

CONTINUOUS ROTATING BIOLOGICAL CONTACTOR
FOR DENITRIFICATION

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FOR DENITRIFICATION

By

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ABSTRACT :

This work presents the results of a pilot plant study where a completely submerged rotating biological contactor was used for denitrification of domestic wastewater.

The literature review emphasizes the existing knowledge on RBC operation and design fundamentals. A brief summary of the theory of denitrification has been presented as this subject already has been presented in detail by various authors.

Dye studies were performed for 4 stage series operation and single stage operation. A two CSTR with interchange flow model seemed to fit the hydraulic model for single stage operation. The dye response was observed

to be function of biological growth which in turn varied with temperature.

"Zero order" kinetics were shown to represent the data obtained for denitrification rates at hydraulic loadings of 2.5 l/min and 4.0 l/min best for the range of $\text{NO}_3^- + \text{NO}_2^-$ -N concentrations studied.

An energy of activation of 16,500 kcal/mole was calculated for hydraulic loading range of 2.5-4.0 l/min.

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ABBREVIATIONS AND SYMBOLS

A_o : frequency factor
BOD : biochemical oxygen demand
BOD₅ : 5 day biochemical oxygen demand
BST : batch stirred tank

°C : degrees Centigrade
cal/mole: calories per mole
cm : centimeter
COD : chemical oxygen demand
CSTR : continuous stirred tank reactor

DO : dissolved oxygen
E : activation energy
Et : dimensionless residence time distribution
F : area under RTD curve
°F : degrees Fahrenheit
ft : feet
ft² : square feet
g : gram
gpd : gallons per day
hr : hour
hrs : hours

K : reaction rate or reaction rate constant for
 $n=0$ or $n=1$, respectively

$^{\circ}K$: degrees Kelvin

l : liter

l/min : litre/minute

MLSS : mixed liquor suspended solids

MLVSS: mixed liquor volatile suspended solids

m : meter

m^2 : meter square

mg : milligram

min : minute

mg/l : milligram per litre

mm : millimeter

NO_2-N : nitrite nitrogen

NO_3-N : nitrate nitrogen

NH_3-N : ammonia nitrogen

pH : logarithm of hydrogen ion concentration

ppb : parts per billion

R : gas constant

RTD : residence time distribution

rpm : revolution per minute

SS : suspended solids

SRT : sludge retention time

T : temperature

TKN : total Kjeldahl nitrogen

TOC : total organic carbon

t : time

θ : dimensionless time
 α : confidence level
 \int : integral
% : percent
 $>$: greater than
 $>>$: considerably greater than

1. INTRODUCTION

The Rotating Biological Contactor (RBC) has in the past been used for removal of carbonaceous BOD extensively. In addition, some investigators have proposed its use for nitrification and algae nutrient removal. Only one reported attempt, Pretorius (1971) has been located that uses the Rotating Biological Contactor (RBC) system for denitrification.

Although the Rotating Biological Contactor treatment system is basically a biological treatment system and the basic principles of biological treatment have been thoroughly studied, the exact mechanisms of removal by the RBC is not known. Different design techniques have been proposed by different authors.

The aim of this study is to review existing knowledge about RBC operation and design formulation, and to study the effects of various operating variables (temperature, hydraulic loading, and mode of operation) on denitrification by the completely submerged RBC.

2. REVIEW OF EXISTING LITERATURE

Since basic principles of biological treatment for carbonaceous BOD removal and nitrification have been well documented by various investigators and authors, they will not be repeated. Rather emphasis will be placed on the Rotating Biological Contactor.

2.1 Rotating Biological Contactors

2.1.1 The Definition and Physical Description of RBC

Partially Submerged RBC Systems

The RBC unit is a series of discs closely spaced on a shaft which is rotated just above the surface of waste liquid to be treated. The discs are believed to perform five specific functions (Antonie, 1974):

- a) provide a large amount of available surface area for the development of a fixed biological growth,
- b) provide contact of this growth with the wastewater,
- c) efficiently aerate the wastewater by carrying a thin film of liquid from the tank,
- d) provide a controlled, mechanical means of stripping excess biomass from the surfaces, and

- e) provide thorough mixing of each stage of treatment.

Various disc diameters are being used in practice; Antonie (1974) employed a unit of 3.7 m, Steels (1974) used diameter of 3.0 m, while Bruce (1973) described a package plant developed for small communities with 1.0 m disc diameter. McAliley (1974) used a diameter of 3.7 m while Summer (1973) and Antonie (1970) indicated that discs up to 3.0 m diameter are available. Chittenden (1971) described his system consisting of 3.3 m discs. It is apparent that the diameter varies with the company producing the RBC although 3.0 m diameter is normal except for specific package plants.

The typical disc spacings used in practice have been reported by Chittenden (1971) who indicated that typical center to center distance of discs to be 2.5 to 3.8 cm. Bruce (1973) indicated that a 10.0 mm gap between adjacent surfaces for a package plant was used while Antonie (1970) employed 2.5 cm center to center spacing for a package plant of 1.8 m diameter discs. Normally the disc spacings are fixed by the manufacturers of the RBC.

The third property fixed by the manufacturer is the rotational speed; Beak (1973) gave 1 rpm as typical for a 3.0 m diameter disc. One rpm for a 3.0 m diameter disc was also quoted by Steels (1974), and Bruce (1973) for a package plant.

It must be concluded that various combinations of different disc diameters, speeds (rpm's) and disc spacings are possible depending upon the specific field of applications.

Totally Submerged RBC Systems

This type of system is employed when aeration is not essential and/or desirable. The current literature does not give any application of this type of RBC.

2.1.2 Historical Development and Various Applications

A rotating disc was first used by Doman (1929). Maximum BOD₅ removal achieved was 27%. The apparatus that he used was a series of partially submerged discs rotated at 0.5 rpm. The detention time was 2.4 hrs and maximum microbial film thickness was less than 0.8 mm.

Thirty years passed before the RBC was again used to treat wastewater. Grieves (1972) in his literature survey stated that, Hartmann had experimented extensively with a rotating disc plant in 1958. The following conclusions were reported:

- a) the optimal rpm was 5 to 7 for a 1.0 m diameter disc,
- b) the optimum disc spacing was at 15.0 mm,
- c) the discs should be submerged as deeply as possible,

- d) best performance occurred when the disc rotation was in the direction of the wastewater flow,
- e) staging was advisable from a "construction view point", but did not affect efficiency,
- f) the system was especially suited to small plants, because of its low maintenance costs and insensitivity to fluctuations in load, and
- g) the operational cost was "considerably less" than those for activated sludge or trickling filter processes.

Grieves (1972) cited further contributions by Hartmann and Popel in Europe around the mid-sixties. Most of the existing literature in the English language appeared after 1968.

2.1.3 Mechanism of Removal by RBC

After commencement of operation with any substrate, a layer of microbial film develops on the surfaces of the discs. The film is submerged in the substrate to absorb food and emerges into the air to absorb oxygen. During submersion of the disc, oxygen is transferred from the liquid film to substrate and substrate removal by the film continues. Hartmann (Steel, 1974) believed that the organisms which are suspended in the liquid also aid in substrate removal.

2.1.4 Factors Affecting the Performance of RBC

To describe mechanistically the performance of any RBC system would be cumbersome. The success of any such description is doubtful, since there are many independent and interdependent operational variables which affect the ultimate performance of system.

The main factors that effect the performance of an RBC system under steady state conditions are:

I) PHYSICAL FACTORS:

- a) the temperature,
- b) the physical dimensions of discs (disc diameter, disc spacings),
- c) the rotational speed of discs,
- d) the physical dimensions of the tank,
- e) the physical configuration of the tank,
- f) the inlet and outlet arrangements,
- g) the oxygen content of surrounding atmosphere, etc.,
- h) the hydraulic loading
- i) the existence of staging, and
- j) the temperature of substrate.

II) CHEMICAL AND BIOLOGICAL FACTORS:

- a) the type of substrate, and
- b) the concentration of substrate.

The following variables should be added to the above list when non-steady state performance is being evaluated:

- a) the pattern of change in the hydraulic loading with time,
- b) the pattern of change of the organic loading to the system, and
- c) the change of temperature of air and temperature of incoming waste.

As one can visualize, there are many factors affecting the performance of any RBC system. That is why many authors resort to an experimental evaluation by changing some of the important operating variables and observing the response in the RBC performance. Few authors have attempted to develop an overall formula for RBC systems under steady or non-steady operation.

PHYSICAL FACTORS

Temperature

Since the RBC treatment system is basically a biological treatment system, it is not surprising that it is temperature dependent.

Many investigators have studied the effects of temperature on the performance of the RBC. Antonie (1970) as a result of his pilot plant study, reported that for any specified operating conditions, as the wastewater temperature

was reduced, the performance also was reduced. This reduction was not significant between 10°C and 21°C but was drastic below 10°C . Similar trends have been observed for Kjeldahl and $\text{NH}_3\text{-N}$ removal and the author suggest the use of higher residence times to achieve the desired performance levels.

Antonie (1974) in his introduction of the "Bio-Surf" process as an $\text{NH}_3\text{-N}$ removal device reported that the process efficiency was reduced drastically below 55°F . For 95% removal, Q_{10} (between $55\text{-}45^{\circ}\text{F}$) is given as 1.70.

Pretorius (1974) reported that the energy of activation was equivalent to 3658 cal/mole giving Q_{10} (between $10^{\circ}\text{C}\text{-}20^{\circ}\text{C}$) equal to 1.25. This was based upon a laboratory study of nitrification by RBC system.

Davies and Pretorius (1975) studied denitrification by RBC systems. Their anaerobic unit was only partially submerged. They reported 3 different temperature sensitivity zones for denitrification:

$$Q_{10} = 13.06 \text{ for } T < 10^{\circ}\text{C}$$

$$Q_{10} = 1.38 \text{ for } 10^{\circ}\text{C} \text{--} 30^{\circ}\text{C}$$

$$Q_{10} = -2.65 \text{ for } T > 30^{\circ}\text{C}$$

In another pilot plant nitrification study, Wilson (1974) reported a Q_{10} of 2.36 between 10°C and 20°C , and an energy of activation equal to 14,200 kcal/mole.

Oxygen Availability

Authors have different thoughts as to whether or not oxygen enrichment of the atmosphere surrounding the discs helps the performance of the RBC.

The first attempt to enrich the atmosphere to which the discs are exposed, was made by Torpey and his co-workers (1972). Using a 10 stage laboratory unit, they observed the detention time needed for 90% removal of BOD_5 was reduced to 18 minutes, when oxygen enrichment was applied to the first stage of their system. When the first and third stages of the 3 operating stages were enriched, they observed that substantial reduction in NOD was achieved.

No attempts have been made in practice to enrich the atmosphere around the discs to increase the performance. Hartmann (Beak, 1973) believed that oxygen transfer to the biomass is not a limiting factor, and claimed that oxygen enrichment would not be advantageous.

A second study was made by Pretorius (1974) with an enriched surrounding atmosphere to observe its effects on nitrification. For fixed operational conditions, the following results were obtained:

- a) enrichment to 40% oxygen increased the reaction rate 1.75 times,
- b) at 60% oxygen, the reaction rate was 2 times higher, and
- c) at higher values, increases in the reaction rate levelled off.

There does not appear to be sufficient evidence concerning the relative effect of oxygen enrichment in comparison to the other factors affecting the RBC's performance to justify a conclusion. In addition, no attempts have been reported .. whether or not it is feasible to enrich the atmosphere.

Rotational Speed

The important effects of rotational speed will be on:

- a) the oxygen transfer from the atmosphere to the disc,
- b) the oxygen transfer from the atmosphere to the liquid,
- c) the oxygen transfer from the disc to the liquid,
- d) the substrate transfer from and to the disc, from and to the bulk liquid, and
- e) the organisms.

Many authors have investigated the effects of rotational velocity for their RBC systems performance.

Antonie (1970) as a result of a prototype application showed that, an increase in rpm from 2 to 3.2 increased the percentage of BOD, and NH₃-N removal. A change from 3.2 to 4.6 rpm did not increase the removal efficiency for either NH₃-N or BOD. The COD and SS removal percentages did not show any particular change with increased rpm but increases in rpm resulted in a drastic power consumption increase.

Antonie's observation indicated that there is a certain optimum level of rpm as far as BOD_5 and NH_3-N removal rates concern. His data indicated that it is approximately 3 rpm. This would be specific for only the particular system employed and could be different for other systems or other operating conditions.

In the literature, another optimum value of 5-7 rpm was given by Hartmann (Grieves, 1972) for his 0.90 m diameter disc.

Weng and Molof (1974) studied the effects of rpm values on nitrification with a laboratory scale disc unit. They obtained no nitrification at 10 and 20.5 rpm but at 30 rpm, the NH_3-N concentration was reduced from 26.6 to 22.7 mg/l. When the rpm was increased from 30 to 42, NH_3-N decreased to 17.1 mg/l in the last stage. The authors attributed the cause for the lack of nitrification at 10 and 20 rpm to the presence of excess COD which they claimed had a toxic effect on nitrification.

Another study on the effect of rpm on performance was presented by Welch (1968). The author employed a pilot scale RBC unit on a synthetic sewage. He claimed there was a definite improvement if the rpm were increased for a given fixed hydraulic and organic loading. The author failed to explain whether this increase was due to an increase in D.O. level or due to increased mass transfer.

These studies indicate that there is a definite relationship between rpm and performance values. Any laboratory or pilot plant procedure which is to be used for generation of information for actual full size plant should be operated at optimum rpm values.

Hydraulic Loading

The hydraulic loading is one of the variables which has been most thoroughly studied and is one of the most important design parameters for RBC systems.

Welch (1968) in his pilot-scale study of a synthetic sewage prepared from dairy waste, found that the hydraulic loading affected the plant performance under fixed rpm and influent COD concentrations. Antonie (1970) observed a similar trend for BOD removal, Kjeldahl and $\text{NH}_3\text{-N}$ removal, and COD removal for domestic sewage. The data examined from both of these references indicated that, to increase the detention time above a certain value does not lead to further improvement.

Similar observations were made by Summer and Bennett (1973) as a result of their pilot plant study on pulp, paper and fireboard effluents, Gillespie *et al* (1974) from a pilot-scale RBC study on pulp and paper wastes, and McAliley (1974) from a pilot study on unbleached kraft mill waste.

Staging of RBC Units

One of the first comments on the effects of staging was by Hartmann (Grieves, 1972). It was indicated that staging was advisable from "a construction viewpoint", but did not affect efficiency.

In a translation of Hartmann's paper by Steels (1976), the author explained how to design a two stage treatment plant for domestic sewage and pointed out that if the stage number is more than 2, a correction factor should be applied to reduce the required area needed.

Antonie (1970) in his description of the "Bio-Disc" process, claimed that increasing the number of stages, increased the efficiency. Since, in his belief, BOD reduction by the RBC process is concentration dependent, staging should help the system to operate as plug flow reactor.

Joost (1969) claimed that biochemical reaction is concentration dependent following a "first order" equation, therefore, staging would lead to increased economy.

NCASI (1974) reported, that for certain types of wastes at fixed hydraulic loadings, as the number of stages was increased, the cumulative removal increased. Beyond a certain number of stages, it was observed that the removal curve levels off. An interesting observation is that 6 stages at $2 \text{ gpd}/\text{ft}^2$ and 4 stages at $3 \text{ gpd}/\text{ft}^2$ gave the same performance. Therefore a reduction in the hydraulic loading

to a fixed number of stages may do the same job, as increasing number of stages at a fixed hydraulic loading.

Aacken and Antonie (1971) claimed that the most effective use of disc surface area was achieved in successive rows or stages of discs. They stated that this kind of operation resulted in an improved residence time distribution which yielded a greater BOD removal rate. Further it allowed the development of a mixed culture in each stage that is especially suited to treating the wastewater present in that stage.

Torpey et al (1972) claimed that another advantage of staging was that it separated the carbon oxidation phase and nitrification phase thus giving the most efficient results.

Hydraulic Surging and Organic Shock Loading

This is another parameter which has been examined by different authors for their particular systems.

Antonie (1970) simulated the flow pattern of the influent to a pilot plant, from an industrial plant operating 8 hrs and giving an intermittent flow, with constant concentration. During the early hours of the operating period, the quality of the pilot plant effluent was poor. It took several hours to reach a steady performance level. When a small recycle was maintained during the 24 hour period, the removal results were

improved.

Popel according to Steels (1974) reported that for a peak load which was 4 times the average load and with a continued duration of 2.5 hrs, the BOD removal rate was higher than it would have been for an uniform load, and that if the loading was extended to 5 hrs instead of 2.5 hrs, the results were worse but still better than the uniform load efficiency.

Popel's observations can be explained if at uniform conditions, the bacteria were not working at their maximum capacity, so it would be possible to remove more substrate in the same system when the peak loading was applied. It would be useful if a parameter which could show the optimum substrate removal capacities of the RBC system could be developed.

A hydraulic surge study conducted by Marki and Antonie (1970) indicated that a RBC system subjected to a ten-fold increase in hydraulic loading for 8 hrs each week still gave significant BOD removals during the shock loading periods. A rapid recovery to steady state condition was observed.

In another study reported by NCASI (1974) on pulp and paper wastes, it was observed that when a hydraulic loading which was 4 times the original loading and of 4 hrs duration was applied, the effect was much less than when influent BOD concentration was raised 4 times the

original. Recovery occurred over a shorter period and there was continuing BOD removal activity. Again, the total BOD removal rates were higher than the steady state removal rates. This last observation by NCASI (1974) and Popel's observation are in agreement.

2.1.5 Substrate Removal Kinetics

Joost (1969) claimed that "a first order" reaction is applicable and thus staging should prove beneficial and economical. He gave the following general formula for the RBC system performance:

$$\% \text{ BOD}_5 \text{ Reduction/Stage} = K \times C^a \times R^b \times T^c \times S^d \quad (1)$$

where: K is the treatability constant,
C is the concentration of waste material,
R is the physical configuration of system
T is the temperature of waste,
S is the reactor residence time, and
a, b, c, and d are exponents.

The practical range of applicability was not discussed in detail:

A design method, applicable to a particular domestic sewage, has been developed by Hartmann (Steels 1974). The author has presented design graphs. By knowing only the inlet BOD and using these graphs, the disc area/flow rate required for desired BOD removal efficiencies can be found. The above solution is applicable only for RBCs with two stages. The author also gives factors if more than 2 stages

are to be used, and if variations in either hydraulic or organic loadings are anticipated.

It should be emphasized that this method is applicable only, to Germany conditions. Similar methods or charts could be developed for any local conditions.

Another attempt to develop a design method based on pilot-plant data was made by McAliley(1974). He presented the following formula for the specific removal rate:

$$\frac{F(S_i - S_o)}{A} = P \frac{S_o}{K + S_o} \quad (2)$$

where: F is the flow rate,

S_i is the influent substrate concentration,

S_o is the effluent substrate concentration,

A is the area respectively, and

P and K are the constants to be developed from pilot-plant work.

The above equation indicates the existence of two different rate dependency regions where $K \gg S_o$ the expression is first order and where $S_o \gg K$ it is zero order. The above method seems to be applicable for design of RBC systems, but more work in this direction is needed to show its validity. Further, one can anticipate scale up problems resulting from the applicability of P obtained from pilot plant work to predict the actual full scale removal rate.

McAliley introduced another useful concept in RBC design, that is the oxygen limiting point of operation where oxygen transfer rate is less than the reaction rate.

Grieves (1972) attempted to develop a steady state and dynamic model to represent the RBC performance. He tried to demonstrate the validity of his model by concurrent laboratory studies. He found that the experimental results and results predicted by the model gave good conformance. NCASI used Grieves' model without success but they indicated that the failure could have been caused by scale up difficulties.

3. A SHORT REVIEW OF ESSENTIALS OF DENITRIFICATION

Biological denitrification is the last step of biological nitrogen removal from wastewaters following the nitrification step.

Biological denitrification is basically an anaerobic process employing heterotrophic micro-organisms. These organisms utilize the NO_3^- as a hydrogen acceptor to derive energy for their activities, in the presence of an acceptable organic energy source.

The various mechanisms involved in the biological denitrification process has been reviewed by various authors and these reviews will not be repeated as excellent literature reviews are already available. Rather the theory and the important results of the basic denitrification processes will be presented.

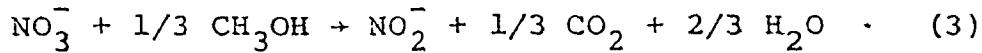
3.1 Basic Reactions Involved in Denitrification

3.1.1 Reactions:

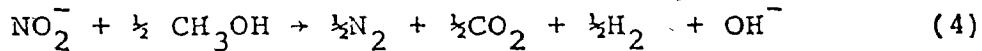
Denitrification can be defined as a two step process. Methanol is generally used as a carbon or energy source.

The denitrification reactions can be written in the following manner:

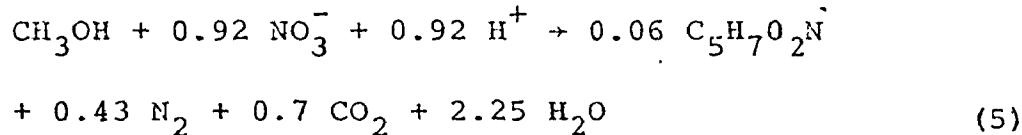
Step 1: NO_2^- formation:



Step 2: N_2 gas formation:



The above equations represent only the nitrate dissimilation part of denitrification. McCarty (1973) presented the following resultant equation for both nitrate dissimilation and bacterial production:



Therefore theoretically it is natural to expect, 0.95 g' CH_3OH -C (methanol carbon) utilization for denitrification of each g' of NO_3^- -N, with the production of 0.53 g of micro-organism.

McCarty, Bech, and St. Amant (1969) gave the following formulas for calculating actual micro-organism production and methanol requirement based upon their denitrification studies.

For biomass production:

$$C_b = 0.53 N_o + 0.32 N_1 + 0.19 DO \quad (6)$$

where: C_b is the biomass produced in mg/l,
 N_0 is the initial $\text{NO}_3\text{-N}$ concentration in mg/l,
 N_1 is the initial $\text{NO}_2\text{-N}$ concentration in mg/l, and
 D_o is the dissolved oxygen concentration of influents in mg/l.

Dissolved oxygen is another consumer of $\text{CH}_3\text{-OH}$, as in the presence of oxygen and a carbon source growth of aerobic bacteria will occur.

For methanol consumption:

$$C_m = 0.53 N_0 + 0.32 N_1 + 0.19 D_o$$

where: C_m is the required methanol (mg/l)

The experimental results obtained by Sutton, Murphy and Dawson (1974) confirmed the estimates from the above formulas.

3.1.2 Kinetics of Denitrification

Various authors have indicated the denitrification rate to be zero order with respect to $\text{NO}_3\text{-N}$ concentration for suspended growth systems if $\text{CH}_3\text{-OH}$ is supplied in sufficient quantities and the $\text{NO}_3\text{-N}$ concentration is above 1.0 mg/l.

These authors include Moore and Schroeder (1970), and Murphy and Sutton (1974).

In addition, Sutton, Murphy and Dawson (1975) and Requa and Schroeder (1973) indicated the denitrification rate to be zero order with respect to NO_3^- -N concentration for supported growth systems. Harremoes (1975) reported that the denitrification rate was half order with respect to NO_3^- -N, owing to diffusion limitations in supported growth systems.

3.1.3 Temperature Dependency

The temperature dependency of denitrification processes is one of the most thoroughly studied aspect of denitrification since it is essential factor for process design and volumetric determination.

Table 1 gives the type of processes and the corresponding temperature dependency relationship presented by various authors. The authors listed in Table 1 presented the temperature dependency of denitrification as an empirical expression

$$K = A_0 e^{-E/RT} \quad (8)$$

where: K is the zero order reaction rate,
 A_0 is the constant
 E is the activation energy,
 R is the gas reaction constant, and
 T is the temperature in $^{\circ}\text{K}$.

TABLE 1

TEMPERATURE DEPENDENCY OF VARIOUS DENITRIFICATION UNITS

Type of Systems	Frequency Factor A_0	Activation Energy E	Author
BST SRT = 3 days	6.78×10^{14}	20,400	Sutton and Jank (1976)
BST SRT = 6 days	4.78×10^9	13,900	"
CSTR SRT = 3 days	8.76×10^8	12,900	"
CSTR SRT = 6 days	1.08×10^{11}	15,900	"
CSTR / SRT = 9 days	8.76×10^{10}	15,650	"
BST + CSTR SRT = 3 days	5.42×10^{10}	15,300	"
BST + CSTR SRT = 6 days	1.11×10^{11}	15,900	"
Pall Rings	2.89×10^6	7,025	"
Intalox Saddle /	2.97×10^9	11,090	"
Batch Laboratory	3.195×10^{11}	+16,800	Dawson and Murphy (1972)

Pretorius and Davies (1974) define 3 different temperature dependency regions for denitrification rates achieved by RBCs namely:

$$Q_{10} = 13.06 \text{ for } T < 10^{\circ}\text{C}$$

$$Q_{10} = 1.38 \text{ for } 30^{\circ}\text{C} > T > 10^{\circ}\text{C}$$

$$Q_{10} = -2.65 \text{ for } T > 20^{\circ}\text{C}$$

where: Q_{10} is the rate increase for 10°C temperature increase, and

T is the temperature in $^{\circ}\text{C}$.

