))^2$

= $(E[(X-\mu_x)^+] + E[(Y_{e}+\mu_y)^+] + \delta V(X)V(Y_{i})/(V(X)+V(Y_{i}))^2$.

(c) The Sum of a Triangular, and a Normal Variable

Kristofferson (1976) proposed that efferent latency variability could be described as normal and that the synchronization latency distribution was the convolution of independent triangular and normal variables. Hopkins (1982) originally used the logistic as an approximation to the normal, though the logistic Kurtosis is substantially larger than the normal's. From the equation above, the Kurtosis coefficient of this distribution is

 $\beta_2 = \{2.4q^{4}/36 + 3\sigma^{4} + q^2\sigma^2\}/\{q^2/6 + \sigma^2\}^2.$

This has a minimum of 2.4 when the normal variance is zero and

a maximum of 3 when the triangular variance is zero. It increases monotonically between these two extremes. (I have checked this by examining the higher derivatives of the function.)

The kurtosis coefficients of the present data were generally larger than 3. In all of the experiments, averages across 5 sessions of the kurtosis coefficients were almost always larger than 3. This distribution does not describe the present data.

(d) <u>Convolution</u> of a Triangular and a Logistic Variable

Hopkins (1982) derived the density and distribution functions of this convolution. Unfortunately, there were two copying errors in his final equations for the cumulative distribution function. The integrations involved were not easy, so a final corrected formula should be made available. I have checked the following expression with him (Hopkins, personal communication, November 1, 1982) and by simulation. Let Z be the convolution of the two variables and F(Z) be its cumulative distribution function. Then, in the notation of Hopkins (1982),

F(z) = 1 + A + B + C, where

 $A = (k/w^2) \int \ln(1 + \exp((a-z)/k) dz$

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 $B = (k/w^2) \int \ln(1 + \exp((2w + a - z)/k) dz, and \forall$

 $C = (2k/w^2) \int \ln(1+\exp(w+a-z)/k) dz$

Translating notations, w = q, the quantum size, a is the logistic mean, and $k = \sqrt{3}b/\pi$, where b is the logistic standard deviation. For evaluation of the integrals, see Hopkins (1982, p.

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The variance of this distribution is $b^2 + q^2/6$. It is symmetrical. From the formula above for the Kurtosis of a sum of two variables, we obtain

 $\beta_2 = 2.4(q^4 + 63b^4 + 15q^2b^2)/(q^2 + 6b^2)^2$.

For later reference, the fourth central moment can be obtained by multiplying the Kurtosis by the square of the variance of the convolution. It is

 $\mu_{4} = (q^{4} + 63b^{4} + 15q^{2}b^{2})/15.$

The kurtosis is between 2.4 (when the logistic variance is zero) and 4.2 (when the triangular variance is zero). Let V(T) be the triangular variance and V(L) be the logistic variance. The kurtosis of the convolution is a strictly monotonically increasing function of V(L)/(V(T)+V(L)). The kurtosis is independent of scale. That is, suppose that you multiply q and b by a non-zero constant, c. The variances V(T) and V(L) will each change by a factor of c^2 . (If V(T) was the variance before multiplication, the new variance is $c^2V(T)$.) The numerator and denominator of the expression for kurtosis both change by a factor of c^4 and this constant factor divides out. The kurtosis depends only on the relative values of V(L) and V(T). If V(L)/(V(L)+V(T)) is constant, so is the kurtosis, whatever the value of V(L).

In Chapter 3 I tabled (Table 3) some examples of the Kurtosis of the logistic-plus-triangle convolution as a function of $V(L)/(V(L)+V(\bar{T}))$ (which I called b²/V there). The values were obtained by setting V(L) to an arbitrary constant (100, except when the ratio was Q), adjusting V(T) to obtain ratios of 0.1, 0.2, etc., and computing the Kurtosis from the formula above. Since the Kurtosis is independent of scale, any other values of V(L) would have yielded the same results.