One of the main purposes of this study is to determine the temperature dependency relationships for denitrification by the RBC under different hydraulic loadings.

4. EQUIPMENT AND PROCEDURES

4.1 Pilot Plant and Reactor Descriptions

The RBC unit used in this study comprises part of a pilot plant facility located at the Wastewater Technology Centre at CCIW, Burlington, Ontario. The pilot plant was installed expressly for nitrification and denitrification research. Figure 1 indicates a simplified flow diagram of the pilot plant. The screened raw sewage is obtained continuously from Burlington Skyway Sewage Treatment Plant. The nitrified effluent passes through a temperature control unit. A NO_3^- -N spike is applied when necessary. The nitrified effluent then enters the denitrification units.

The RBC used for denitrification in this study was an Autotrol Bio-Surf, Serial No. 7415. The unit was originally manufactured as a 4 stage reactor (Figure 2). There are a total of 44 discs (11 in each stage) with a total surface area of 306 ft^2 (28.4 m^2). Rotation was at 13 rpm. The diameter of discs was 47.3 cm (18-5/8 inch). The total volume of the total unit was 408 l for 4 stage operation. This measurement was done when biological growth existed on the discs. The volume for single stage operation was measured as 100 l.

FIGURE NO. 1 SIMPLIFIED FLOW DIAGRAM OF NITRIFICATION / DENITRIFICATION PILOT PLANT

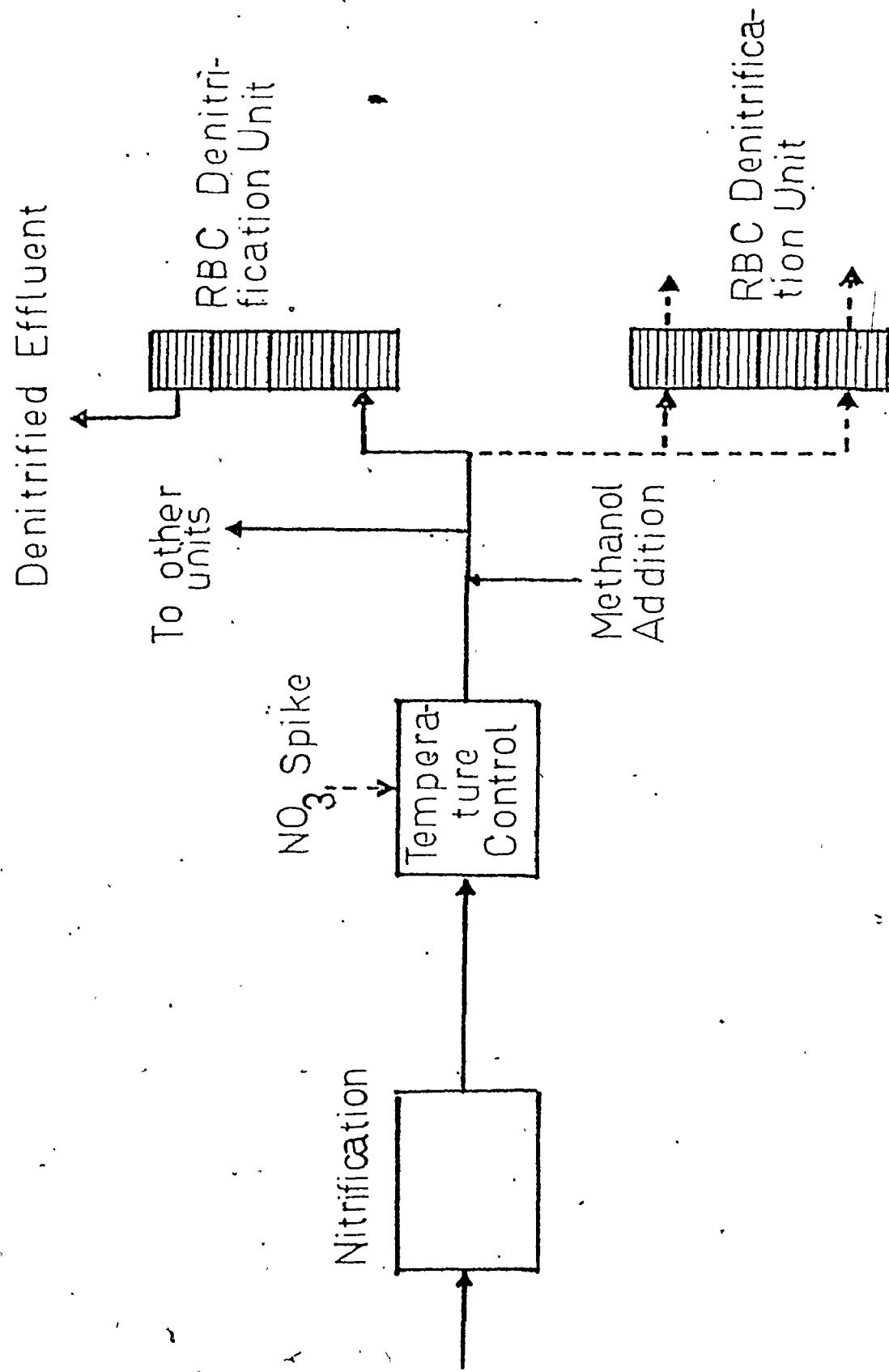
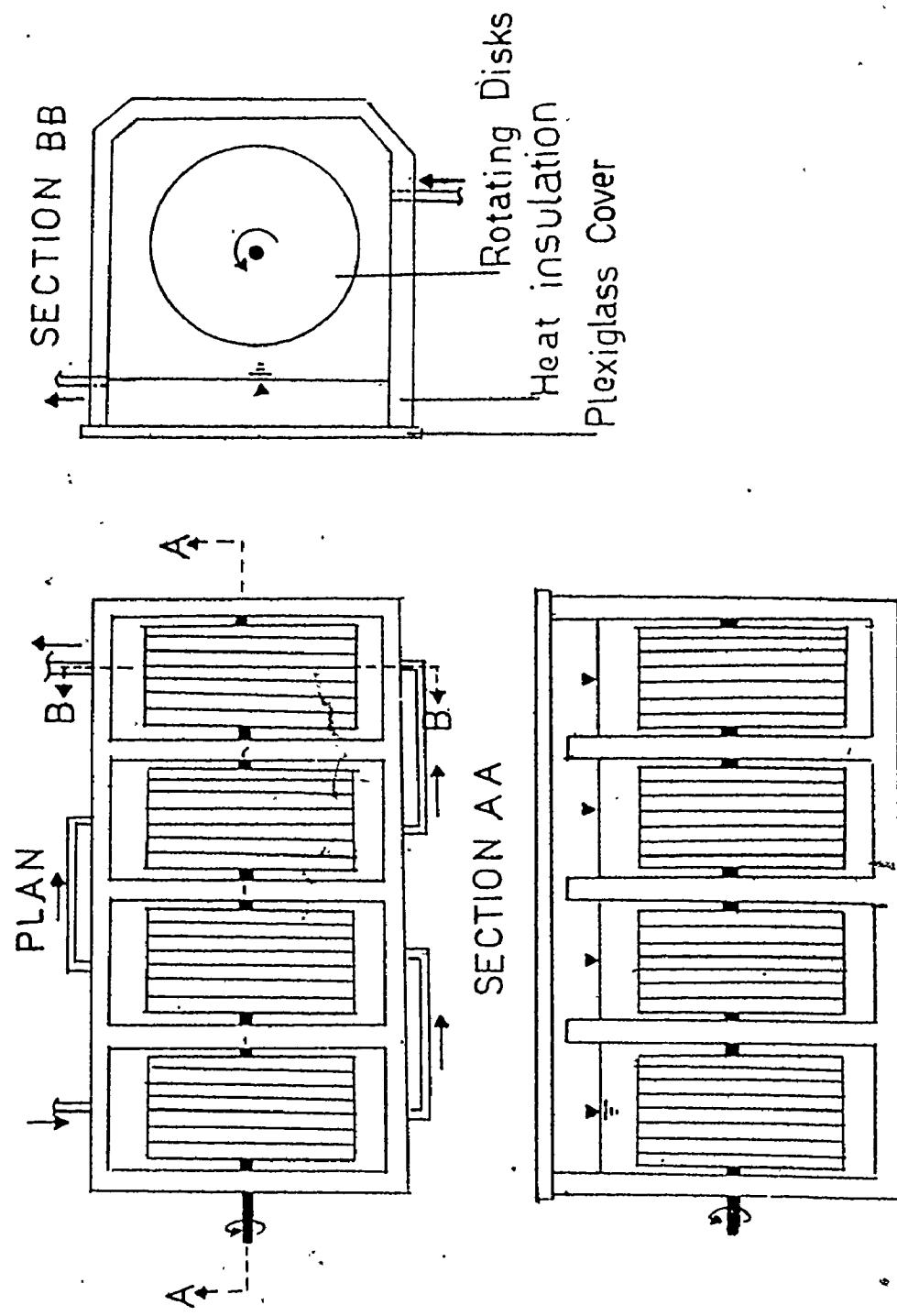


FIGURE NO. 2 COMPLETELY SUBMERGED RBG



Variable speed positive displacement pumps delivered nitrified secondary effluent to the submerged RBC unit.

4.2 Start Up

The RBC had been operated two months before the initiation of the experimental program. No attempt was made to evaluate the minimum start up time required. The original unit provided by Autotrol was covered with one half inch (1.3 cm) styrofoam on the sides to reduce the heat transfer and to obtain a relatively constant temperature in all stages of RBC.

4.3 Experimental Program and Work

Since the purpose was mainly to investigate the basic design criteria of RBC as a denitrification unit, the effects of the following design variables or controlled factors were investigated - temperature, hydraulic loading and mode of operation (single stage or 4 stage in series).

It was planned to run the RBC under temperature levels of 5°C, 10°C, 15°C, 20°C, and 25°C (two runs for each temperature level) for 4 stage in series operation. This original program was altered later, since unreasonable and impractical amounts of NO₃-N spike were needed to have measurable quantities of NO₃-N in the effluent to determine removal rates. Only single runs were employed with the 4 stage in series operation.

Similarly, two runs were planned for temperature levels of 5°, 10°, 15°, 20°, and 25°C, at hydraulic loadings of 2.5 l/min and 4.0 l/min of single

stage operation.

As it was expected that the dye response would be a function of the biological growth which existed on the discs, dye studies at 4 different temperature levels with biological growths specific to that temperature level were carried out. A separate dye study was made after all growth on the discs was stripped off.

Detailed experimental designs for single stage operation and 4 stage series operation are included in the appendix. The plant was operated at least 5 days before taking samples for the determination of denitrification rates at any specified operating condition.

4.4 Feed Characteristics

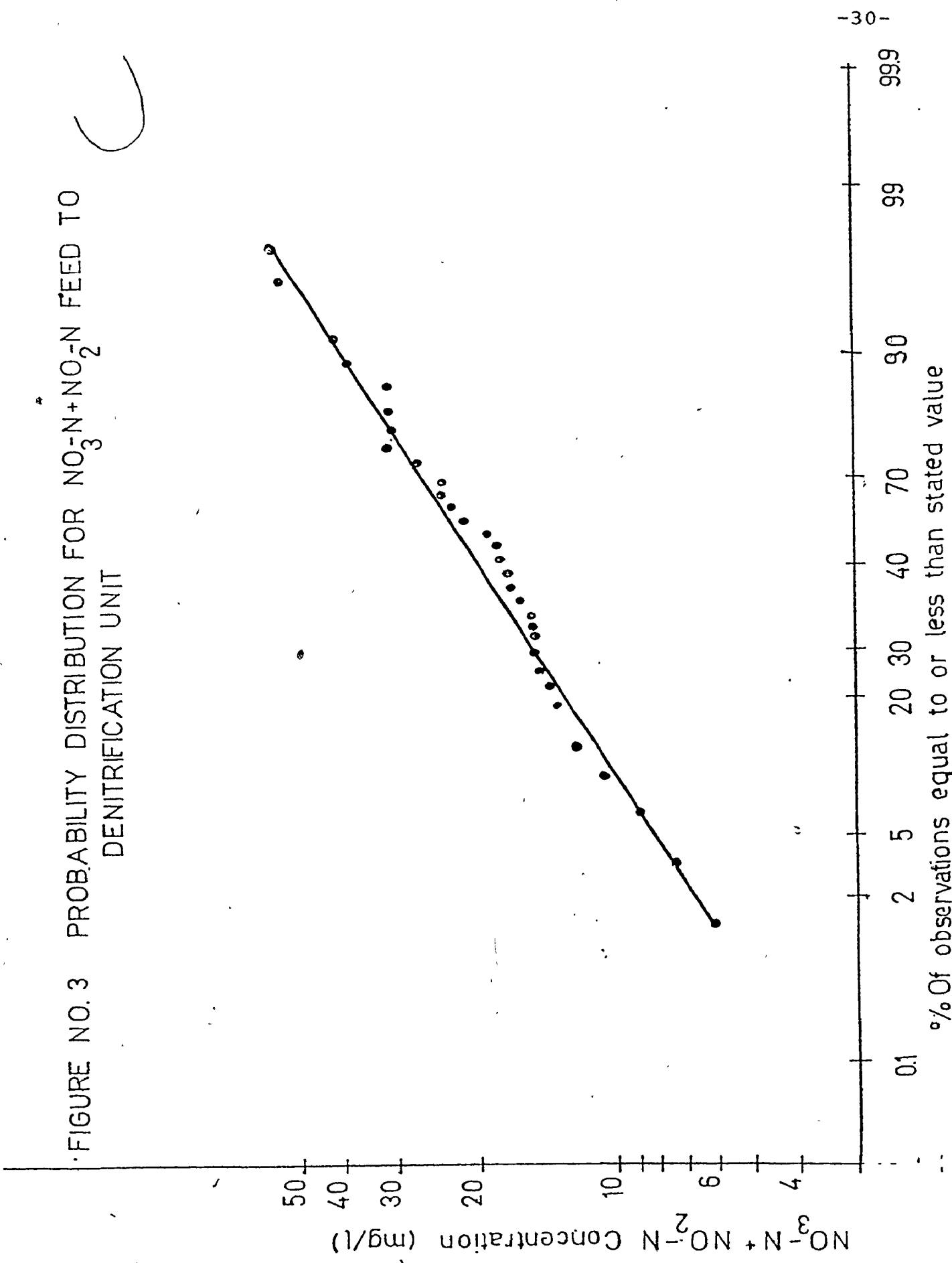
The NO_3^- -N concentration of the feed fluctuated and was dependent upon the efficiency of the nitrification step, the strength of raw sewage and the spike addition if any.

NO_3^- -N spikes in the form of KNO_3 were necessary especially at high temperature levels to obtain NO_3^- -N residual in the effluent in order to measure the denitrification rate.

Carbon was added in the form of methanol. The C:N ratio in the feed was kept about 1.5:1.0 so that microbiological activity was not limited by carbon limitations.

Figure 3 shows the probability distributions of $\text{NO}_3^- + \text{NO}_2^-$ -N concentration in the feed throughout the study. The resulting carbon nitrogen ratio from the addition of

FIGURE NO. 3 PROBABILITY DISTRIBUTION FOR NO_3^- -N + NO_2^- -N FEED TO DENITRIFICATION UNIT



methanol is given in Figure 4.

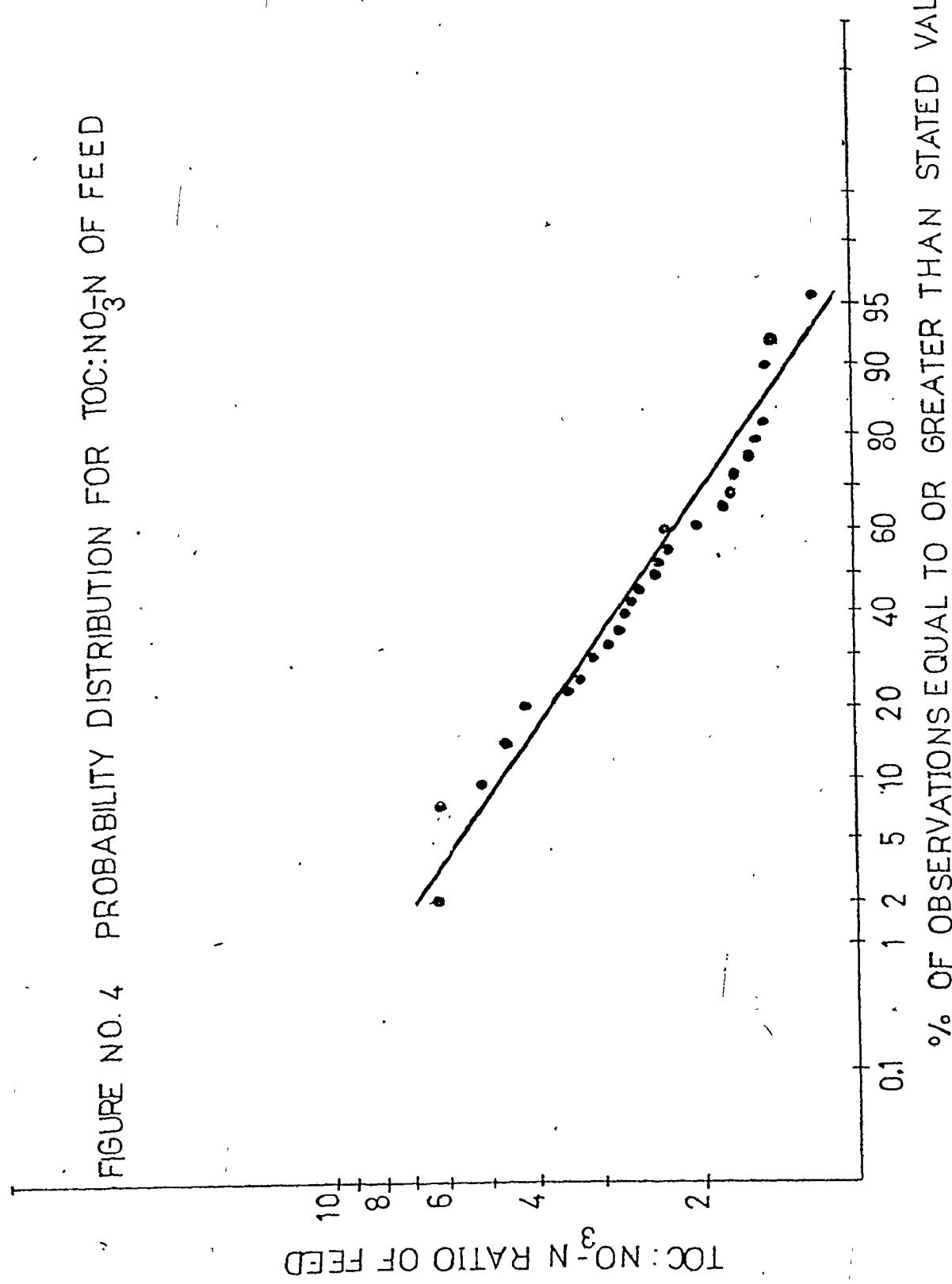
4.5 RBC Operation and Sampling

The operation of the RBC was controlled closely. The operation of the unit was easy and generally trouble free. The following operating problems were noted.

- a) Continuous running of the disc shaft was essential as a considerable amount of bio-mass was lost after each power failure which effected the performance-capacity of the unit.
- b) The biomass activity stopped when raw water which contained chlorine was used as make up water due to possible chlorine toxicity. After this fact was observed, dechlorinated water was used as make up water.
- c) Occasional bubbling within the reactor due to air entrainment was observed. This caused some of the biomass on the rotating discs to be stripped off.
- d) With the hydraulic loading employed, and with the reduced microbial activity at low temperature, a measurable concentration of DO was monitored in the first compartment of four stage in series operation. This reduced the rate of denitrification in this compartment.

Sampling was done after 5 days of continuous operation for each run. The total duration of sampling for each run was 2 hours, with one sample for every half an hour.

FIGURE NO. 4 PROBABILITY DISTRIBUTION FOR TOC:NO₃-N OF FEED



The samples were analyzed directly or kept refrigerated and analyzed later.

The dye samples were collected with 1 min and 2 min intervals for hydraulic loadings of 4.0 l/min and 2.5 l/min respectively, after pulse dye injection.

5. RESULTS AND DISCUSSION

5.1 Hydraulic Characterization

5.1.1 Simulation of Single Stage Operation

Three separate dye studies were conducted to simulate the flow through a single stage RBC. Dye studies were made on a single stage with the unit operating at 15°C and hydraulic loadings of 4 l/min, 2.5 l/min with biomass specific to the operating conditions. An additional dye study was performed at 15°C with all the biomass removed from the discs.

Rhodamine WT dye was injected as a pulse input and the effluent response was measured. The measured dye concentration for the above operating conditions are listed in Appendix 2.

The dye concentrations obtained from all tracer response studies (Figures 5,6, and 7) indicate that the responses for the single stage operation is intermediate to that of a single CSTR with 40 l volume, i.e., the volume not occupied by rotating discs and that of a single CSTR with 100 l volume, i.e., the total volume of the single stage.

A computer program (Appendix 2) calculated the dye recovery, mean dye residence time, residence time distribution (E), dimensionless residence time distribution (Et), and F curves.

FIGURE NO. 5 DYE CONCENTRATION VERSUS TIME
Single stage operation
Hydraulic loading= 2.5 l/min.

- 1 CURVE for CSTR of 100 l. Volume
- 2 " " " 40 l. Volume
- 3 Actual dye measurements

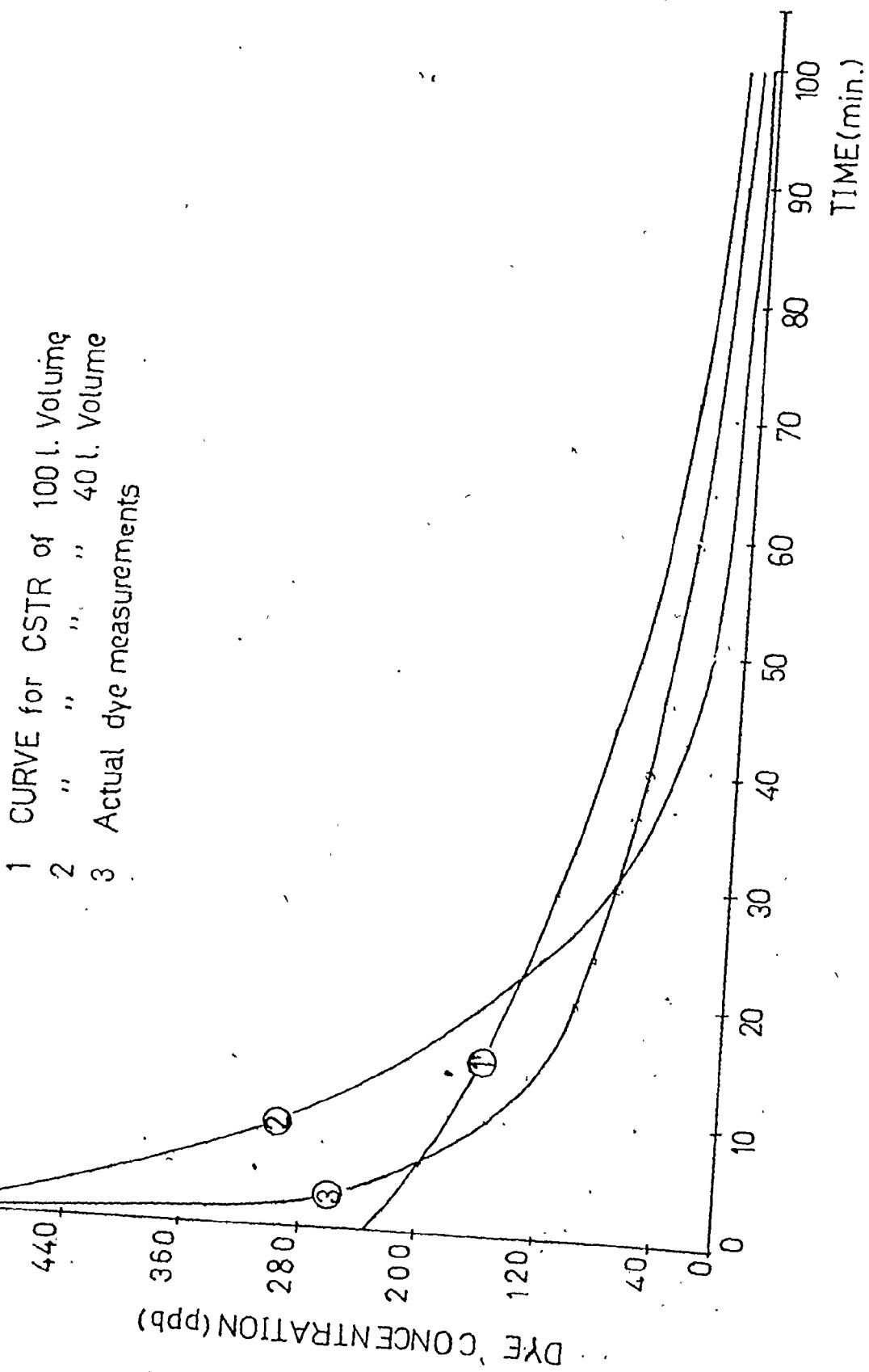


FIGURE NO. 6 DYE CONCENTRATION VERSUS TIME
Single stage operation
Hydraulic loading=4.0 l./min.

- 1 Curve for CSTR of 100 l Volume
- 2 " " 40 l Volume
- 3 Actual dye measurements

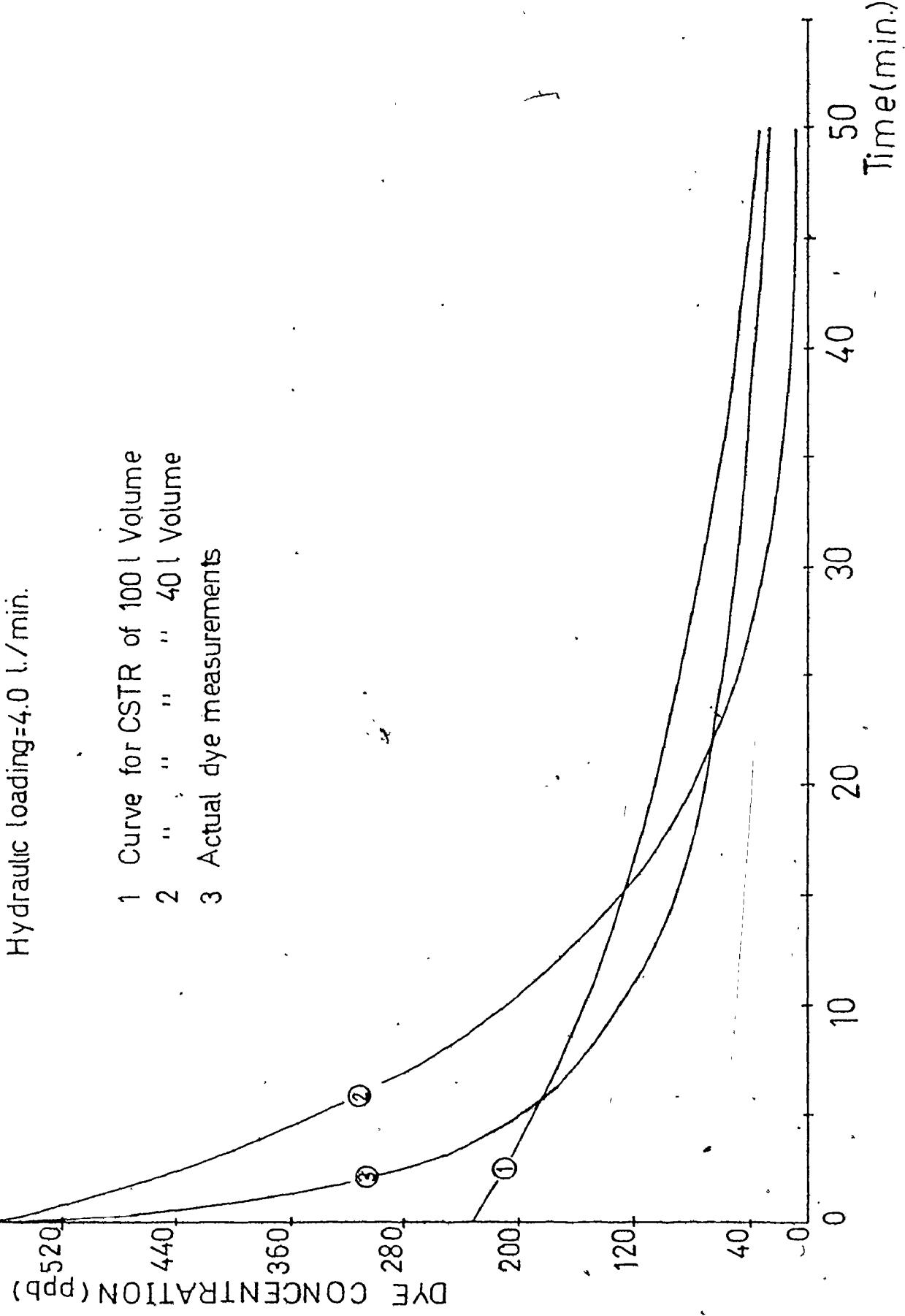


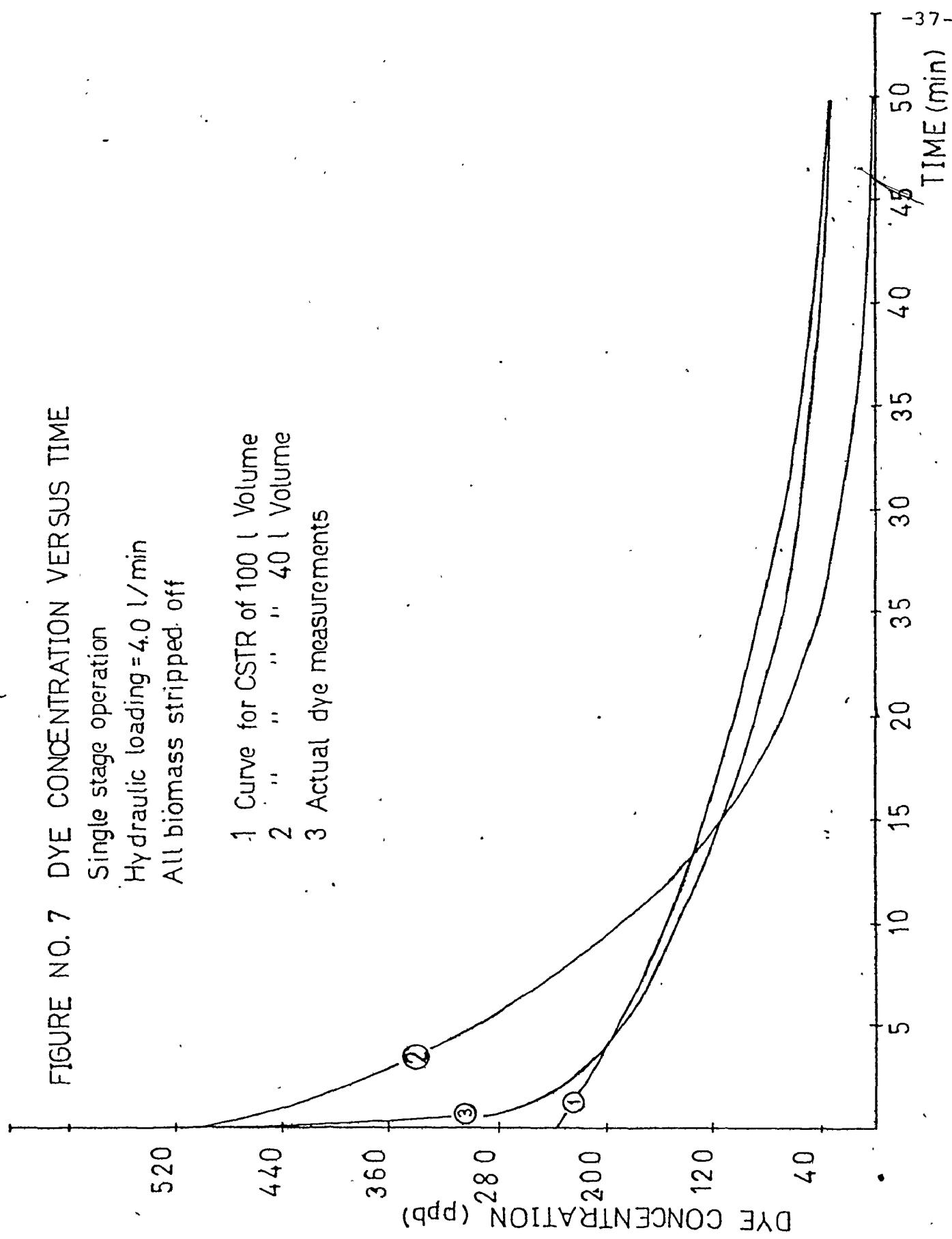
FIGURE NO. 7 DYE CONCENTRATION VERSUS TIME

Single stage operation

Hydraulic loading = 4.0 l/min

All biomass stripped off

- 1 Curve for CSTR of 100 l Volume
- 2 " " " 40 l Volume
- 3 Actual dye measurements



The effect of the accumulated biomass in modifying the tracer response curve is illustrated by dimensionless residence time distribution curves (E_t) (Figure 8) for the single stage tracer studies.

Both RTD (residence time distribution) and concentration curves implied that the actual flow pattern differed from that of a single CSTR. A model employing two CSTRS with interchange flow was proposed (Fig. 9). The volume not occupied by disc drum was considered to be 100% active and basically a CSTR of 40 l volume, and the liquid volume within the disc drum was considered to be a CSTR of 60 l volume with no dead space. An interchange flow between the two CSTR's (v_B) was postulated. No shortcirculating was considered to exist.

The method uses a material balance of the dye to determine the flow rate between 2 CSTR's.

At any time t , after a pulse dye injection:

$$CA_t VA = CA_0 VA - \int_{t=0}^t CA v dt - \int_{t=0}^t CA v_B dt + \int_{t=0}^t CB v_B dt \quad (9)$$

and

$$CB_t VB = \int_{t=0}^t CA v_B dt - \int_{t=0}^t CB v_B dt \quad (10)$$

FIGURE NO 8 DIMENSIONLESS RESIDENCE TIME

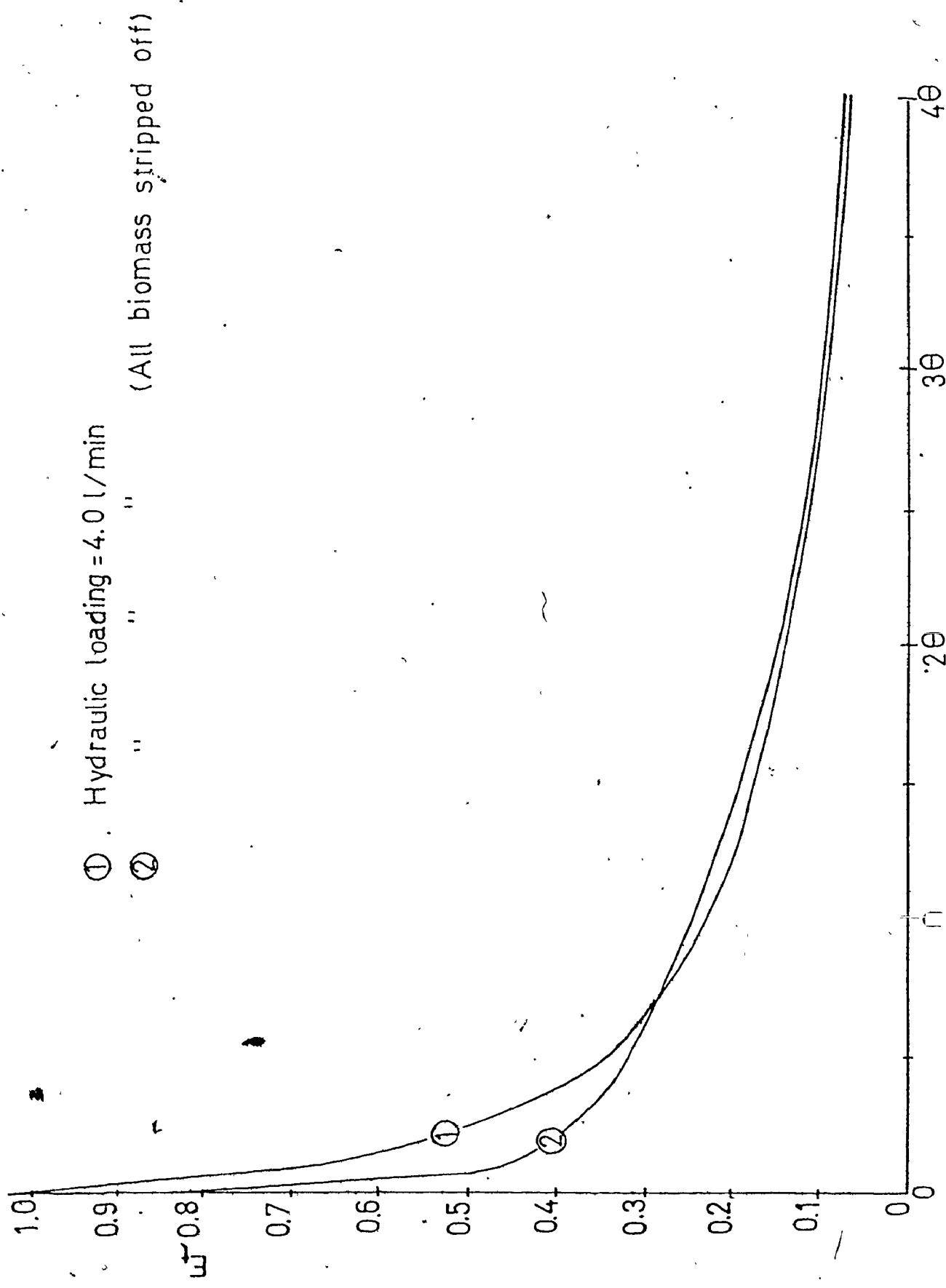
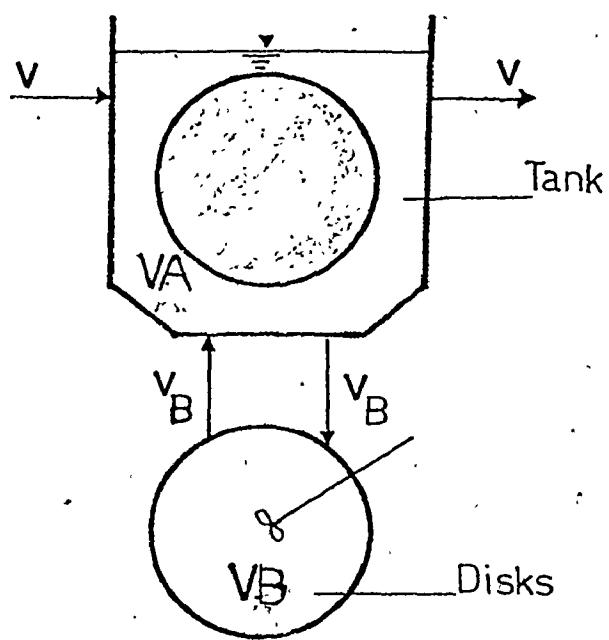


FIGURE NO. 9 PROPOSED FLOW PATTERN
FOR SINGLE STAGE OPE-
TION



where:

CA_0 = concentration of dye at time $t = 0$ in outer CSTR, ppb

CA_t = concentration of dye at time t after pulse injection in outer CSTR, in ppb

CA = concentration of dye in outer CSTR, ppb

CB_t = concentration of dye at time t after pulse injection in inner CSTR, in ppb

CB = concentration of dye in inner CSTR, ppb

v = hydraulic flow rate, l/min

v_B = the flow exchange rate between inner and outer CSTR, l/min

VA = volume of outer CSTR, l

VB = volume of inner CSTR, l

From equations 9 and 10:

$$CA_t VA = CA_0 VA - \int_{t=0}^t CA v dt - CB_t VB \quad (11)$$

and

$$CB_t VB = CA_0 VA - \int_{t=0}^t CA v dt - CA_t VA \quad (12)$$

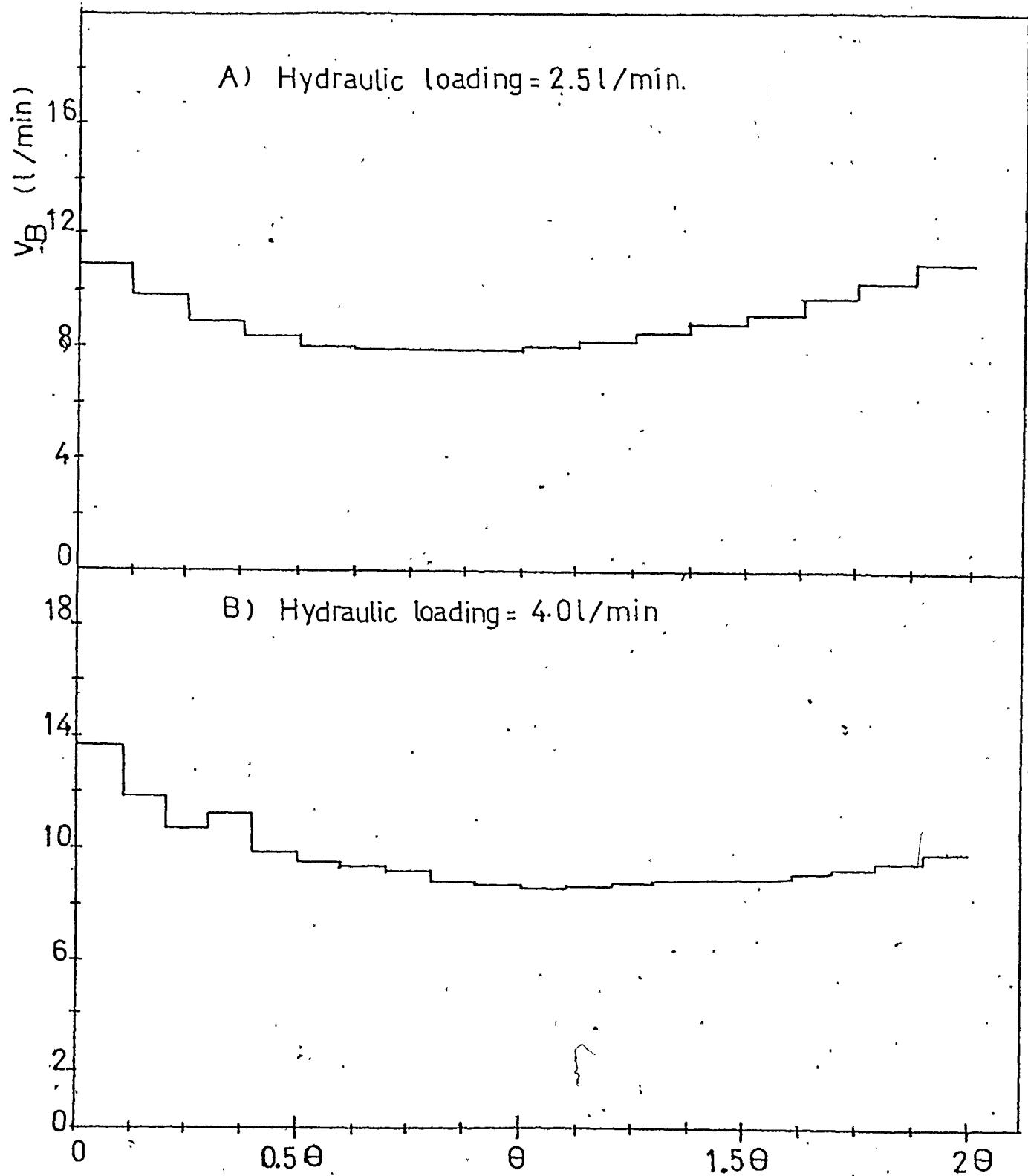
The only unknown of equation (12) is CB_t , since CA and CAt are known as a result of the dye study. VA is known due to the geometry of unit, and v , the hydraulic load is selected. Therefore, CB_t , the dye concentration in the disc drum, can be calculated using equation (12).

Once CB_t , at any time, is calculated, the determination of flow exchange rate between inner and outer CSTR's is possible by using the modified form of equation (10):

$$v_B = \frac{CB_t V_B}{\int_{t=0}^t CA dt - \int_{t=0}^t CB dt} \quad (13)$$

This method was applied to the data obtained from the pulse dye study on the single stage mode using the computer program presented in Appendix 2. The results are presented in Figure 10. Figure 10 indicates that there is relatively a constant flow exchange rate between the inner and outer CSTR's for 2 detention times of the outer CSTR at a hydraulic loading of 2.5 l/min. Figure 10 indicates that the flow exchange rate between inner and outer CSTR's is relatively constant except in the early period at a hydraulic loading of 4.0 l/min. If the proposed model of 2 CSTR's with interchange flow fitted perfectly, the v_B values calculated would be constant over the entire duration. Figure 10 indicates that there are only slight deviations from the proposed model.

FIGURE NO. 10 INTERCHANCE FLOW RATES(CALCULATED USING MATERIAl BALANCE ON PULSE DYE INJECTION)



Figures 11 and 12 give the dye concentrations of both inner and outer CSTR's. CB values are calculated by using the above numerical method and flow exchange rate as presented in Figure 10.

While this method is a powerful tool to determine if there is a relatively constant interflow rate between the inner and outer CSTR, its sensitivity depends upon the consistency of pumping rate and accuracy of concentration measurement. It is natural to expect high propagation of errors after a prolonged integration period.

The validity and degree of deviation of the proposed model can be checked by comparing the actual data to calculated curves using the formula given by Dedrick and Bischoff (1968). Their formula calculates the dye concentration in the outer CSTR for pulse input for the postulated flow model.

Dedrich and Bischoff's Formula for the concentration in the outer CSTR of a compartment model is:

$$\frac{CA_t}{CA_0} = \frac{(r_2 + \frac{v_B}{V_B})e^{r_2 t} - (r_1 + \frac{v_B}{V_B})e^{r_1 t}}{2R} \quad (14)$$

where:

$$r_1, r_2 = -\frac{1}{2} \left[\frac{v_A}{V_A} + \frac{v_B}{V_A} \left(1 + \frac{V_A}{V_B} \right) \right] \mp R \quad (15)$$