In Chapter 6 I estimated logistic variances of Kristofferson's (1976) data from Kurtosis coefficients and overall variances of the latency distributions. This was done iteratively. The Kurtosis of the logistic-plus-triangle convolution is strictly monotonically increasing with V(L)/(V(L)+V(T)). By virtue of monotonicity, there is a one-to-one relationship between Kurtosis and this variance ratio. I wrote a short computer program that tried different variance ratios until it found one that yielded a Kurtosis value extremely close to the sample kurtosis. This yielded, then, an estimate of V(L)/(V(L)+V(T)). According to Hopkins' model, synchronization latency variance is V(L)+V(T), the variance of a logistic variable plus that of a triangular variable, so by multiplying the estimated ratio V(L)/(V(L)+V(T)) by the response latency variance (V(L)+V(T)), we obtain an estimate of V(L), the logistic variance.

The estimate of V(L) so obtained is a "method of moments" estimate. This is an old and intuitively reasonable procedure but there is no guarantee that it is unbiased and such estimates, when they involve high moments, are often quite variable. Hopkins (1982) provided kurtosis coefficients, sample variances, and minimum chi-square estimators of the logistic variances of some distributions. He provided me with a few further groups of estimates (persona) communication). I compared his estimates with those obtained in the way outlined above. The two estimation procedures yielded different
values for the logistic variance but there was no tendency for one to yield higher values than the other. Minimum chi-square estimation may (or, given the flatness of its parameter space, as discussed in Chapter 3, may not) yield lower variance estimates of the logistic variance but it appears that with samples of 1500 observations (5 sessions) the two estimation procedures will on average yield the same estimates. (The sample size is important because the bias of a consistent estimator diminishes as the sample size increases. The distributions chosen from Kristofferson (1976) all involved at least 5 sessions.)

(e) <u>Timing From WS</u> or From P1

First, suppose that the subject times from P1. Let random variable ZP1 be the interval between P1 onset and the completion of the response and suppose that ZP1 is distributed as the convolution of two independent random variables, one logistic, the other triangular. Let the duration of the interval between the offset of the warning stimulus (WS) and the onset of P1, i.e. the foreperiod duration, be represented by FP. FP is geometrically distributed with a known parameter. Because timing units were 1 msec on the PDP-8 and .2 msec on the Apple, the geometric parameter differed across computers, resulting in very small differences in the shape of the geometric distribution on each machine. In what follows I will use the Apple values since most sessions were run on the Apple.

The interval from WS offset to the completion of the response is the sum of two independent (\underline{ex} <u>hypothesis</u>) random variables, ZP1 and FP.

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 $V(ZP1 + FP) = V(ZP1) + V(FP) = q^2/6 + b^2 + 90$. From Section (b) of this appendix, the symmetry coefficient is

 $\int B_1 = (EI(ZP1-EIZP1J)^3] + EI(FP-EIFPJ)^3J)/V(ZP1+FP)^{3/2}$ where EIZP1J and EIFPJ are the respective means of ZP1 and FP. ZP1 is
symmetrical so its third central moment is 0. The third central moment
of FP, as tabled in Section (a) of this appendix, is approximately
1708. Thus

 $\int B_1 = 1708/V(ZP1 + FP)^{3/2}$.

The Kurtosis of the sum of ZP1 and FP can also be found using the general Kurtosis formula of Section (b) and the Kurtosis and 4th moment of the sum of a logistic and a triangle obtained in Section (d). The calculations are straightforward but longer than those for symmetry. The final result is

 $\beta_2 = \frac{(q^4 + 63b^4 + 15q^2b^2)/15 + 90(q^2 + 6b^2) + 72904}{(b^2 + q^2/6)^2 + 180(b^2 + q^2/6)^2 + 8100}$

This is an unwieldy result. It may be worth noting that I have checked it (and almost all of the others in this appendix) by simulation.

These are the statistics of the distribution of response latencies obtained if the subject times exclusively from P1, with logistic-plus-triangular variability, and if we measure the response latency from the offset of WS.