FIGURE NO 11 DYE CONCENTRATION OF INNER AND OUTER CSTR'S

Hydraulic loading=2.5 l/min

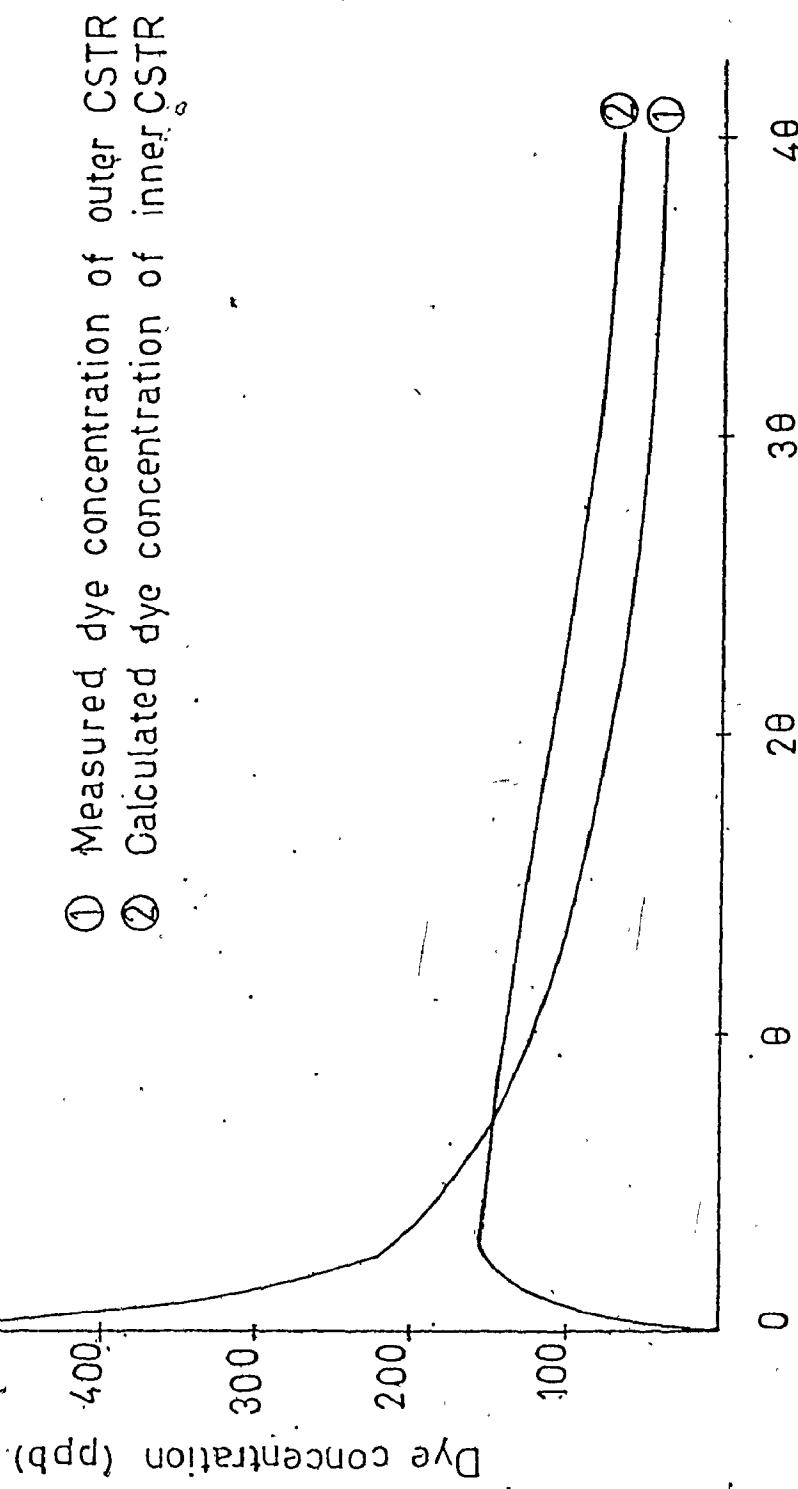
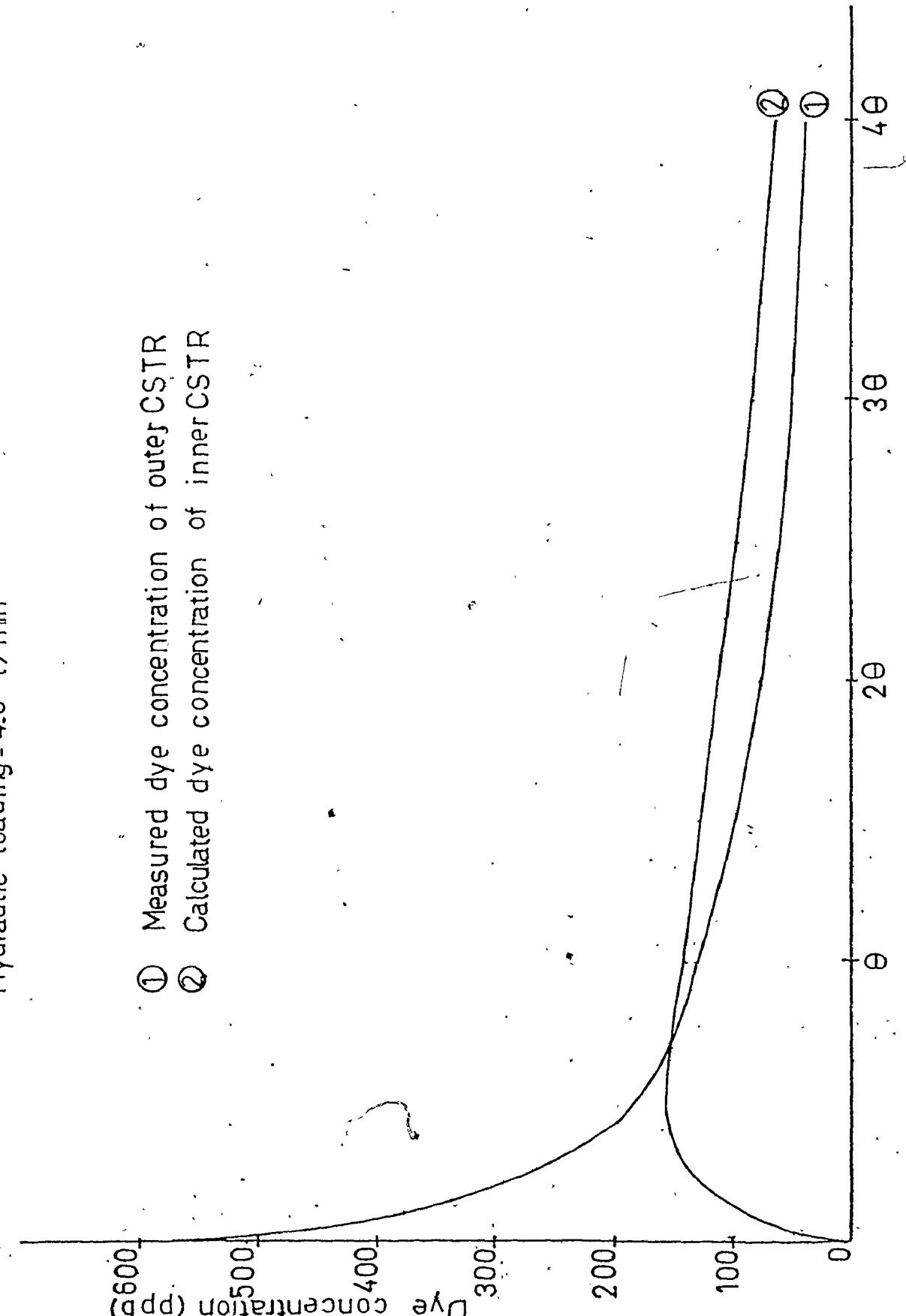


FIGURE NO 12 DYE CONCENTRATION OF INNER AND OUTER CSTR'S

Hydraulic loading = 4.0 l/min

- ① Measured dye concentration of outer CSTR
- ② Calculated dye concentration of inner CSTR

Dye concentration (ppm)



and

$$R = \sqrt{\frac{1}{4} \left[\frac{V_B}{V_A} + \frac{V_B}{V_A} \left(1 + \frac{V_A}{V_B} \right) \right]^2 - \frac{V_B}{V_A V_B}} \quad (16)$$

Figures 13 and 14 give the comparison of the curves for the predicted responses using Equations 14, 15, and 16 (Appendix 2, Program DEDBIS), and the measured dye concentrations, for hydraulic loadings of 2.5 l/min and 4.0 l/min.

5.1.2 Simulation of 4 Stage Operation

Pulse dye injection was applied at a flow different operating conditions, namely, rate of 4 l/min and at temperatures of 10, 15, 20, and 25°C. For all the above operating conditions, there was biological growth on the discs.

A corresponding dye injection at 4 l/min flow rate and temperature of 15°C was made and all the biomass stripped from the discs.

The tracer concentrations are given in Appendix 2. Also E_t versus θ curves were calculated for different operating conditions, using the computer program of

FIGURE NO 13

COMPARISON OF ACTUAL DYE MEASUREMENTS
WITH THE FITTED CURVE
(NONLINEAR TO DEDRICK AND BISCHOFFS FORMULA)
FITTING TECHNIQUE IS APPLIED

Fitted curve ($y_B = 10.3l/min, v = 2.6l/min.$)

Actual measurements

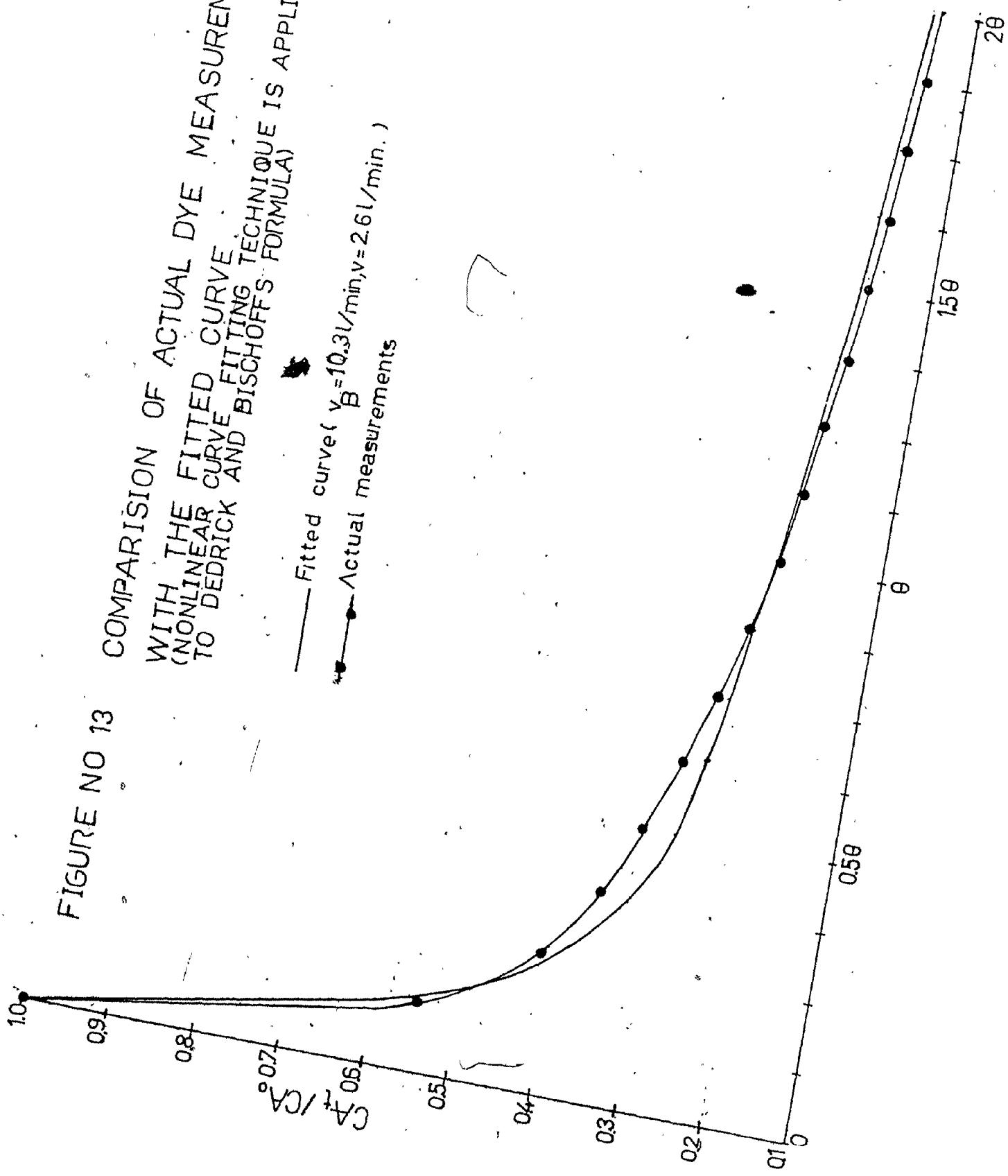
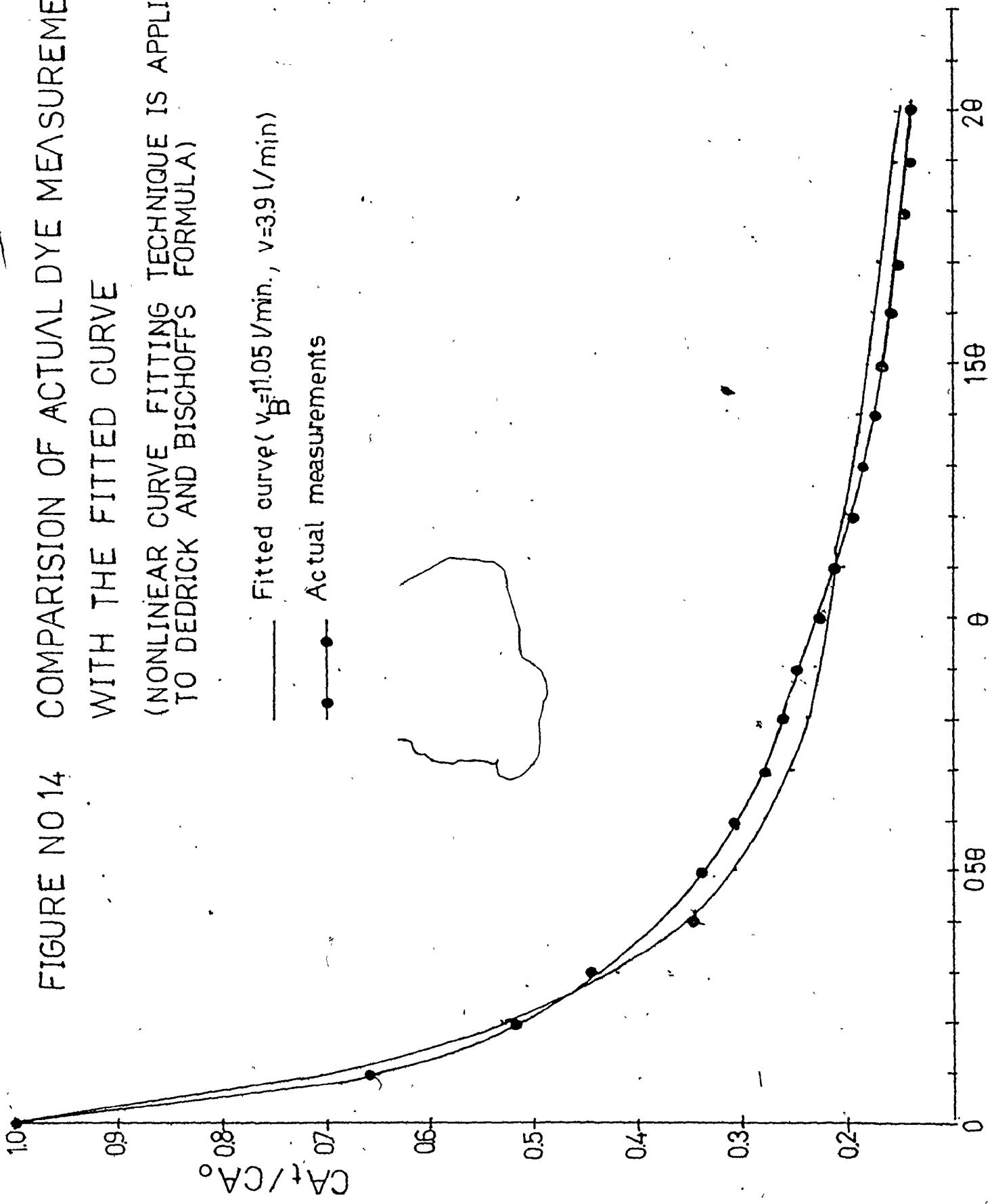


FIGURE NO 14

COMPARISION OF ACTUAL DYE MEASUREMENTS
WITH THE FITTED CURVE

(NONLINEAR CURVE FITTING TECHNIQUE IS APPLIED
TO DEDRICK AND BISCHOFF'S FORMULA)

— Fitted curve ($V_B = 11.05 \text{ V/min.}$, $v = 3.9 \text{ l/min.}$)
● Actual measurements



Appendix 2. These curves are plotted on Figure 15 (see Table 2).

Figure 15 shows that the tracer response curve is a function of the biomass that exists at a given operating temperature. This conclusion is supported by comparison of Figures 4, 5, and 6.

5.2 EVALUATION OF A TOTALLY SUBMERGED RBC FOR DENITRIFICATION

It has been demonstrated that each compartment of an RBC can be simulated as two CSTR's with interchange flow. Geometric properties of reactors dictates that the outer surface area (1.40 m^2) of a total surface area of discs (7.10 m^2) would be available to the outer CSTR, while the remainder 5.70 m^2 is available to the inner CSTR. The reactor walls which are equivalent to 0.5 m^2 should also be included as an active area as they attached heavy biomass throughout the study. Therefore, the effective surface area of the outer CSTR would be 1.90 m^2 .

Using a model similar to Figure 9, a material balance for the substrate in the outer CSTR gives:

$$vS_o - K A_A S_A^n - v_B S_A + v_B S_B - v S_A = 0 \quad (17)$$

where: S_o = the substrate concentration in influent, mg/l

v = hydraulic loading, l/min

K = substrate removal rate, $\text{mg/m}^2 \cdot \text{min}$

A_A = available area of outer CSTR, m^2

FIGURE NO. 15 DIMENSIONLESS RESIDENCE TIME DISTRIBUTION FOR FOUR STAGE IN SERIES

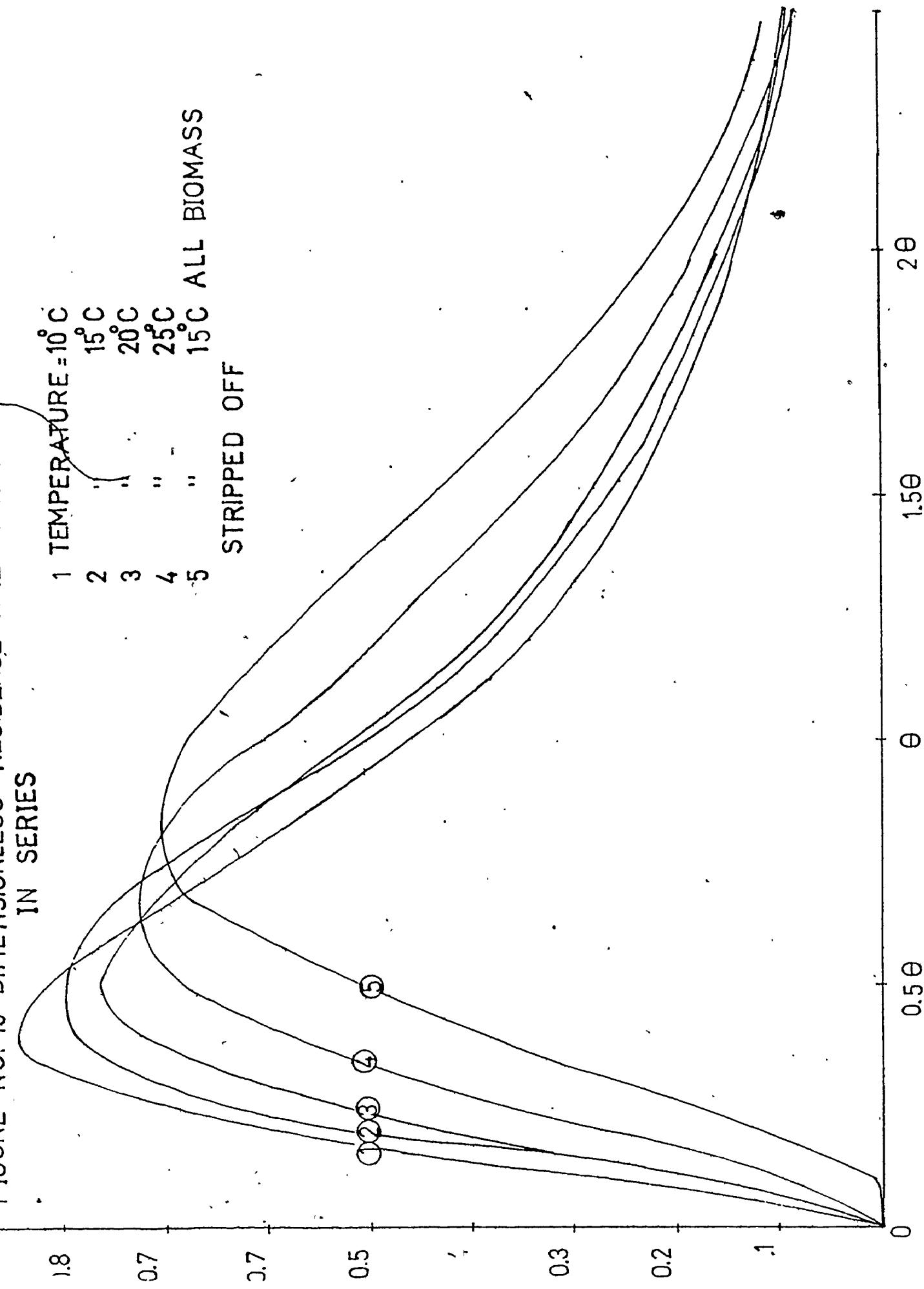


TABLE 2

THE OBSERVED BIOMASS UNDER DIFFERENT OPERATING TEMPERATURE
FOR 4 STAGE IN SERIES MODEL OF OPERATION

Temperature under which dye study performed	Stages containing heavy biomass
25°C	Only first stage
20°C	First two stages
15°C	First three stages
10°C	All stages
15°C (stripped)	None

v_B = flow exchange rates between inner and outer CSTR, l/min

S_A = substrate concentration in outer CSTR, mg/l

S_B = substrate concentration in inner CSTR, mg/l, and

n = order of denitrification rate, dimensionless.

A material balance on the inner CSTR gives:

$$v_B S_A - K A_B S_B^n - v_B S_B = 0 \quad (18)$$

where:

A_B = the area available in inner CSTR,
 m^2

For a zero order reaction rate, $n=0$, was inserted and S_B and K values were calculated using equations 17, and 18. Table 3 gives the calculated K , S_A , and S_B values for different operating temperatures and hydraulic loadings, using the data tabulated in Appendix 1. These calculations are summarized in Appendix 3.

Similarly for the first order reaction rate, $n=1$ was inserted, and S_A , S_B , and K values were calculated using equations 17 and 18 (Table 4 and Appendix 3).

The empirical Arrhenius temperature dependency describing the denitrification for the totally submerged RBC may be calculated using the K values from Tables 3 and 4.

THE CALCULATED DENITRIFICATION RATES AND SUBSTRATE CONCENTRATIONS OF INNER AND OUTER CSTR'S ASSUMING ZERO ORDER KINETIC MODEL (K's are in mg/m²-hr)

Hydraulic loading l/min	K at Temperature (°C)	S _O mg/l	S _A mg/l	S _B mg/l	K at Temperature (°C)	S _O mg/l	S _A mg/l	S _B mg/l
2.5	K ₅ °C=66.1	13.6	10.3	9.7	K ₂₀ °C=455.4	30.0	6.9	3.3
	K ₅ °C=159.6	25.2	17.1	15.8	K ₂₀ °C=315.6	29.0	13.0	10.5
	K ₁₀ °C=92.4	17.0	12.3	11.6	K ₂₅ °C=528.8	32.0	6.2	2.2
	K ₁₀ °C=100.2	18.0	12.9	12.1	K ₂₅ °C=375.6	22.9	3.9	0.9
	K ₁₅ °C=177.6	13.8	4.7	3.5	K ₂₅ °C=1007.4	29.0	2.8	0.0
	K ₁₅ °C=169.2	13.3	4.7	3.4				
4.0	K ₅ °C=30.0	13.6	12.7	12.5	K ₁₀ °C=240.0	17.0	9.4	7.7
	K ₅ °C=82.1	26.2	23.6	23.1	K ₁₅ °C=145.3	19.0	14.4	13.4
	K ₅ °C=123.7	15.4	11.5	10.6	K ₂₀ °C=631.6	30.0	10.0	5.7
	K ₅ °C=53.7	17.0	15.3	14.9	K ₂₅ °C=705.9	30.0	7.7	2.9
	K ₅ °C=116.8	14.1	10.4	9.6	K ₂₅ °C=735.8	40.0	16.7	11.7
	K ₁₀ °C=66.3	17.0	14.9	14.5	K ₂₅ °C=378.9	54.0	42.0	39.4
	K ₁₀ °C=94.7	18.0	15.0	14.4				

TABLE 6

THE CALCULATED DENITRIFICATION RATES AND SUBSTRATE CONCENTRATIONS OF INNER AND OUTER CSTR's ASSUMING FIRST ORDER KINETIC MODEL (K' 's are in $1/m^2 \text{-min}$)

Hydraulic Loading 1/min	K at Temperature (°C)	S_O (mg/l)	S_A (mg/l)	S_B (mg/l)	K at Temperature (°C)	S_O (mg/l)	S_A (mg/l)	S_B (mg/l)
2.5	0.11	13.6	10.3	9.7	1.68	30.1	6.9	3.8
	0.16	25.2	17.1	15.8	0.47	29.0	13.0	10.6
	0.12	17.0	12.3	11.6	2.23	32.0	6.2	3.0
	0.13	18.0	12.9	12.1	2.89	22.9	3.8	1.6
	0.79	13.8	4.7	3.4	7.53	29.0	2.8	0.6
	0.76	13.3	4.7	3.4				
4.0	0.04	13.6	12.6	12.4	0.46	17.0	9.4	7.8
	0.05	26.2	23.6	23.0	0.17	19.0	14.4	13.4
	0.19	15.4	11.4	10.6	1.46	30.0	10.0	6.2
	0.07	17.0	14.9	14.4	2.47	30.0	7.6	3.8
	0.19	14.1	10.4	9.6	0.43	40.0	23.3	19.8
	0.06	17.0	15.3	14.9	0.15	54.0	42.0	39.4
	0.10	18.0	15.0	14.3				

The Arrhenius temperature dependency was calculated by using reparameterized equation, Sutton (1971):

$$K = K^* (e^{-E/RT_0} A_0) \quad (19)$$

where:

$$K^* = e^{-\frac{E}{R}(\frac{1}{T} - \frac{1}{T_0})} \quad (20)$$

A computer program (Appendix 2), calculates K^* and E values, once K values and corresponding temperatures are inserted. This program utilizes a least square fitting technique using linear regression.

Another program (Appendix 2), calculates the 95% confidence limits for predicting of actual reaction rate values, utilizing the output of previous program. Computational examples are illustrated in Appendix 4 and the results are given in Table 5.

As there did not appear to be much difference in the models for 2.5 l/min and 4.0 l/min hydraulic loadings, assuming zero order reaction rates, the results for both flow rates were combined and analyzed together. The ANOVA results (Table 5) indicated no lack of fit for the combined data at the 95% significance level. The fitted curve for the combined data is presented in Figure 20. Similar data for a "first order" rate is also presented in Table 5. Using a first order model, the analysis of variance indicated the two flow rates could not be pooled as with the zero order model.

TABLE 5
RESULTS OF LINEAR REGRESSION FOR ARRHENIUS TEMPERATURE DEPENDENCY

Operating Condition	-E/R	Activation Energy cal/mole	$\frac{-E}{RT_0}$	A_0	ANOVA RESULTS		
					lnA ₀	ϵ	Mean Square Lack of Fit Mean Square Pure Error
ZERO ORDER							
Hydraulic loading = 2.5 l/min	-7800	15400	5.35	1.21×10^{14}		1.04	4.75
Hydraulic loading = 4.0 l/min	-8800	17300	5.06	3.35×10^{15}		0.52	4.06
Combined Data	-8400	16500	5.34	8.42×10^{14}		2.52	2.70
FIRST ORDER							
Hydraulic loading = 2.5 l/min	-13800	27000	-0.57	3.65×10^{20}		2.01	4.75
Hydraulic loading = 4.0 l/min	-8000	15800	-1.34	3.03×10^{20}		0.64	4.06
Combined Data	-11200	22000	-0.92	3.08×10^{16}		4.23	2.70

5.3 Discussion of Concentration Dependency of Denitrification by RBC

Figures 16, 17, 18, and 19 show that denitrification rates achieved for hydraulic loadings of 2.5 l/min and 4.0 l/min can be represented by narrower 95% confidence levels for zero order kinetics.

In order to statistically evaluate the models, the ratios of residual mean squares to variances of observations were calculated.

$$\lambda = \frac{\sum s^2 / n}{\sigma^2}$$

where:

σ^2 = variance of observations,

$\sum s^2$ = sum of squares of residuals,

λ = ratio, and

n = degrees of freedom.

While there is no rigorous statistical test for the null hypothesis, larger deviations from unity indicate a real lack of fit for the model. On this basis, the smaller values of λ with zero order model for 2.5 l/min hydraulic loading and combined data tend to indicate superiority for the zero order model (Table 6).

TABLE 6
EVALUATION OF MODELS FOR ZERO ORDER AND
FIRST ORDER ASSUMPTIONS

Hydraulic Loading	Zero Order			First Order			
	$\frac{\Sigma S^2}{n}$	σ	λ	$\frac{\Sigma S^2}{n}$	σ^2	λ	F
2.5 l/min	0.16	0.16	1.00	0.37	0.27	1.33	4.09
4.0 l/min	0.27	0.31	0.87	0.90	0.99	0.90	3.34
Combined Data	0.20	0.13	1.54	0.82	0.37	2.17	2.34

FIGURE NO 16 TEMPERATURE DEPENDENCY OF
DENITRIFICATION RATE
First order kinetics
Hydraulic loading=2.5 l/min

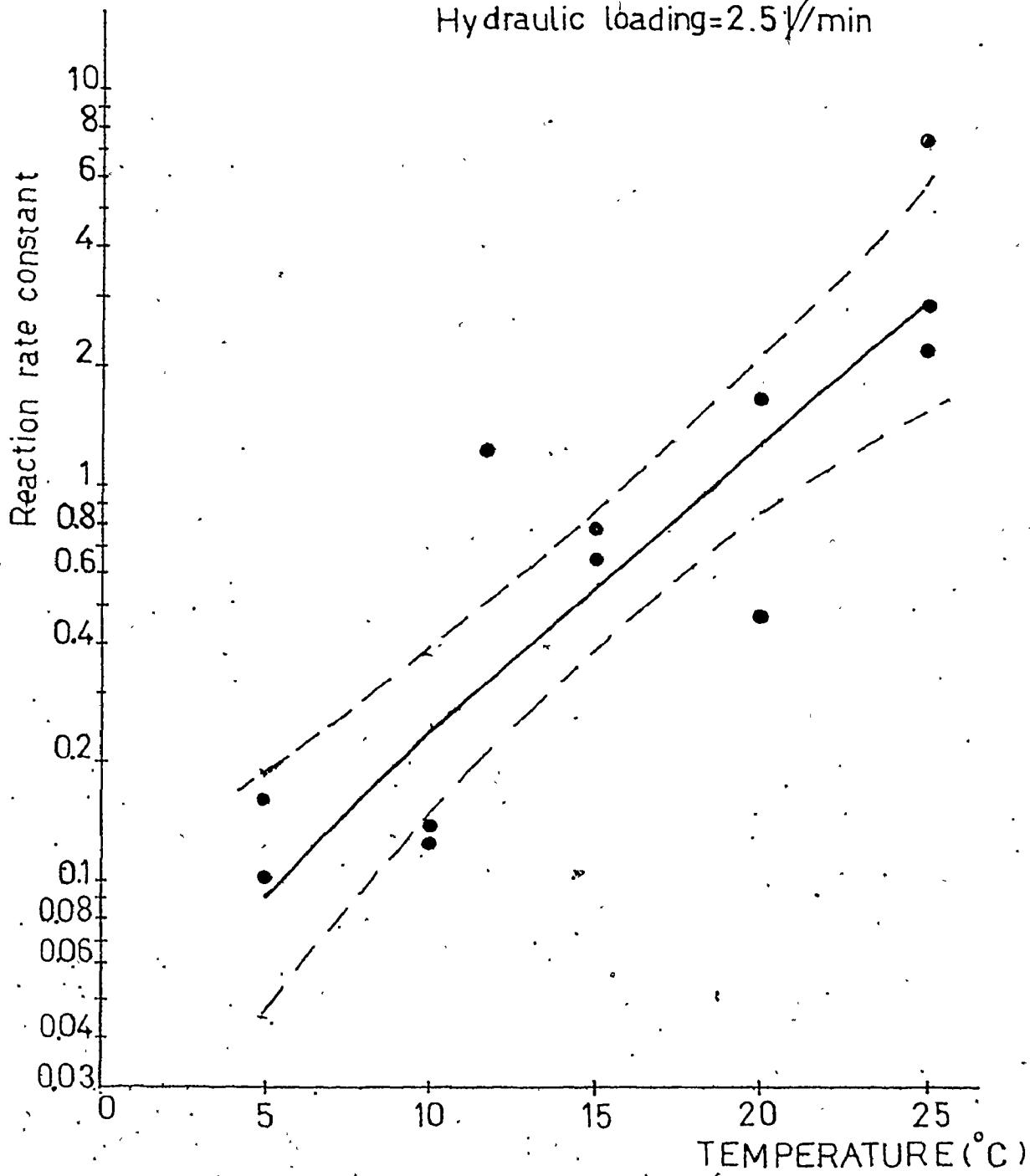


FIGURE NO 17 TEMPERATURE DEPENDENCY OF
DENITRIFICATION RATE
First order kinetics
Hydraulic loading=4.00 l/min

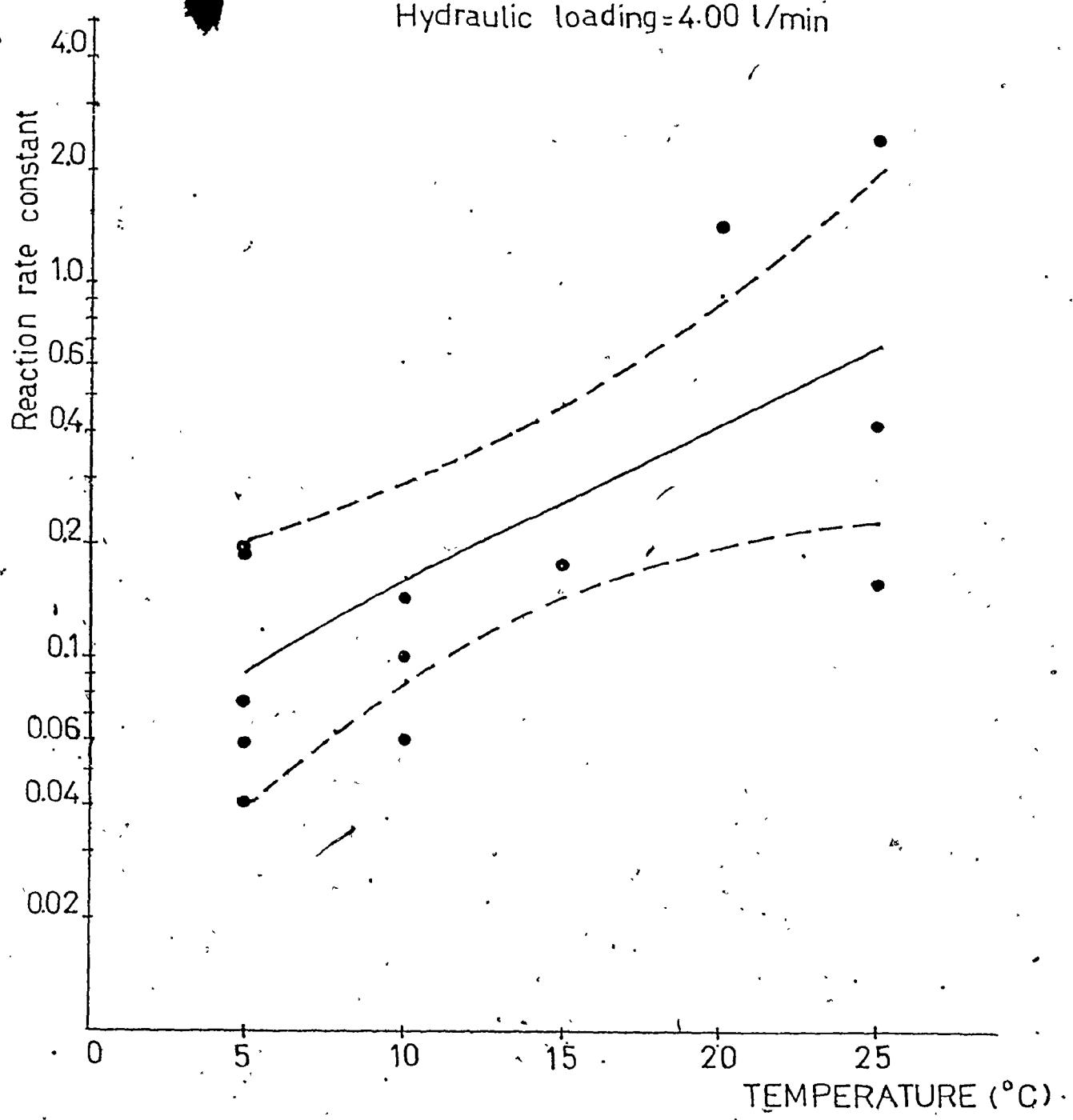


FIGURE NO. 18 TEMPERATURE DEPENDENCY OF
DENITRIFICATION RATE
Zero order kinetics
Hydraulic loading=2.5 l/min

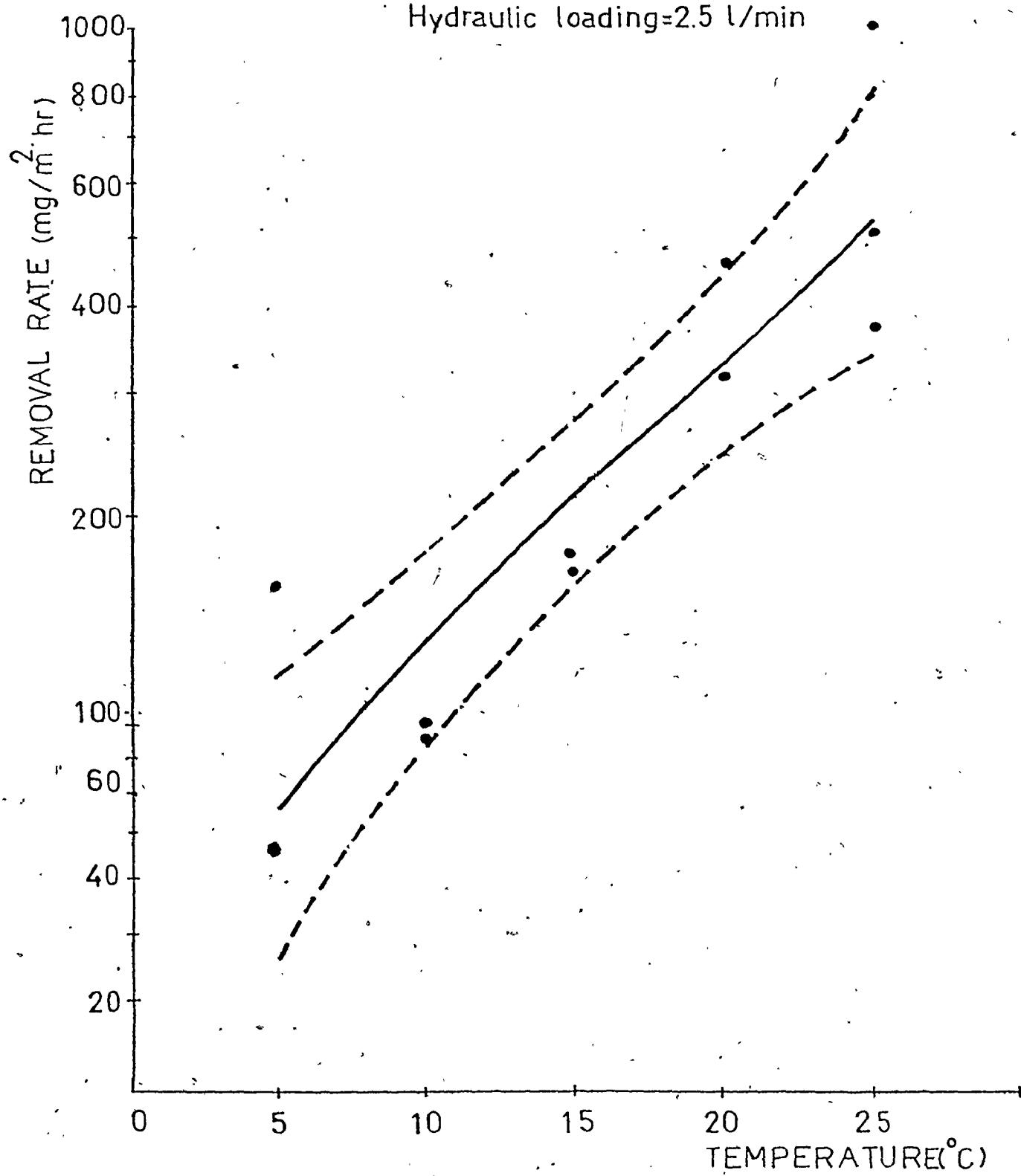


FIGURE NO 19 TEMPERATURE DEPENDENCY OF
DENITRIFICATION RATE
Zero order kinetics
Hydraulic loading = 4.0 l/min

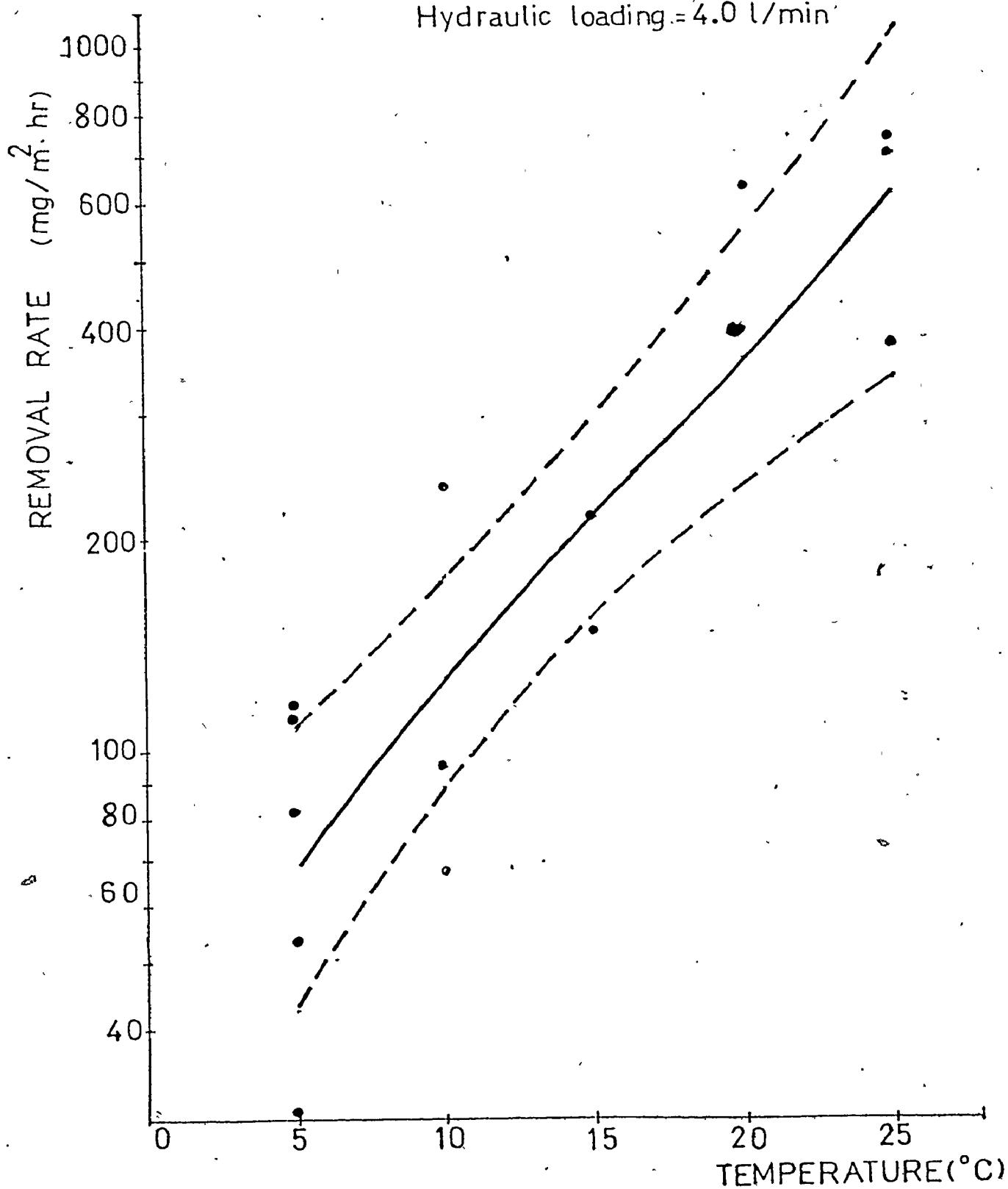


FIGURE NO. 20 TEMPERATURE DEPENDENCY OF DENITRIFICATION BY RBC

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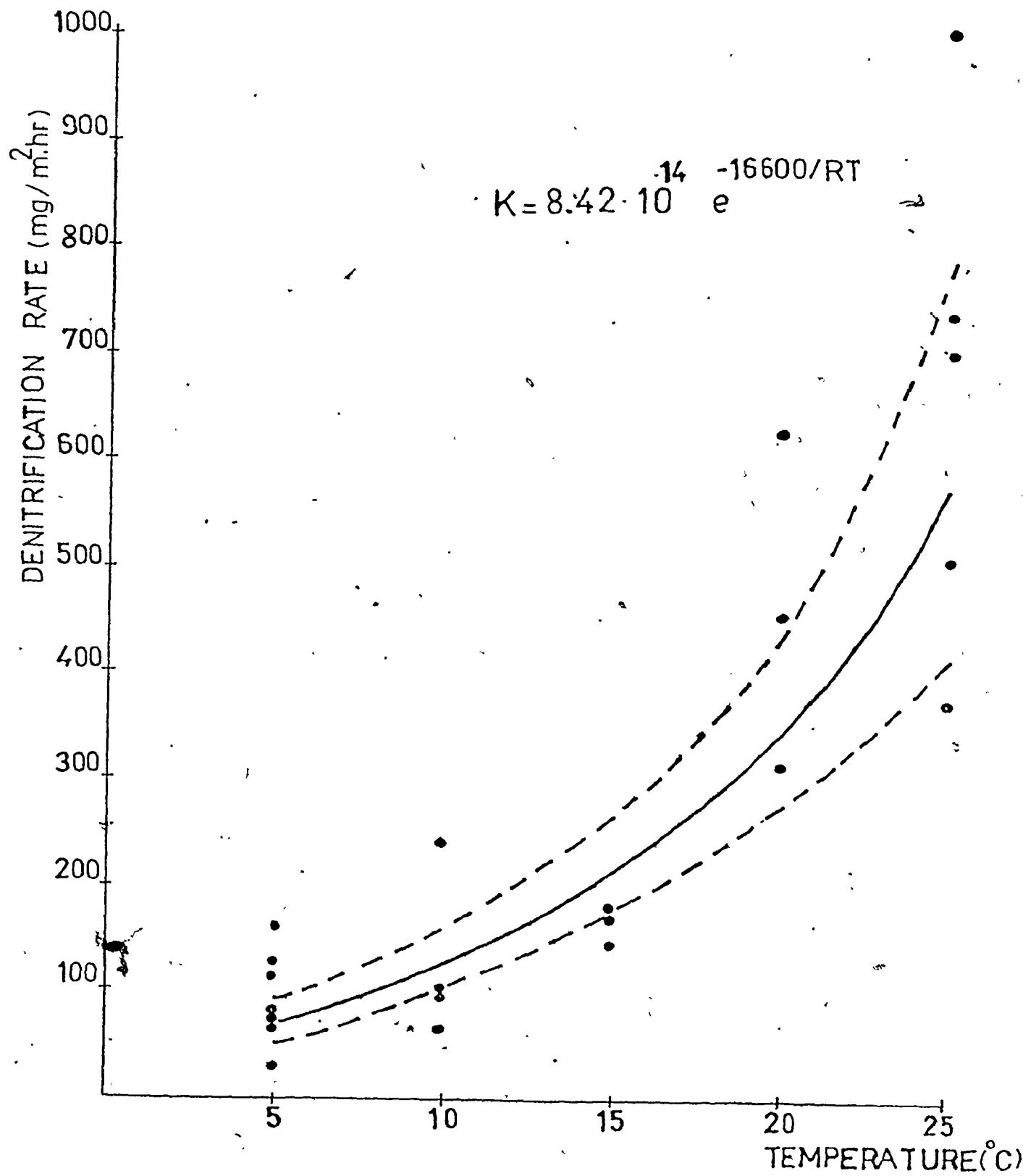
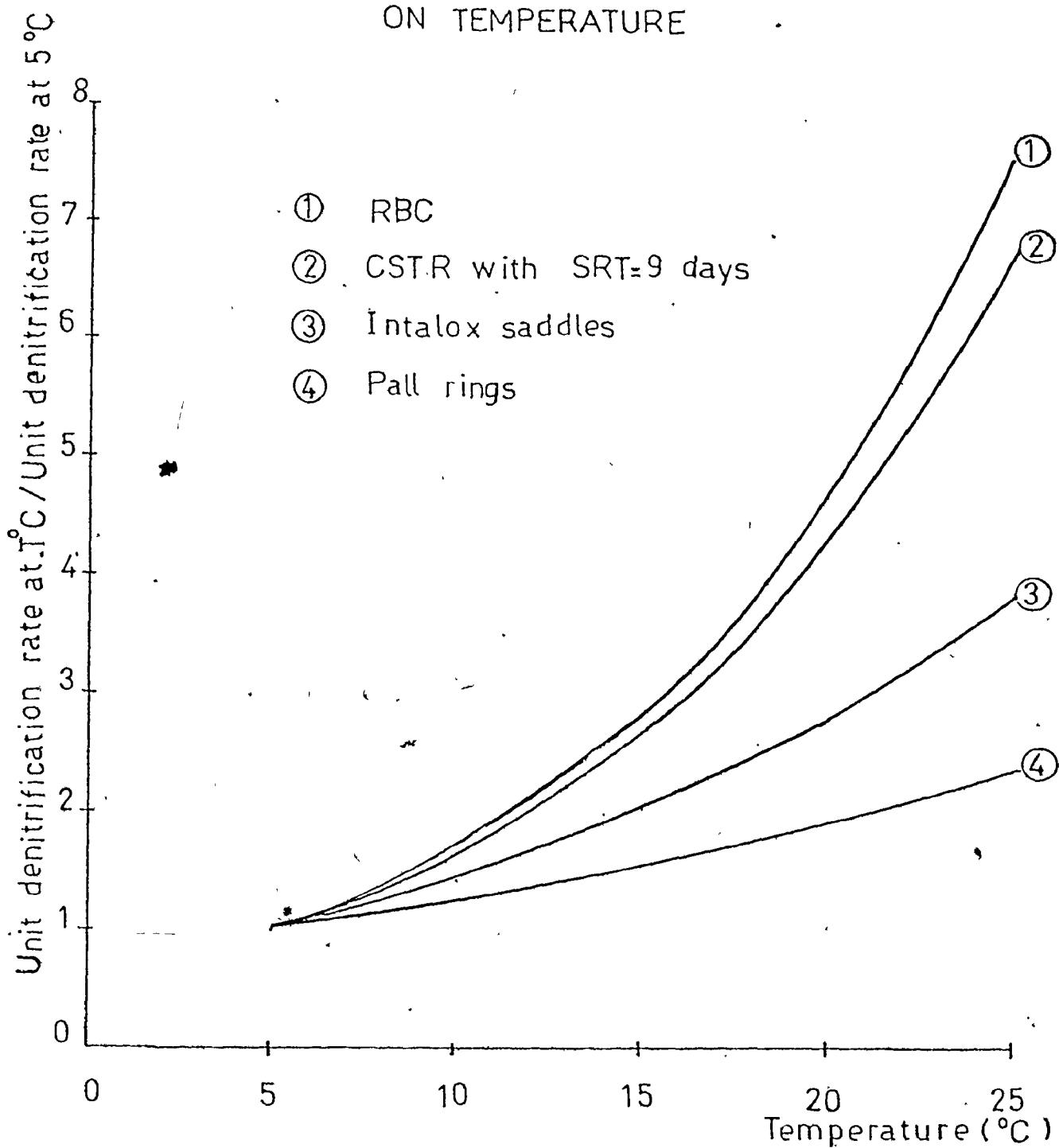


FIGURE NO 21 DENITRIFICATION RATE DEPENDENCY ON TEMPERATURE



5.4 Comparison of Temperature Dependency Curve with Other Literature Values

It is not surprising that, the energy of activation for denitrification by the RBC which is 16,600 cal/mole is not far different from the laboratory results of Dawson and Murphy (1971), energy of activation of 16,800 cal/mole, and Sutton's (1976), hydraulically well defined CSTR and Batch, energy of activation values of 15,300 and 15,900 cal/mole for different SRT values. Since the reactions are basically the same for different systems, only the constant A_0 should differ (Figure 20).