Now suppose that the subject times from WS offset instead of from P1 onset. Let ZWS represent his response latency, i.e. the interval from WS offset to completion of the response, and suppose that ZWS is distributed as the sum of a logistic and a triangular variable

(independent), just as we supposed for ZP1. Finally, suppose that the subject's timing is uninfluenced by the foreperiod duration, i.e. that ZWS and FP are independent.

If we measure the response latency relative to P1 in this case, on each trial we are subtracting the foreperiod duration from ZWS to obtain the final latency estimate (re P1): That is, we are measuring ZWS = FP.

All of the theorems above refer to addition of random variables rather than to subtraction. It is easy enough to recast this in terms of addition. Define a new variable, MFP, such that MFP = -FP. For example, if the foreperiod on a given trial is 999 msec, the value of MFP for that trial is -999. Multiplication by -1 has no effect on variance or kurtosis. The variances of MFP and FP, and their Kurtoses, are the same. The symmetry coefficients of FP and MFP are the same in absolute magnitude, but they differ in sign. The symmetry coefficient of FP is approximately 1708. The symmetry coefficient of MFR is -1708.

 $V(ZWS-FP) = V(ZWS+MFP) = b^2+q^2/6$.

This formula is identical to that for V(2P1+FP) but I do not mean to imply here that the logistic (b²) and triangular (q²+8) variances are identical when the subject times from P1 and when the subject times from WS. The values of q and b may differ in the two cases. Given the values of q and b, the <u>expressions</u> for computing V(2WS-FP) and V(2P1+FP) are the same.

The symmetry coefficient for 2WS-FP is $-1708/V(2WS-FP)^{3/2}$, the same as that for ZP1+FP except for the reversal in sign.

The expressions for Kurtosis of 2P1+FP and 2WS-FP are the same.

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The formulas in this Section are the basis for most of the entries in Table 5 of Chapter 4, which described the shapes of latency distributions when timing was from WS or from P1 and latencies were measured relative to WS or P1. Tables 5 and 6 and other sections of Chapter 4 also made reference to correlations between FP and response latency under the two models above. These are dealt with in the next and final section of this appendix.

(f) <u>Response Latency and Foreperiod Correlations</u>

Suppose that the subject times from P1, and let ZP1 be defined i as in the previous Section. ZP1 is triangular-plus-logistic and is assumed independent of FP, the foreperiod duration.

Let COV(X,Y) be the symbol for the covariance of two random variables, X and Y and denote their correlation by CORR(X,Y). Since ZP1 is independent of FP ex hypothesis,

COV(ZP1,FP) = CORR(ZP1,FP) = 0.

If we measure response latencies from WS offset rather than from P1 onset, and correlate these with foreperiod duration, we are computing the correlation between ZP1+FP and FP. Casting this in terms of Freund's theorem 6.4, presented in Section (a) of this appendix, let $Y_1 = ZP1+FP$ and let $Y_2 = FP$. Then

 $COV(ZP1+FP,FP) = COV(Y_1,Y_2)$

 $= 0 \times V(ZP1) + 1 \times V(FP) + COV(ZP1,FP)$

= V(FP)

Thus $CORR(ZP1+FP,FP) = V(FP)/(V(ZP1+FP)*V(FP))^{1/2}$

 $= (V(FP)/V(ZP1+FP))^{1/2}$.

Now, suppose instead that the subject times from WS rather than from P1, represent the subject's response latency (from WS offset to response completion) by ZWS, assumed to be the sum of a logistic and a triangular variate, independent of FP. <u>Ex hypothesis</u> CORR(ZWS,FP)=0.

If we measure response latencies from P1 rather than from WS, we are measuring ZWS-FP and the correlation of these measured latencies with the foreperiod durations is CORR(ZWS-FP,FP). Freund's theorem 6.4 is again directly applicable, with the same result as above except for a reversal in sign:

 $CORR(ZWS-FP,FP) \cong (V(FP)/V(ZWS-FP))1/2$.