5.5 Comparison of Temperature Dependency of Various Denitrification Units

Figure 21 gives the Arrhenius temperature dependency of various denitrification units which were run under similar operating conditions. Details concerning the CSTR and Intalox packed column denitrification results are available from Sutton (1976) and the Pall Ring packed column results are from Wilson (1975). Figure 21 shows that the denitrification dependency of RBC system is greater than either the CSTR or packed columns.

An advantage of the RBC over other denitrification units arises from a volumetric comparison. The greater volumetric efficiency of the RBC over the packed columns is indicated in Figure 22 and over suspended growth systems in Figure 23. This coupled with the ease of operation makes the RBC one of the most competitive type of processes for denitrification.

FIGURE NO. 22 VOLUMETRIC EFFICIENCIES OF
VARIOUS DENITRIFICATION UNITS

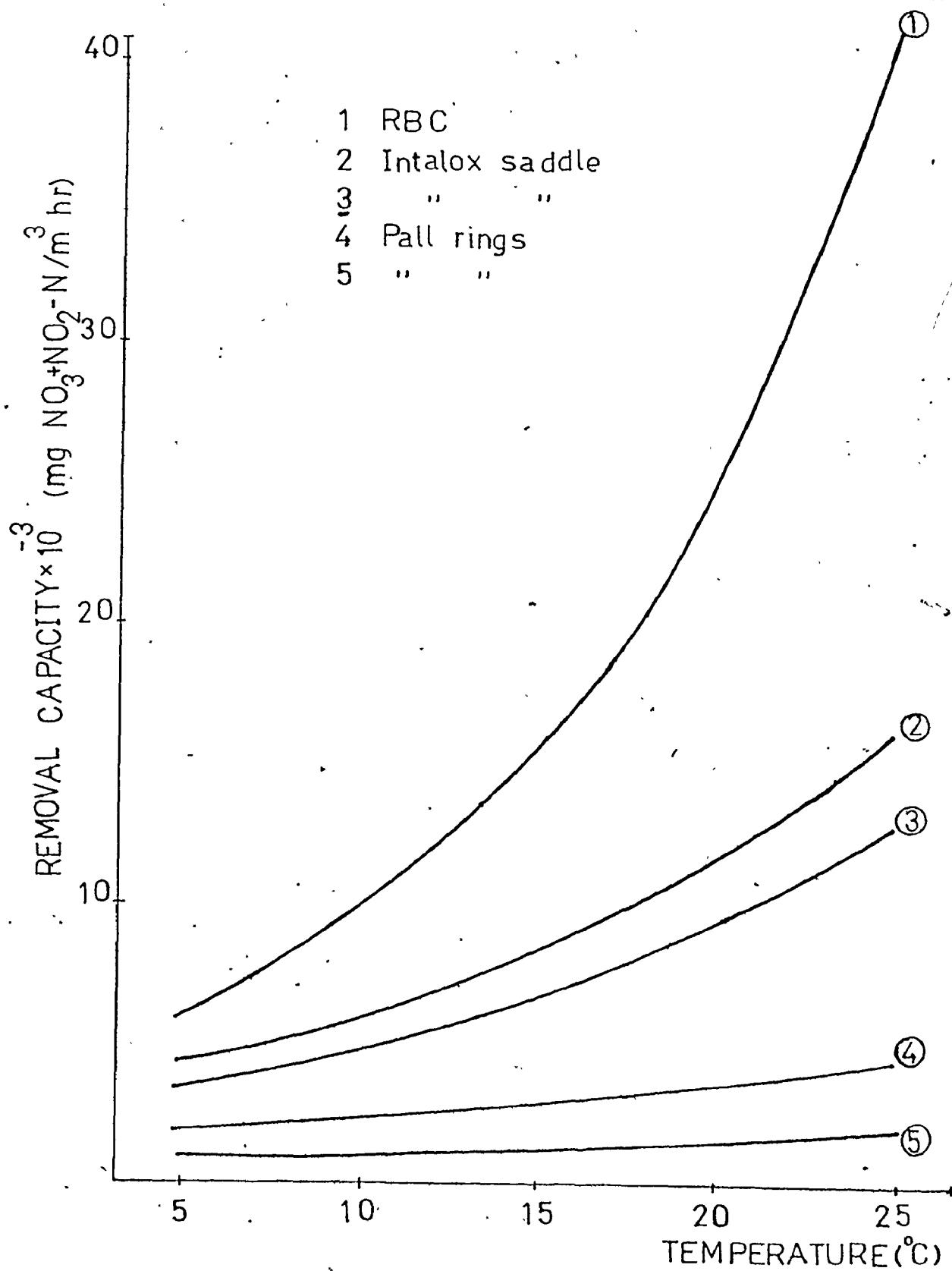
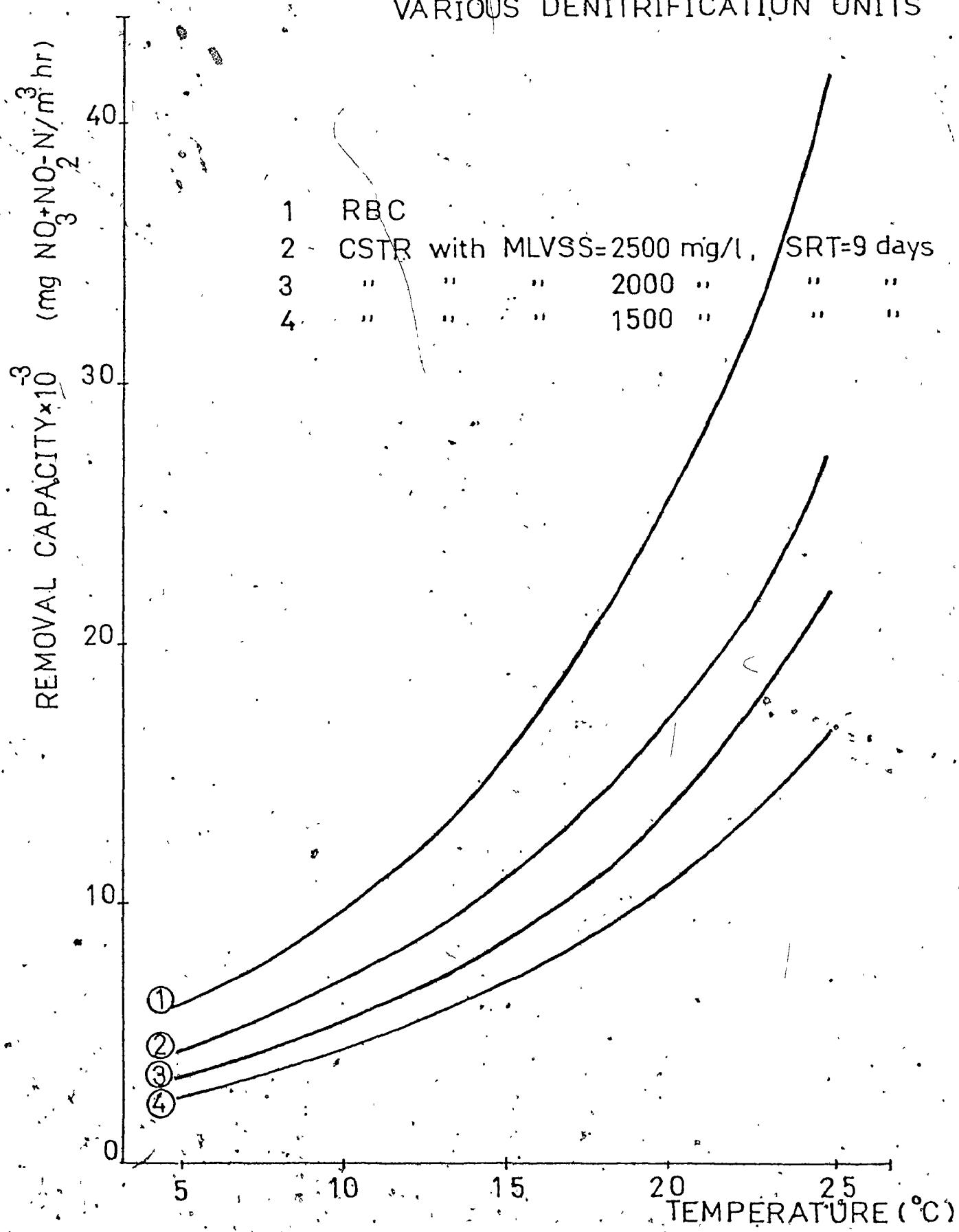


FIGURE NO 23 VOLUMETRIC EFFICIENCIES OF
VARIOUS DENITRIFICATION UNITS -67-



6: CONCLUSIONS

- (1) A general design method, based on hydraulic simulation can be derived, as a result of this study. Hydraulically, each compartment of completely submerged RBC can be modelled as 2 CSTR's with interchange flow. If the flow between the CSTR's can be determined, it is possible to predict whether the entire disc volume will be effective for denitrification.
- (2) The denitrification reaction rates can be expressed as zero order reactions, rather than first order reactions, for the $\text{NO}_3 + \text{NO}_2 - \text{N}$ concentration ranges studied for RBC.
- (3) The temperature dependency of denitrification obtained with an RBC is independent of the hydraulic loading for the range studied (2.5 to 4.0 l/min). It can be empirically described using an Arrhenius relationship with an apparent energy of activation of 16500 cal/g-mole.

- (4) The denitrification rates of the RBC are more sensitive to temperature changes than the corresponding rates for CSTR or packed columns under similar operating conditions.
- (5) The RBC has higher denitrification rates on a volumetric basis compared with either suspended growth systems (for MLSS range of 1500 to 2500 mg/l) or packed columns.

7. RECOMMENDATIONS

- (1) Further research is required to optimize the disc spacing to provide a sufficient area for growth and at the same time provide sufficient interflow transfer into the volume occupied by the discs.
- (2) Further experiments should be carried out to assess the effect of DO on denitrification rates, under actual pilot plant conditions. Nitrification and denitrification units might be operated more efficiently by keeping the oxygen level of the nitrification step at minimum so that the influent level of dissolved oxygen in the denitrification unit is minimized.
- (3) Further experiments should be done to study the denitrification rates at other hydraulic loadings.

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APPENDIX 1

TABLE 1-1

EXPERIMENTAL PROGRAM FOR 4 STAGES IN SERIES
(Altered experimental program due to
unreasonable amount of NO_3^- -N needed as spike)

Temperature	Hydraulic Loading	Date of Run	Run No.
5	4	10/7	R-5
10	4	17/7	R-6
15	4	3/7	R-4
20	4	23/7	R-7
20	4	24/7	R-8
20	4	30/7	R-9
25	4	17/6	R-1
25	4	19/6	R-2
25	4	16/8	R-11**
25	4	17/8	R-12**
25	4	21-22/8	R-12-13**
25	2	26/6	R-3

** These runs were for rate comparisons under natural pH and controlled pH conditions

TABLE 1-2
EXPERIMENTAL PROGRAM FOR SINGLE STAGE OPERATION

Temperature	Hydraulic Loading	Date	Run No.
5	2.5 l/min	17/9/75	R-19
5	"	18/9/75	R-20
10	"	22/10/75	R-31
10	"	23/10/75	R-32
15	"	11/9/75	R-17
15	"	12/9/75	R-18
20	"	10/10/75	R-27
20	"	11/10/75	R-28
25	"	25/9/75	R-22
25	"	26/9/75	R-23
25	"	28/9/75	R-24

TABLE 1-3

EXPERIMENTAL PROGRAM FOR SINGLE STAGE OPERATION

Temperature	Hydraulic Loading	Date	Run No.
5	4.0 l/min	17/9	R-19
5	"	19/9	R-20
5	"	21/9	R-21
5	"	22/10	R-31
10	"	22/10	R-31
10	"	23/10	R-32
20	"	10/10	R-27
25	"	3/10	R-25

* Due to operational difficulties double run for each temperature level was not obtained.

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FOUR STAGE IN SERIES OPERATION

UNFILTERED

FILTERED

ALL CONCENTRATIONS ARE IN PPM

Date	Designation	COD	D.O.	TKN	COD	TOC	NH ₃ -N	NO ₂ -N	NO ₃ -N	TKN
17-04	I.JNAF-C*PM	182	2.3	2.8	37	37	0.2	0.4	7.3	1.8
17-05	EE-1-C*PM		0.3		33	22	0.1	0.0	0.0	1.5
17-05	EE-2-C*		0.1		34	19	0.0	0.0	0.0	1.2
17-05	EE-3-C*PM		0.1		28	17	0.0	0.0	6.0	0.5
17-05	EE-C*PM	52	0.1	2.3	30	16	0.1	0.0	0.0	2.3
19-05	I.JNAF-C*PM	468	5.7	1.7	52	27	0.2	0.3	20.7	1.0
19-05	EE-1-C*PM		0.6		106	53	0.3	0.0	0.0	1.7
19-05	EE-2-C*PM		0.2		76	39	0.1	2.2	3.1	1.0
19-05	EE-3-C*PM		0.2		89	31	0.0	0.0	0.0	0.8
19-05	EE-C*PM	145	0.0	2.3	84	28	0.1	0.0	0.0	0.3
26-06	I.JNAF-C*PM	158		2.9	35	26	0.2	0.5	17.3	2.3
26-06	EE-1-C*				81	12	0.3	0.3	2.8	1.1
26-06	EE-2-C*				88	13	0.2	0.2	0.3	1.1
26-06	EE-3-C*				96	13	0.5	0.0	0.0	1.5
26-06	EE-C*	32		1.5	94	10	0.0	0.2	0.1	0.6
03-07	I.JNAF-C*PM	185	5.1	3.9	111	27	0.3	1.3	17.7	2.7
03-07	EE-1-C*		1.5		42	24	0.2	3.4	11.0	3.0
03-07	EE-2-C*		0.4		33	16	0.2	3.7	3.0	2.2
03-07	EE-3-C*		0.2		31	14	0.3	0.9	2.0	2.3
03-07	EE-C*	71	0.2	3.7	31	12	0.2	0.6	0.6	3.4
16-07	I.JNAF-C*PM	191	7.2	1.0	48	39	0.0	0.8	13.3	2.3
16-07	EE-1-C*		4.1		52	38	0.0	0.3	10.1	1.0
16-07	EE-2-C*		1.9		35	36	0.0	1.2	10.3	1.0
16-07	EE-3-C*		0.8		51	35	0.2	1.9	7.3	1.2
16-07	EE-C*	155	0.4	1.8	59	36	0.1	0.0	0.0	1.3
17-07	I.JNAF-C*PM	102	6.3	2.1	53	26	0.1	0.4	16.0	1.2
17-07	EE-1-C*		2.9		53	23	0.1	0.1	3.3	0.7
17-07	EE-2-C*		1.2		35	21	0.1	0.1	4.5	0.0
17-07	EE-3-C*		0.5		36	22	0.1	0.1	1.5	0.6
17-07	EE-C*	32		1.1	30	13	0.1	0.7	1.3	0.0
23-07	I.JNAF-C*PM	27+	7.2		51	19	0.0	0.0	33.4	1.0 CL
23-07	EE-1-C*PM		1.4		39	12	0.1	1.4	32.0	1.8 CL
23-07	EE-2-C*PM		0.2		36	10	0.1	1.0	21.4	1.4 CL
23-07	EE-3-C*PM		0.2		36	14	0.1	1.4	22.6	1.2 CL
23-07	EE-C*PM		4.4		26	10	0.2	0.9	25.0	2.0 CL
29-07	I.JNAF-C*PM	320	5.4	1.3	345	140	1.0	0.1	12.5	0.5 OP
29-07	EE-1-C*PM		1.1		465	65	0.5	0.6	9.3	1.3 OP
29-07	EE-2-C*PM		0.5		338	58	0.4	0.5	0.0	0.3 OP
29-07	EE-3-C*PM		0.15		326	45	0.6	0.2	2.1	1.3 OP
29-07	EE-C*PM	232	0.10	2.4	192	34	0.5	0.0	0.0	1.1 OP
15-08	I.JNAF-C*PM	301	4.3	1.7	980	30	0.3	0.1	44.0	1.1
15-08	EE-1-C*PM		1.7		620	85	0.3	0.3	23.0	0.8
15-08	EE-2-C*PM		0.3		530	53	0.3	0.4	13.0	1.0
15-08	EE-3-C*PM		0.1		630	57	0.3	0.4	5.3	0.9
15-08	EE-C*PM	303	0.6	1.9	390	40	0.3	0.3	0.7	1.0
17-08	I.JNAF-C*PM	300		2.2	625	55	0.3	0.1	54.0	1.0

FOUR STAGE IN SERIES OPERATION

1	03	EE-1C* ^{PL}	51	6	63	5.3	0.4	42.0	6.3			
1	03	EE-2C* ^{PL}	350		51	3.4	0.6	13.6	1.3			
1	03	EE-3C* ^{PL}	277		39	3.3	0.5	8.3	1.3			
17	03	EE-C* ^{PL}	212		41	0.3	0.3	1.3	0.3			
21	03	IIONEE-C* ^{PL}	525	3.5	11.3	560	5.8	8.6	0.2	32.0	3.9	PC
21	03	EE-1C* ^{PL}	0.62		474	37	7.0	1.3	13.6	0.2	PC	
21	03	EE-2C* ^{PL}	0.4		385	42	5.5	1.5	5.6	0.3	PC	
21	03	EE-3C* ^{PL}	0.26		355	34	5.0	1.0	1.0	6.1	PC	
21	03	EE-C* ^{PL}	265	0.16	41.2	31	5.3	0.1	0.2	0.8	PC	
21	03	IIONEE-C* ^{PL}	565	12.0	625	70	8.8	0.2	43.0	3.7	PC	
21	03	EE-1C* ^{PL}	530		55	5.3	1.3	21.0	7.1	PC		
21	03	EE-2C* ^{PL}	475		45	6.4	1.1	6.5	7.1	PC		
21	03	EE-3C* ^{PL}	480		39	4.9	0.8	1.2	5.5	PC		
21	03	EE-C* ^{PL}	330	10.2	395	34	5.3	0.4	3.2	5.9	PC	

CL STANDS FOR CARBON LIMITATION IN THE SYSTEM

PD STANDS FOR PUMPING DISORDER

PO STANDS FOR VARIOUS OPERATIONAL PROBLEMS

PC STANDS FOR PH CONTROL BY BUFFER ADDITION

PO STANDS FOR OXYGEN LEVEL CONTROL BY STRIPPING BY '12 GAS

IIONEE-C*^{PL} AND IONAF-C* STANDS FOR THE COMPOSITE SAMPLE TAKEN FROM FEED AFTER METHANOL ADDITION

EE-1C*, EE-2C*, EE-3C*, AND EE-C* STAND FOR THE SAMPLES TAKEN FROM FIRST, SECOND, THIRD AND FOURTH STAGES, RESPECTIVELY

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FOUR STAGE IN SERIES OPERATION

Date	Designation	MLSS	MLVSS	Flow Rate	Alkalinity	pH	HACH-NO ₃ -N	Temp.
17-05	I DNAF-C+P1			3.9	7.56	14.0	24.0	
	EE-1C+PM			3.9	7.56	4.75	25.2	
	EE-2C+PM			3.9	7.65	1.65	25.4	
	EE-3C+PM			3.9	7.55	0.60	25.4	
	EE-4C+PM			3.9	7.50	0.40	25.4	
19-05	I DNAF-C+P1			4.0		18.00	25.8	
	EE-1C+PM			4.0		5.10	25.4	
	EE-2C+PM			4.0		2.90	25.5	
	EE-3C+PM			4.0		1.10	25.5	
	EE-4C+PM			4.0		0.60	25.5	
20-05	I DNAF-C+P1	15	11	2.0	125	7.28	20.0	
	EE-1C+PM	143	107	2.0	168	7.70	25.1	
	EE-2C+PM	35	27		194	7.75	25.1	
	EE-3C+PM	136	28	2.0	202	7.72	25.1	
	EE-4C+PM	29	22	2.0	240	7.70	25.1	
23-05	I DNAF-C+PM			4.0	125	7.48	14.4	
	EE-1C+PM	68	48	4.0	138	7.72	14.4	
	EE-2C+PM	36	28	4.0	171	7.84	14.4	
	EE-3C+PM	29	24	4.0	175	7.92	14.4	
	EE-4C+PM	24	21	4.0	181	7.92	14.4	
18-07	I DNAF-C+PM	14	13	4.0	101		19.0	
	EE-2-C+PM	43	33	4.0	123	7.68	19.0	
	EE-3-C+PM	28	21	4.0	130	7.22	19.0	
	EE-4C+PM	26	19	4.0	143	7.78	19.0	
23-07	I DNAF-C+	15	7	4.1	69	7.52	19.0	
	EE-1C+	12	8	4.1	127	7.62	20.0	
	EE-2C+	18	14	4.1	167	8.22	20.0	
	EE-3C+	19	15	4.1	181	8.32	21.0	
	EE-4C+	12	9	4.1	183	8.45	21.0	
29-07	I DNAF-C+			4.0	127	7.65	19.0	
	EE-1C+			4.0	241	7.72	20.0	
	EE-2C+			4.0	300	7.83	20.0	
	EE-3C+			4.0	317	7.65	20.0	
	EE-4C+			4.0	327	7.75	20.0	
16-08	I DNAF-C+PM			4.0	108	7.38	22.0	
	EE-1C+PM			4.0	181	7.30	23.0	
	EE-2C+PM			4.0	243	8.20	23.0	
	EE-3C+PM			4.0	277	8.25	23.0	
	EE-4C+PM			4.0	281	8.25	23.0	
18-08	I DNAF-C-P1	76	50	+0.1	65	7.30	47.5	21.5
	EE-1C+PM	35	32	+0.1	125	7.95	31.25	21.5
	EE-2C+PM	66	50	+0.1	230	8.40	15.0	21.5
	EE-3C+PM	54	43	+0.1	271	8.50	3.00	21.5
	EE-4C+PM	26	19	+0.1	281	8.48	1.75	21.5
21-08	I DNAF-C+P1	45	27	+0.2	156	7.28	42.50	23.2
	EE-1C+P1	69		+0.0	271	7.53	20.00	

ROTATING BIOLOGICAL CONTACTOR
FOUR STAGE IN SERIES OPERATION

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EE-2C*PM	74	53	+0.0	265	7.62	0.00	24.00
EE-3C*PM	56	37	+0.0	344	7.62	3.00	24.0
EE-4C*PM	64.	31.	+0.0	371	7.60	0.00	24.0

I DHEBE-C AND IDNAF-C STANDS FOR THE COMPOSITE SAMPLE TAKEN FROM FEED AFTER METHANOL ADDITION

EE-1C+, EE-2C+, EE-3C+ AND EE-4C+ STAND FOR THE SAMPLES TAKEN FROM FIRST, SECOND, THIRD AND FORTH STAGES RESPECTIVELY
TEMPERATURES ARE GIVEN IN DEGREE CENTRIGRADE

FLOW RATES ARE GIVEN IN LITER/MINUTE

MLSS AND MLVSS MEASUREMENTS ARE GIVEN IN MILIGRAM/LITER

HACH NO3-N AND ALKALINITIES ARE GIVEN IN MILLIGRAM /LITER

SINGLE STAGE OPERATION

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DATE	SAMPLE NAME	COD OO	UNFILTED		FILTERED		TOC	NH3N	NO2N	NO3N	TKN	NOTE
			TKN OO	COD	TKN OO	COD						
29 08	IONEBE-C*PM	278	2.16	13.9	294	60	8.8	0.6	13.0	10.6		
29 08	EE-1C*PY	193	0.45		161	33	7.2	0.4	2.1	11.1		
29 08	EE-4	56	0.46	11.7	145	13	8.5	1.6	12.4	10.4	P0	
03 09	IONBE-C*PY	261	5.50	3.7	274	53	6.4	0.4	3.6	7.0		
03 09	EE-1C*	40	1.35	6.3	72	11	5.5	0.5	7.2	6.3	P0	
03 09	EE-4C*PY	213	1.45	6.5	221	45	5.1	0.3	4.4	5.5		
11 09	IONAF-C*1PM	184	5.7	8.2	210	42	6.8	0.3	15.0	7.5		
11 09	EE-1C*1PM	46	9.2		88	18	0.9		26.4	8.1	P0	
11 09	EE-4-C*1PM	124	1.30	6.9	149	30	5.2	0.7	3.1	6.3		
11 09	IONAF-C*3PM	126	5.6	8.8	184	43	7.2	0.2	12.0	8.1		
11 09	EE-1C*3PM	54	9.1		82	13	8.1	0.7	25.3	8.1	P0	
11 09	EE-4-C*3PM	122	1.30	7.1	173	30	5.9	0.7	5.0	7.1		
12 09	IONAF-C*3PM	195	6.2	6.3	183	68	4.4	0.6	14.0	5.7		
12 09	EE-1C*3PM	50	2.3	5.3	30	45	3.9	1.5	47.5	5.6	P0	
12 09	EE-4-C*3PM	118	1.5	3.9	71	51	1.7	1.9	3.3	3.5		
12 09	IONAF-C*1PM	178	6.3	5.3	164	71	3.4	0.9	11.1	4.5		
12 09	EE-1C*1PM	38	2.20	4.7	33	41	3.3	1.3	42.7	4.3	P0	
12 09	EE-4C*1PM	104	1.30	5.8	69	63	1.4	1.9	2.3	2.6		
17 09	IONAF-C*1PM	166	6.9	4.8	50	40	3.2	0.3	12.9	4.6		
17 09	EE-1-C*1PM	56	3.9	4.3	35	15	2.8	0.6	11.0	3.1		
17 09	EE-4-C*1PM	139	3.5	4.2	41	32	2.5	1.0	3.2	3.0		
17 09	IONAF-C*3PM	37	6.9	5.1	49	37	3.2	0.3	13.7	3.9		
17 09	EE-1-C*3PM	55	3.9	4.9	33	15	3.1	0.5	13.1	3.9		
17 09	EE-4-C*3PM	138	3.5	4.9	35	31	2.9	1.0	9.3	3.6		
18 09	IONAF-C*3PM	149		3.0	45	43	0.1	0.2	24.0	1.8		
18 09	EE-1C*3PM	59		2.4	31		0.2	0.6	32.4	1.3		
18 09	EE-4C*3PM	120		3.1	35	28	0.1	1.6	12.6	1.7		
18 09	IONAF-C*1PM	157	6.70	2.6	43	37	0.1	0.2	25.0	1.6		
18 09	EE-1C*1PM	53	4.30	2.1	28	16	0.1	0.6	23.0	1.5		
18 09	EE-4C*1PM	120	2.70	2.2	33	31	0.1	1.6	19.0	1.4		
21 09	IONAF-C*5PM	24	6.30	1.0	16	9	0.0	0.1	15.3	0.8		
21 09	EE-1C*5PM	58	1.70	2.2	20	15	0.0	0.3	12.8	2.0	P0	
21 09	EE-4C*5PM	152	3.50	2.3	31	40	0.0	0.5	9.9	2.0		
21 09	IONAF-C*7PM	40	6.7	1.2	40	9	0.0	0.0	15.3	1.2		
21 09	EE-1C*7PM	55	1.70	1.2	35	15	0.0	0.3	24.7	0.9	P0	
21 09	EE-4C*7PM	197	2.50	1.3	66	41	0.0	0.5	12.5	1.3		
26 09	IONAF-C*1PM	193	5.4	1.1	163	51	0.1	1.0	11.7	0.7		
26 09	EE-1C*1PM	23	0.3	4.3	22	13	0.0	4.1	7.2	0.3	P0	
26 09	EE-4C*1PM	98	0.2	1.6	44	13	0.1	1.4	1.5	1.3		
28 09	IONAF-C*1PM	267	5.7	1.3	149	53	0.1	2.2	23.7	1.6		

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SINGLE STAGE OPERATION

03 10	EE-4C*1P1	9-	3.3	3.4	39	23	1.1	2.6	1.1	1.3
07 10	IONAF-C*1PM	247	5.36	1.1	166	59	0.5	3.6	25.4	1.3
07 10	EE-4-C*1P1	123	0.66	3.2	55	31	0.3	3.6	2.1	2.3
03 10	IONAF-C*3PM	238	4.05	1.3	136	59	0.2	3.8	25.2	1.3
03 10	EE-4-C*3P1	167	0.65	1.6	57	31	0.2	4.2	2.4	1.4
03 10	IONAF-C*2P1	114	2.40	1.6	239	23	0.3	3.1	33.9	1.3
09 10	EE-4-C*1PM	156	0.80	3.8	125	27	0.5	3.5	0.0	3.5
10 10	IONAF-C*4PM	326	5.4	1.7	354	65	0.2	2.8	27.2	1.3
10 10	EE-1-C*4P1	159	0.50	1.8	165	33	0.2	2.2	4.7	1.7
10 10	EE-4-C*2P1	126	-0.8	1.9	120	27	0.3	3.1	6.9	1.6
11 10	IONAF-C*6PM	355	5.3	1.3	160	61	0.2	2.9	26.1	1.3
11 10	EE-10-C*6P1	102	6.5	1.3	171	35	0.2	2.3	13.7	1.7
15 10	IONAF-C*2P1	100	2.5	2.5	156	32	0.0	1.9	22.1	1.2
15 10	EE-10+	218	1.3	1.4	204	41	0.4	4.2	7.9	1.0
15 10	EE-4C*	170	0.7	1.1	58	19	0.4	3.9	2.7	0.0
16 10	IONAF-C*	177		1.1	180	45	0.7	0.6	0.0	0.9
16 10	EE-10+	192		0.7	193	35	0.3	3.5	10.2	0.2
17 10	EE-C*	95		2.0	50	15	0.5	2.1	0.0	1.3
22 10	IONAF-C*	139	5.3	4.8	119	27	0.2	1.1	19.9	4.1
22 10	EE-10+	86	1.7	3.2	108	18	0.2	1.9	13.4	2.6
23 10	IONAF-C*2P1	150		3.0	143	29	0.2	0.5	17.5	2.1
23 10	EE-10+	83		4.2	115	20	0.2	2.0	13.0	3.3
23 10	EE-40+	72		2.6	85	47	0.5	2.6	10.3	1.7

CL STANDS FOR CARBON LIMITATION IN THE SYSTEM

PO STANDS FOR PUMPING DISORDER

OP STANDS FOR VARIOUS OPERATIONAL PROBLEMS

PC STANDS FOR PH CONTROL BY BUFFER ADDITION

OC STANDS FOR OXYGEN LEVEL CONTROL BY STRIPPING BY N₂ GASLVEBE-4 AND IONAF-C* STANDS FOR THE COMPOSITE SAMPLE TAKEN FROM FIRST
FIVE STAGES RESPECTIVELYEE-10+ AND EE-C* STAND FOR THE COMPOSITE SAMPLE TAKEN FROM FIRST
AND FORTH STAGES RESPECTIVELY

SINGLE STAGE OPERATION

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DATE	SAMPLE	MLDS	MLVSS	FLOW RATE	ALK- HACH NO3-N	TEMP- ERATURE
22 03	IDNAF-C PM	27	22		7.13	16.00
	EE-1C*PM		49	1.80	7.25	4.00
	EE-4C*PM	21	12	0.97	7.13	16.00
03 09	IDNAF-C-P1				7.31	9.20
	EE-1C*P1			0.97	7.25	7.00
	EE-4C*P1			1.30	3.35	4.00
11 09	IDNAF-C PM	17			150	12.50
	EE-1C*PM	17		3.90	144	11.50
	EE-4C*PM	15		2.45	186	5.00
12 09	IDNAF-C*P1	25	15		148 7.45	17.00
	EE-1C*PM	13		3.9	148 7.45	16.00
	EE-4C*PM	38	27	2.45	188	5.00
17 09	IDNAF-C*PM					15.00
	EE-1C*P1			2.47		14.00
	EE-4C*PM			3.90		8.50
18 09	IDNAF-C*P1	3	2		108 7.05	17.00
	EE-1C*PM	9	5	2.50	115	15.50
	EE-4C*PM	14	9	4.00	124	17.50
21 09	IDNAF-C*PM	6			7.33	14.40
	EE-1C*PM	12		4.00	7.70	15.50
	EE-4C*PM	18		4.00	7.52	17.00
26 09	IDNAF-C*PM					25.70
	EE-1C*PM			4.00		25.00
	EE-4C*PM			2.50		6.00
03 19	IDNAF-C*PM					25.00
	EE-1C*PM					15.33
	EE-4C*PM			4.10		
08 10	IDNAF-C*P1				7.03	18.00
	EE-1C*P1			+.00	7.35	20.50
	EE-4C*PM				7.15	22.00
10 10	IDNAF-C*P1			2.50		20.00
	EE-1C*P1			+.00	7.45	4.50
	EE-4C*P1					20.00
15 10	IDNAF-C*P1				138 7.10	
	EE-1C*PM			3.20	185 7.40	8.00
	EE-4C*P1			1.80	7.38	4.00
16 10	IDNAF-C PM					
	EE-1C*P1			3.20		15.25
	EE-4C*P1			1.80		15.00
22 10	IDNAF-C P1					
	EE-1C*P1			4.00		
	EE-4C*PM			2.50		13.00

APPENDIX 2

The following are included in this appendix:

Computer Program DYEL:

Calculates dye recovery, mean dye residence time, residence time distribution, and also lists the dye study results.

Program FLBTCS:

Calculates the average flow rates between inner and outer CSTR'S of a single stage RBC. Also it gives the dye concentration of inner CSTR at any time after pulse dye injection.

Program DEDBIS:

Calculates ideal response of two CSTR with interchange flow for pulse dye injection using Dedrick's and Bischoff's formula.

Program LINRG

Program CONF95

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PROGRAM DYEI; {INPUT;TAPE5=INPUT;OUTPUT:TAPE6=OUTPUT}

PROGRAM FOR THE CALCULATION OF DYE RECOVERY, MEAN DYE RESIDENCE TIME, E (RESIDENCE TIME DISTRIBUTIONS) AND FIC/COI VALUES.

DEF = THEORETICAL DETENTION TIME OF THE REACTOR, IN MIN.
 DYE = AMOUNT OF THE DYE ADDED, IN LIT.
 DT = THE SAMPLING INTERVAL, IN MIN.
 DT = RESIDENCE TIME DISTRIBUTION BASED ON DIMENSIONLESS TIME, DIMEN-
 SIONLESS AREA UNDER ET CURVE, DIMENSIONLESS TIME BASED ON THE DYE MEASUREMENTS, IN MIN.
 TBAR = MEAN RESIDENCE TIME
 THETA = DIMENSIONLESS TIME
 VTOA = VOLUME OF THE REACTOR, IN LIT.
 VT = HYDRAULIC LOADING, IN LIT.
 N = THE NUMBER OF SAMPLES COLLECTED
 C(N) = THE MEASURED DYE CONCENTRATIONS AT TIME=N*DT AFTER PULSE
 DYE INJECT, IN PPB

VTO TA, VT, DY IN, DY CON, DT, N, DET INPUTS

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VTOA,VT,DET,DYIN,DYCON,C(N),PER,TEBAR,DT
      OUTPUTS
DIMENSION RE(500),TIME(500),CUL(500),FIT(500)
DIMENSION ET(500)
DIMENSION C(500)
DIMENSION CTBDET(500),THETA(500),F(500),Q(500),E(500),Y(500),X(500)
DIMENSION CI(500),CB(500),T(500)
DIMENSION TE(500)
DIMENSION STE(500)

      READ(5,1) VTOTA,VI,DYIN,DYCON,DT
      1 FORMAT(3F10.4,E10.2,F10.4)
      READ(5,2)N,DET
      2 FORMAT(17,F10.4)
      READ(5,3)(C(I),I=1,N)
      3 FORMAT(5F10.2)

      WRITE(6,51)
      51 FORMAT(10X1'////////////////////////////')
      WRITE(6,997)H INPUT DATA, //)
      997 FORMAT(20X,11H INPUT DATA, //)
      WRITE(6,995,#FOUR STAGE IN SERIES OPERATION#,10X)
      995 FORMAT(10X1#)
      WRITE(6,111)TEMPERATURE=15 DEG. C#,10X)
      111 FORMAT(10X1#VT)
      WRITE(6,306)FLLOW RATE=#,F6.2,#LIT./MIN.,#,5X)
      306 FORMAT(10X1#FLOTA,VOLTA OF THE REACTOR#=,F7.1.#LIT.,10X)
      WRITE(6,305)VOLUME OF THE REACTOR#=,F7.1.#LIT.,10X)
      305 FORMAT(10X1#DETEN(10X7))
      WRITE(6,307)THEORETICAL DETENTION TIME#=,F7.2,#MIN#,10X,1X)
      307 FORMAT(10X1#DT#=,F5.2,#MIN#,10X)
      102 FORMAT(10X1#DT#=,F5.2,#MIN#,10X)

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WRITE(6,308) DYN

308 FORMAT(10X, #THE AMOUNT OF DYE INJECTED= #,F7.4, #LIT, #,10X)

WRITET(6,309) DYCUN

FORMAT(10X, #CONCENTRATION OF THE DYE ADDED= #,F10.2, #PPB#, //)

309 WRITET(6,990) DYE MEASUREMENTS#, //

990 FORMAT(10X, #DYE MEASUREMENTS#, //)

991 FORMAT(5X, 5F10.2, //)

C CALCULATION OF TIME VALUES

TIME(1)=0.0

DO 10 I=2,N

KL=I-1

TIME(I)=TIME(KL)+DT

C CALCULATIONS OF THETA VALUES

THEΤΑ(I)=0.0

DO 90 I=2,N

THEΤΑ(I)=TIME(I)/DEI

CONTINUE

C CALCULATION OF DYE RECOVERY

CUL(1)=0.0

DO 800 I=2,N

LUL=I-1 C(I)=C(LUL)*DT*VTL*10.**(-6)

AΜT=0.5*(C(I)+C(LUL))*DT*VTL*10.**(-6)

CUL(I)=CUL(LUL)+AMT

CONTINUE DΥΙΝ*DΥΙΝ*10.**(-6)

DΥΕ=DΥΙΝ/DΥE*100.

CTBDET(I)=0.0

C1(I)=0.0

DO 78 I=2,N

KL=I-1 C(KL)=C(I)+C(I)**0.50

T(I)=DT*0.5*D

CTBDET(I)=CB(KL)*T(I)*DT+CTBDET(KL)

C1(I)=C(I)+CB(KL)*DT

CONTINUE

TEBAR=CTBDET(N)/C1(N)

CONVERSION OF E CURVE TO F CURVE

C CALCULATION OF VALUES

G(I)=0.0

DO 8 I=2,N

KM=I-1 Q(KM)=0.5*(C(I)+C(KM))*DT

CONTINUE

Q(I)=Q(I)+DTQ#

DE(I)=E(I)-TOTQ#

E(I)=E(I)+DET#

CONTINUE

AMT=0.0

F(I)=AMT

DO 4 I=2,N

ILK=I-1

F(I)=F(ILK)+0.5*(E(I)+E(ILK))*DT

INPUT DATA

FOUR STAGE IN SERIES OPERATION
TEMPERATURE = 10 DEG. C
FLOW RATE = 4.00 LIT / MIN.
VOLUME OF THE REACTOR = 408.0 LIT.
THEORETICAL DETENTION TIME = 102.0 MIN
DT = 4.00 MIN
THE AMOUNT OF DYE INJECTED = 0.220 LIT.
CONCENTRATION OF THE DYE ADDED = 238000.00 PPM

DYE MEASUREMENTS

0.00	6.50	26.00	42.00	60.00
73.00	85.00	95.00	103.00	108.00
108.00	108.00	107.00	104.50	101.00
97.00	93.50	90.00	86.00	82.00
79.00	75.50	72.00	68.50	65.00
61.00	58.00	54.50	51.50	49.00
46.50	44.00	42.00	40.50	38.50
37.00	35.00	33.50	32.00	30.50
29.00	28.00	26.50	25.50	24.50
23.50	22.50	21.75	21.00	20.00
19.50	18.50	18.00	17.50	16.50
16.00	15.25	14.50	14.00	13.25
12.75	12.00	11.50	11.00	10.50
10.00	9.50	9.00	8.50	8.25
8.00	7.50	7.25	7.00	6.75
6.50	6.40	6.25	6.10	6.00
5.75	5.50	5.40	5.25	5.20
5.05	5.00	4.90	4.80	4.50
4.50	4.25	4.00	3.90	3.75
3.50	3.50	3.40	3.25	3.00
3.00	2.90	2.75	2.60	2.50

TEST RESULTS AND CALCULATED VALUES

DYE RECOVERY = 100.17 PER CENT

MEAN DYE RESIDENCE BASED ON C=110.67 MIN
MEAN DYE RESIDENCE BASED ON E=110.67 MIN

THETA	E	ET	F
0.000	0.000	0.000	0.000
.039	.000	.051	.001
.078	.002	.202	.006
.118	.003	.327	.016
.157	.005	.467	.032
.196	.006	.568	.052
.235	.006	.661	.076
.275	.007	.739	.104
.314	.006	.801	.134
.353	.008	.840	.166
.392	.008	.840	.199
.431	.008	.832	.232
.471	.008	.813	.265
.510	.008	.786	.297
.549	.008	.755	.328
.588	.007	.727	.359
.627	.007	.700	.388
.667	.007	.669	.416
.706	.007	.638	.442
.745	.006	.615	.468
.784	.006	.587	.493
.824	.006	.560	.516
.863	.005	.533	.539
.902	.005	.506	.560
.941	.005	.475	.581
.980	.005	.451	.600
1.020	.004	.424	.618
1.059	.004	.401	.635
1.098	.004	.381	.651
1.137	.004	.362	.667
1.176	.004	.342	.681
1.216	.003	.327	.695
1.255	.003		.708
1.294	.003	.315	.721
1.333	.003	.299	.733
1.373	.003	.288	.744
1.412	.003	.272	.755
1.451	.003	.261	.766
1.490	.003	.249	.776
1.529	.003	.237	.785
1.569	.003	.226	.794
1.608	.003	.218	.803
1.647	.003	.206	.811
1.686	.003	.198	.819
1.725	.003	.191	.827
1.764	.003	.183	.834
1.804	.003	.175	.841
1.843	.003	.169	.848
1.882	.002	.163	.854
1.922	.002	.156	.861

THETA	E	ET	F
1.951	.001	.152	.857
2.000	.001	.144	.872
2.039	.001	.140	.878
2.078	.001	.136	.883
2.118	.001	.133	.889
2.157	.001	.124	.894
2.196	.001	.119	.898
2.235	.001	.113	.903
2.275	.001	.109	.907
2.314	.001	.103	.911
2.353	.001	.099	.915
2.392	.001	.093	.923
2.431	.001	.089	.926
2.471	.001	.086	.929
2.510	.001	.082	.933
2.549	.001	.074	.936
2.588	.001	.070	.938
2.627	.001	.066	.941
2.667	.001	.064	.944
2.706	.001	.062	.946
2.745	.001	.060	.948
2.784	.001	.058	.951
2.824	.001	.056	.953
2.863	.001	.054	.957
2.902	.001	.051	.959
2.941	.000	.050	.961
2.980	.000	.049	.963
2.998	.000	.047	.965
3.098	.000	.047	.966
3.137	.000	.045	.968
3.176	.000	.043	.970
3.215	.000	.042	.971
3.254	.000	.041	.973
3.293	.000	.040	.975
3.332	.000	.039	.976
3.371	.000	.038	.978
3.410	.000	.037	.979
3.449	.000	.035	.980
3.488	.000	.033	.982
3.527	.000	.031	.983
3.566	.000	.029	.984
3.605	.000	.027	.986
3.644	.000	.026	.987
3.683	.000	.025	.988
3.722	.000	.024	.989
3.761	.000	.023	.990
3.800	.000	.025	.991
3.839	.000	.023	.992

INPUT DATA

FOUR STAGE IN SERIES OPERATION
TEMPERATURE=15 DEG C
FLOW RATE= 4.00LIT/MIN.
VOLUME OF THE REACTOR= 408.0LIT.
THEORETICAL DETENTION TIME= 102.0GMIN
DT= 4.00MIN
THE AMOUNT OF DYE INJECTED= 0197LIT.
CONCENTRATION OF THE DYE ADDED= 238000.00PPM

DYE MEASUREMENTS

.50	7.50	15.00	26.00	42.00
57.00	69.00	77.00	83.00	86.50
88.50	89.50	90.00	90.00	89.50
88.50	87.00	85.00	83.00	80.00
76.50	73.00	69.00	66.00	62.50
59.00	56.00	53.00	50.00	48.00
46.00	43.50	41.50	39.50	37.50
36.00	34.50	32.50	30.50	29.00
27.50	26.50	25.00	24.00	22.75
21.75	21.00	19.90	18.50	18.00
17.00	16.50	15.75	14.75	14.00
13.00	12.50	11.50	11.00	10.50
10.00	9.75	9.50	9.00	8.50
8.25	8.00	7.50	7.40	7.00
6.90	6.50	6.30	6.00	5.60
5.50	5.20	4.90	4.50	4.50
4.40	4.35	4.25	4.10	4.00
3.90	3.75	3.60	3.50	3.50
3.45	3.30	3.20	3.10	3.00
2.60	2.50	2.25	2.10	2.00

TEST RESULTS AND CALCULATED VALUES

-94-

DYE RECOVERY = 98.48 PER CENT

MEAN DYE RESIDENCE BASED ON C=108.77 MIN
MEAN DYE RESIDENCE BASED ON E=108.77 MIN

THETA	E	ET	F
0.000	.000	.004	0.000
.039	.001	.066	.001
.078	.001	.133	.005
.118	.002	.230	.012
.157	.004	.371	.024
.196	.005	.504	.041
.235	.006	.610	.063
.275	.007	.680	.088
.314	.007	.733	.116
.353	.007	.764	.146
.392	.008	.782	.176
.431	.008	.791	.207
.471	.008	.795	.238
.510	.008	.795	.269
.549	.008	.791	.300
.588	.008	.782	.331
.627	.008	.769	.361
.667	.007	.751	.391
.706	.007	.733	.420
.745	.007	.707	.448
.784	.007	.676	.476
.824	.006	.645	.502
.863	.006	.610	.526
.902	.006	.558	.550
.941	.005	.552	.572
.980	.005	.521	.593
1.020	.005	.495	.613
1.059	.005	.468	.632
1.098	.004	.442	.649
1.137	.004	.424	.666
1.176	.004	.406	.683
1.216	.004	.384	.698
1.255	.004	.367	.713
1.294	.003	.349	.727
1.333	.003	.331	.740
1.373	.003	.318	.753
1.412	.003	.305	.765
1.451	.003	.287	.777
1.490	.003	.270	.788
1.529	.003	.256	.798
1.569	.002	.248	.808
1.608	.002	.234	.817
1.647	.002	.221	.826
1.686	.002	.212	.835
1.725	.002	.201	.843
1.765	.002	.192	.851
1.804	.002	.186	.858
1.843	.002	.176	.865
1.882	.002	.163	.872
1.922	.002	.159	.878

THETA	E	ET	F
1.961	.001	.150	.884
2.060	.001	.146	.890
2.039	.001	.139	.895
2.078	.001	.130	.901
2.118	.001	.124	.906
2.157	.001	.115	.910
2.196	.001	.110	.915
2.235	.001	.102	.919
2.275	.001	.097	.923
2.314	.001	.093	.927
2.353	.001	.088	.930
2.392	.001	.086	.934
2.431	.001	.084	.937
2.471	.001	.080	.940
2.510	.001	.075	.943
2.549	.001	.073	.946
2.588	.001	.071	.949
2.627	.001	.066	.952
2.667	.001	.065	.954
2.706	.001	.062	.957
2.745	.001	.061	.959
2.784	.001	.057	.961
2.824	.001	.056	.964
2.863	.001	.053	.966
2.902	.000	.049	.968
2.941	.000	.049	.970
2.980	.000	.046	.971
3.029	.000	.043	.975
3.059	.000	.040	.975
3.098	.000	.040	.976
3.137	.000	.039	.978
3.176	.000	.038	.979
3.216	.000	.038	.981
3.255	.000	.036	.982
3.294	.000	.035	.984
3.333	.000	.034	.985
3.373	.000	.033	.986
3.412	.000	.032	.988
3.451	.000	.031	.989
3.490	.000	.031	.990
3.529	.000	.030	.991
3.569	.000	.029	.993
3.608	.000	.028	.994
3.647	.000	.027	.995
3.686	.000	.027	.996
3.725	.000	.023	.997
3.765	.000	.022	.998
3.804	.000	.020	.999
3.843	.000	.019	.999
3.882	.000	.018	1.000

INPUT DATA.

FOUR STAGE IN SERIES OPERATION
TEMPERATURE=20 DEG.C
FLOW RATE= 4.00LIT/MIN
VOLUME OF THE REACTOR= 408.0LIT.
THEORETICAL DETENTION TIME= 102.20MIN
DT= 4.00MIN
THE AMOUNT OF DYE INJECTED= 0.236LIT.
CONCENTRATION OF THE DYE ADDED= 23800.00PPB

DYE MEASUREMENTS.

.10	6.00	17.50	27.50	40.00
52.50	61.00	70.00	79.00	85.00
89.50	92.50	94.00	94.00	93.50
92.00	90.50	88.50	86.50	84.00
81.00	77.50	75.00	71.50	68.50
65.50	62.50	60.00	57.50	55.00
52.00	49.00	47.00	44.00	42.00
40.50	39.00	37.00	35.00	34.00
32.50	31.00	29.50	28.00	27.00
26.00	24.50	23.50	22.50	21.50
20.50	20.00	19.00	18.00	17.00
16.00	15.50	14.50	14.00	13.00
12.75	12.00	11.50	11.00	10.50
10.00	9.50	9.00	8.50	8.25
8.00	7.50	7.25	7.00	6.75
6.50	6.40	6.25	6.10	6.00
5.75	5.50	5.40	5.25	5.10
5.05	5.00	4.90	4.80	4.50
4.50	4.25	4.00	3.90	3.75
3.50	3.50	3.40	3.25	3.00
3.00	2.90	2.75	2.60	2.50

TEST RESULTS AND CALCULATED VALUES

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DYE RECOVERY = 90.24 PER CENT

MEAN DYE RESIDENCE BASED ON C=117.18 MIN
MEAN DYE RESIDENCE BASED ON E=117.13 MIN

THETA	E	ET	F
0.000	.000	.001	.000
.039	.000	.049	.001
.078	.001	.142	.005
.117	.002	.223	.012
.157	.003	.324	.023
.196	.004	.425	.037
.235	.005	.494	.055
.274	.006	.567	.076
.313	.006	.640	.100
.352	.007	.688	.126
.391	.007	.725	.153
.431	.007	.749	.182
.470	.007	.761	.212
.509	.007	.761	.241
.548	.007	.757	.271
.587	.007	.745	.301
.626	.007	.733	.329
.665	.007	.717	.358
.705	.007	.701	.386
.744	.007	.680	.413
.783	.006	.656	.439
.822	.006	.628	.464
.861	.006	.607	.488
.900	.006	.579	.511
.939	.005	.555	.533
.978	.005	.531	.555
1.018	.005	.506	.575
1.057	.005	.486	.594
1.096	.005	.466	.613
1.135	.004	.445	.631
1.174	.004	.421	.648
1.213	.004	.397	.664
1.252	.004	.381	.679
1.292	.003	.356	.694
1.331	.003	.340	.707
1.370	.003	.328	.720
1.409	.003	.316	.733
1.448	.003	.300	.745
1.487	.003	.283	.756
1.526	.003	.275	.767
1.566	.003	.263	.778
1.605	.002	.251	.788
1.644	.002	.239	.797
1.683	.002	.227	.807
1.722	.002	.219	.815
1.761	.002	.211	.824
1.800	.002	.198	.832
1.840	.002	.190	.839
1.879	.002	.182	.847
1.918	.002	.174	.854

THETA	E	ET	F
1.957	.002	.166	.860
1.996	.002	.162	.867
2.035	.002	.154	.873
2.074	.001	.146	.879
2.114	.001	.138	.884
2.153	.001	.130	.889
2.192	.001	.126	.894
2.231	.001	.117	.899
2.270	.001	.113	.904
2.309	.001	.105	.908
2.348	.001	.103	.912
2.387	.001	.097	.916
2.427	.001	.093	.920
2.466	.001	.089	.923
2.505	.001	.085	.927
2.544	.001	.081	.930
2.583	.001	.077	.933
2.622	.001	.073	.936
2.661	.001	.069	.939
2.701	.001	.067	.941
2.740	.001	.065	.944
2.779	.001	.061	.949
2.818	.001	.059	.951
2.857	.001	.057	.953
2.896	.001	.055	.957
2.935	.001	.053	.959
2.975	.001	.051	.961
3.014	.000	.049	.963
3.053	.000	.047	.967
3.092	.000	.045	.969
3.131	.000	.044	.970
3.170	.000	.043	.972
3.209	.000	.041	.974
3.249	.000	.041	.975
3.288	.000	.040	.977
3.327	.000	.039	.978
3.366	.000	.036	.980
3.405	.000	.034	.981
3.444	.000	.033	.983
3.483	.000	.032	.984
3.523	.000	.032	.985
3.562	.000	.032	.986
3.601	.000	.032	.987
3.640	.000	.032	.989
3.679	.000	.032	.990
3.718	.000	.032	.991
3.757	.000	.024	.992
3.796	.000		
3.836	.000		
3.875	.000		

INPUT DATA

FOUR STAGE IN SERIES OPERATION
TEMPERATURE=25 DEG.C
FLOW RATE= 4.00LIT./MIN.
VOLUME OF THE REACTOR= 408.0LIT.
THEORETICAL DETENTION TIME= 102.00MIN
DT= 4.00MIN
THE AMOUNT OF DYE INJECTED= 0.250LIT
CONCENTRATION OF THE DYE ADDED= 23800.00PPB

DYE MEASUREMENTS

2.50	5.75	10.50	18.00	28.00
39.50	50.50	60.00	69.50	77.50
84.50	91.00	96.90	100.00	103.00
104.50	105.25	105.50	105.40	104.50
103.50	101.50	99.00	96.00	93.50
90.00	87.00	84.00	80.50	77.50
74.25	71.00	68.00	65.25	62.50
60.00	57.00	54.00	51.00	48.50
46.25	43.75	41.50	39.50	37.50
35.25	33.50	31.75	30.00	28.50
26.90	25.50	24.00	22.75	21.50
20.25	19.00	17.75	16.50	15.50
14.50	13.50	12.75	12.00	11.00
10.50	9.75	9.00	8.50	8.00
7.50	7.00	6.50	6.00	5.75
5.50	5.25	4.90	4.50	4.40
4.10	3.90	3.70	3.50	3.35
3.10	3.00	2.90	2.60	2.50
2.49	2.48	2.45	2.10	2.00
1.75	1.60	1.40	1.25	1.00

TEST RESULTS AND CALCULATED VALUES

DYE RECOVERY = 98.95 PER CENT

MEAN DYE RESIDENCE BASED ON C=116.19 MIN
MEAN DYE RESIDENCE BASED ON E=116.19 MIN

THETA	E	ET	F
0.000	.000	.017	0.000
.039	.000	.040	.001
.078	.001	.073	.003
.118	.001	.125	.007
.157	.002	.194	.013
.196	.003	.274	.023
.235	.003	.350	.035
.275	.004	.416	.050
.314	.005	.482	.067
.353	.005	.537	.087
.392	.006	.586	.109
.431	.006	.631	.133
.471	.007	.672	.159
.510	.007	.693	.186
.549	.007	.714	.213
.588	.007	.724	.241
.627	.007	.729	.270
.667	.007	.731	.299
.706	.007	.730	.327
.745	.007	.724	.356
.784	.007	.717	.384
.824	.007	.703	.412
.863	.007	.686	.439
.902	.007	.665	.466
.941	.006	.648	.491
.980	.006	.624	.516
1.020	.006	.603	.540
1.059	.006	.582	.564
1.098	.005	.558	.586
1.137	.005	.537	.607
1.176	.005	.515	.628
1.216	.005	.492	.648
1.255	.005	.471	.667
1.294	.004	.452	.685
1.333	.004	.433	.702
1.373	.004	.416	.719
1.412	.004	.395	.735
1.451	.004	.374	.750
1.490	.003	.353	.764
1.529	.003	.336	.777
1.569	.003	.321	.790
1.608	.003	.303	.803
1.647	.003	.288	.814
1.686	.003	.274	.825
1.725	.003	.260	.836
1.765	.003	.244	.846
1.804	.002	.232	.855
1.843	.002	.220	.864
1.882	.002	.208	.872
1.922	.002	.198	.880

THETA	E	ET	F
1.961	.002	.186	.888
.000	.002	.177	.895
.039	.002	.166	.901
.078	.002	.158	.908
.118	.001	.149	.914
.157	.001	.140	.919
.196	.001	.132	.925
.235	.001	.123	.930
.275	.001	.114	.934
.314	.001	.107	.939
.353	.001	.100	.943
.392	.001	.094	.947
2.431	.001	.068	.950
.471	.001	.083	.954
.510	.001	.073	.957
.549	.001	.068	.960
.588	.001	.066	.967
.627	.001	.055	.970
.667	.001	.055	.972
.706	.001	.045	.974
.745	.001	.042	.975
.784	.000	.038	.977
.824	.000	.036	.979
.863	.000	.034	.980
.902	.000	.031	.982
.941	.000	.030	.983
.979	.000	.027	.984
.999	.000	.024	.986
1.37	.000	.021	.987
.176	.000	.018	.988
.216	.000	.017	.990
.255	.000	.017	.991
.294	.000	.016	.992
.333	.000	.015	.993
.412	.000	.015	.994
.451	.000	.015	.995
.490	.000	.017	.995
.529	.000	.017	.996
.569	.000	.017	.997
.608	.000	.017	.997
.647	.000	.014	.998
.686	.000	.012	.998
.725	.000	.011	.999
.765	.000	.010	.999
.804	.000	.009	1.000
.843	.000	.007	1.000
3.882	.000		

INPUT DATA

FOUR STAGE IN SERIES OPERATION AND ALL BIOGAS WAS STRIPPED OFF
TEMPERATURE=15 DEG. C
FLOW RATE= 4.0LIT./MIN.
VOLUME OF THE REACTOR= 408.0LIT.
THEORETICAL DETENTION TIME= 102.20MIN
DT= 5.0MIN
THE AMOUNT OF DYE INJECTED= .0200LIT.
CONCENTRATION OF THE DYE ADDED=2380000.00PPB

DYE MEASUREMENTS

0.00	.05	2.00	7.00	12.50
22.50	28.00	35.00	41.50	47.50
54.00	60.00	64.50	70.00	75.00
77.00	77.50	78.00	78.00	77.50
76.50	75.00	72.00	69.00	67.00
64.00	61.50	59.00	56.00	53.50
51.00	48.00	46.00	43.00	40.00
37.50	35.00	32.50	30.00	28.00
26.00	24.00	22.50	20.50	19.00
17.50	16.00	14.50	13.50	12.50
11.50	10.50	9.50	8.50	7.50
7.00	6.50	6.00	5.00	4.50
4.00	3.50	3.50	3.50	3.50
3.00	3.00	2.50	2.50	2.00
2.00	2.00	2.00	2.00	1.50
1.50	1.50	1.50	1.50	1.50
1.50	1.00	1.00	1.00	1.00
1.00				

TEST RESULTS AND CALCULATED VALUES

-103-

DYE RECOVERY = 95.32 PER CENT

MEAN DYE RESIDENCE BASED ON C=130.54MTN
MEAN DYE RESIDENCE BASED ON E=130.54MIN

THETA	E	ET	F
0.000	0.000	0.000	0.000
.049	.000	.000	.000
.098	.000	.018	.000
.147	.001	.063	.002
.196	.001	.113	.007
.245	.002	.203	.014
.294	.002	.252	.026
.342	.003	.315	.039
.391	.004	.374	.056
.440	.004	.428	.076
.489	.005	.487	.098
.538	.005	.541	.123
.587	.006	.581	.151
.636	.006	.631	.181
.685	.007	.676	.212
.734	.007	.694	.246
.783	.007	.698	.280
.832	.007	.703	.314
.881	.007	.703	.349
.930	.007	.698	.383
.978	.007	.689	.417
1.027	.007	.676	.450
1.076	.006	.649	.483
1.125	.006	.622	.514
1.174	.006	.604	.544
1.223	.006	.577	.573
1.272	.006	.554	.600
1.321	.005	.532	.627
1.370	.005	.505	.652
1.419	.005	.482	.676
1.468	.004	.460	.699
1.517	.004	.432	.721
1.566	.004	.414	.742
1.614	.004	.387	.762
1.663	.003	.360	.780
1.712	.003	.338	.797
1.761	.003	.315	.813
1.810	.003	.293	.828
1.859	.003	.270	.842
1.908	.002	.252	.854
1.957	.002	.234	.866
2.006	.002	.216	.877
2.055	.002	.203	.887
2.104	.002	.185	.897
2.153	.002	.171	.906
2.202	.002	.158	.914
2.250	.001	.144	.921
2.299	.001	.131	.928
2.348	.001	.122	.934
2.397	.001	.113	.940

THETA	E	ET	F
2.446	0.001	104	945
2.495	0.001	095	950
2.544	0.001	086	954
2.593	0.001	077	958
2.642	0.001	068	962
2.691	0.001	059	965
2.740	0.001	045	968
2.789	0.001	041	971
2.838	0.000	036	973
2.886	0.000	032	975
2.935	0.000	032	979
2.984	0.000	032	980
2.992	0.000	027	982
2.993	0.000	027	983
2.994	0.000	023	985
2.995	0.000	018	986
2.996	0.000	018	987
2.997	0.000	018	988
2.998	0.000	014	989
2.999	0.000	014	990
3.000	0.000	014	991
3.001	0.000	014	992
3.002	0.000	014	993
3.003	0.000	014	994
3.004	0.000	014	995
3.005	0.000	014	996
3.006	0.000	014	997
3.007	0.000	014	998
3.008	0.000	009	999
3.009	0.000	009	000
3.010	0.000	009	000
3.011	0.000	009	000

INPUT DATA

SINGLE STAGE OPERATION
TEMPERATURE=15 DEG. C
FLOW RATE= 2.50LIT./MIN.
VOLUME OF THE REACTOR= 100.0LIT.
THEORETICAL DETENTION TIME= 16.00MIN
DT= 2.00MIN
THE AMOUNT OF DYE INJECTED= 0.009ALIT.
CONCENTRATION OF THE DYE ADDED= 238000.00PPM

DYE MEASUREMENTS

540.00	295.00	225.00	195.00	175.00
158.00	142.00	130.00	120.00	112.00
105.00	98.00	94.00	88.00	84.00
80.00	76.00	73.00	69.00	66.00
64.00	62.00	60.00	60.00	58.00
46.50	44.50	43.00	41.00	39.50
38.50	37.00	35.50	34.00	32.50
31.50	30.00	28.50	27.00	26.00
24.50	23.50	22.30	21.50	20.50
19.50	18.75	18.00	17.30	16.70
16.00	15.30	14.90	14.15	14.00
13.40	13.00	12.80	12.30	12.00
11.70	11.20	10.90	10.60	10.20
10.00	9.80	9.60	9.30	9.00
8.70	8.40	8.20	8.00	7.80
7.60	7.30	7.00	6.85	6.40
6.20	5.90	5.60	5.40	5.20
5.00	4.80	4.60	4.40	4.20
4.00	4.00	4.00	4.00	4.00
4.00	4.00	4.00	4.00	4.00
4.00	4.00	4.00	4.00	4.00
3.60	3.60	3.60	3.60	3.60
3.30	3.20	3.10	2.90	2.70
2.60	2.45	2.30	2.10	1.95
1.75	1.60	1.50	1.35	1.25
1.75	1.60	1.50	1.35	1.25

TEST RESULTS AND CALCULATED VALUES

-106-

DYE RECOVERY = 97.65 PER CENT

MEAN DYE RESIDENCE BASED ON C = 43.75 MIN
MEAN DYE RESIDENCE BASED ON E = 43.75 MIN

THETA	E	ET	F
0.000	.065	1.033	0.000
.125	.035	.564	.100
.250	.027	.430	.162
.375	.023	.373	.212
.500	.021	.335	.256
.625	.019	.302	.296
.750	.017	.272	.332
.875	.016	.249	.368
1.000	.014	.229	.394
1.125	.013	.214	.422
1.250	.013	.201	.448
1.375	.012	.187	.472
1.500	.011	.180	.495
1.625	.011	.168	.517
1.750	.010	.161	.538
1.875	.010	.153	.557
2.000	.009	.145	.576
2.125	.009	.140	.594
2.250	.008	.132	.611
2.375	.008	.126	.627
2.500	.008	.122	.642
2.625	.007	.119	.657
2.750	.007	.115	.672
2.875	.007	.115	.686
3.000	.007	.111	.700
3.125	.006	.089	.713
3.250	.005	.085	.724
3.375	.005	.082	.734
3.500	.005	.078	.744
3.625	.005	.076	.754
3.750	.005	.074	.763
3.875	.004	.071	.772
4.000	.004	.068	.781
4.125	.004	.065	.789
4.250	.004	.062	.797
4.375	.004	.060	.805
4.500	.004	.057	.812
4.625	.003	.055	.819
4.750	.003	.052	.826
4.875	.003	.050	.832
5.000	.003	.047	.838
5.125	.003	.045	.844
5.250	.003	.043	.849
5.375	.003	.041	.855
5.500	.003	.039	.860
5.625	.002	.037	.864
5.750	.002	.036	.869
5.875	.002	.034	.873
6.000	.002	.033	.878
6.125	.002	.032	.882

INPUT DATA

SINGLE STAGE OPERATION
TEMPERATURE = 16 DEG. C
FLOW RATE = 4.0 LIT./MIN.
VOLUME OF THE REACTOR = 40.0 LIT.
THEORETICAL DETENTION TIME = 10.00 MIN
DT = 1.00 MIN
THE AMOUNT OF DYE INJECTED = 0.0100 LIT.
CONCENTRATION OF THE DYE ADDED = 23800.00 PPM

DYE MEASUREMENTS

570.00	376.00	296.00	254.00	198.00
195.00	175.00	160.00	150.00	142.00
130.00	122.00	112.00	105.00	100.00
96.00	92.00	88.00	84.00	79.00
74.00	73.00	71.00	68.00	65.00
63.00	61.00	59.00	58.00	54.00
53.00	52.00	51.00	48.00	41.00
45.00	44.00	43.00	41.00	40.00
39.00	37.00	36.00	35.00	33.00
32.00	31.00	30.00	29.00	28.00
27.00	26.00	25.00	24.00	23.00
22.50	21.80	21.00	20.30	20.00
19.00	18.60	18.00	17.60	17.00
16.70	16.00	15.90	15.30	15.00
14.70	14.50	14.00	13.80	13.50
13.30	12.90	12.60	12.30	12.00
11.80	11.60	11.20	11.00	10.70
10.50	10.30	10.00	9.80	9.60
9.40	9.20	8.50	8.30	7.60
7.50	7.40	7.20	7.20	7.20
6.80	6.60	6.40	6.20	6.30
6.00	5.90	5.70	5.60	5.50
5.40	5.20	5.30	5.30	5.10
5.00	4.50	3.90	3.90	3.90
3.90	3.90	3.80	3.70	3.60
3.50	3.50	3.50	3.50	3.40
3.00	3.00	2.50	2.50	2.60
2.00	2.00	2.00	2.00	2.00

TEST RESULTS AND CALCULATED VALUES

DYE RECOVERY = 93.54 PER CENT

MEAN DYE RESIDENCE BASED ON C = 25.74 MIN
MEAN DYE RESIDENCE BASED ON E = 25.74 MIN

THETA	E	ET	F
0.000	.102	1.024	0.000
.100	.068	.676	.0885
.200	.053	.532	.145
.300	.046	.456	.195
.400	.036	.356	.235
.500	.035	.350	.271
.600	.031	.314	.304
.700	.029	.287	.334
.800	.027	.270	.362
.900	.026	.255	.389
1.000	.023	.234	.413
1.100	.022	.219	.435
1.200	.020	.201	.456
1.300	.019	.189	.476
1.400	.018	.172	.494
1.500	.017	.165	.512
1.600	.017	.158	.529
1.700	.016	.151	.545
1.800	.015	.142	.560
1.900	.014	.133	.575
2.000	.013	.121	.589
2.100	.013	.128	.602
2.200	.012	.122	.615
2.300	.012	.117	.627
2.400	.011	.113	.639
2.500	.011	.110	.651
2.600	.011	.106	.662
2.700	.011	.104	.673
2.800	.010	.097	.683
2.900	.010	.095	.693
3.000	.009	.093	.703
3.100	.009	.092	.712
3.200	.009	.086	.721
3.300	.008	.084	.730
3.400	.008	.081	.739
3.500	.008	.079	.747
3.600	.008	.077	.755
3.700	.008	.074	.763
3.800	.007	.074	.771
3.900	.007	.070	.778
4.000	.007	.066	.785
4.100	.007	.065	.792
4.200	.006	.065	.798
4.300	.006	.063	.805
4.400	.006	.059	.811
4.500	.006	.057	.817
4.600	.006	.056	.822
4.700	.005	.054	.828
4.800	.005	.052	.833
4.900	.005	.050	.838

THETA	E	ET	F
5.000	.005	.049	.843
5.100	.005	.047	.848
5.200	.004	.045	.852
5.300	.004	.043	.857
5.400	.004	.041	.861
5.500	.004	.040	.865
5.600	.004	.039	.869
5.700	.004	.038	.873
5.800	.004	.036	.877
5.900	.004	.034	.880
6.000	.004	.033	.884
6.100	.004	.032	.887
6.200	.003	.031	.890
6.300	.003	.030	.894
6.400	.003	.029	.897
6.500	.003	.028	.900
6.600	.003	.027	.903
6.700	.003	.026	.906
6.800	.003	.025	.911
6.900	.003	.024	.914
7.000	.003	.023	.916
7.100	.003	.022	.922
7.200	.003	.021	.924
7.300	.003	.020	.926
7.400	.003	.019	.929
7.500	.003	.018	.931
7.600	.003	.017	.933
7.700	.003	.016	.935
7.800	.003	.015	.938
8.000	.002	.014	.940
8.100	.002	.013	.942
8.200	.002	.013	.944
8.300	.002	.013	.946
8.400	.002	.013	.948
8.500	.002	.013	.949
8.600	.002	.013	.951
8.700	.002	.013	.953
8.800	.002	.013	.955
8.900	.002	.013	.956
9.000	.002	.013	.958
9.100	.002	.013	.960
9.200	.002	.013	.961
9.300	.001	.013	.963
9.400	.001	.013	.964
9.500	.001	.013	.965
9.600	.001	.013	.967
9.700	.001	.013	.968
9.800	.001	.013	.969
9.900	.001	.013	

INPUT DATA

SINGLE STAGE OPERATION AND ALL BIOMASS STRIPPED
TEMPERATURE=15 DEG. C
FLOW RATE= 4.00LIT./MIN.
VOLUME OF THE REACTOR= 100.0LIT.
THEORETICAL DETENTION TIME= 25.00 MIN
DT= 1.00 MIN
THE AMOUNT OF DYE INJECTED= .0100LIT.
CONCENTRATION OF THE DYE ADDED=2380000.COPPB

DYE MEASUREMENTS

498.00	264.00	242.00	214.00	200.00
190.00	180.00	170.00	162.00	154.00
146.00	140.00	132.00	125.00	120.00
114.00	108.00	102.00	99.00	95.00
88.00	84.00	80.00	76.00	74.00
70.00	67.00	65.00	63.00	61.00
58.00	57.00	55.00	53.00	51.00
49.00	48.00	47.00	45.00	44.00
43.00	42.00	40.00	39.00	38.00
37.00	35.00	34.00	33.00	32.00
32.00	31.00	30.00	29.00	28.00
28.00	27.00	26.00	26.00	25.00
24.00	24.00	23.00	22.00	22.00
22.00	21.00	20.00	20.00	19.00
19.00	18.00	18.00	17.00	16.00
16.00	16.00	15.00	14.00	14.00
14.00	15.00	13.00	12.00	12.00
12.00	11.00	11.00	10.00	10.00
10.00	10.00	10.00	9.00	9.00
9.00	8.00	8.00	8.00	8.00
8.00	7.80	7.70	7.60	7.50
7.30	7.10	7.00	6.90	6.70
6.60	6.50	6.30	6.20	6.00
5.90	5.80	5.60	5.50	5.30
5.20	5.00	4.90	4.80	4.60
4.50	4.40	4.20	4.00	3.94
3.80	3.70	3.50	3.40	3.20
3.10	2.90	2.80	2.70	2.50
2.40	2.20	2.10	2.00	1.80
1.70	1.60	1.40	1.30	1.10

TEST RESULTS AND CALCULATED VALUES

-111-

DYE RECOVERY = 99.37 PER CENT

MEAN DYE RESIDENCE BASED ON C = 28.57 MIN
MEAN DYE RESIDENCE BASED ON E = 28.57 MIN

THETA	E	ET	F
0.000	.084	2.106	0.000
.040	.045	1.116	.064
.080	.041	1.023	.107
.120	.036	.955	.146
.160	.034	.846	.181
.200	.032	.803	.214
.240	.030	.761	.245
.280	.029	.719	.275
.320	.027	.685	.303
.360	.026	.651	.329
.400	.025	.617	.355
.440	.024	.592	.379
.480	.022	.558	.402
.520	.021	.529	.424
.560	.020	.507	.444
.600	.019	.482	.464
.640	.018	.457	.483
.680	.017	.431	.501
.720	.017	.419	.518
.760	.016	.402	.534
.800	.015	.372	.550
.840	.014	.355	.564
.880	.014	.338	.578
.920	.013	.321	.591
.960	.013	.313	.604
1.000	.012	.295	.616
1.040	.011	.283	.628
1.080	.011	.275	.639
1.120	.011	.266	.650
1.160	.010	.245	.660
1.200	.010	.241	.670
1.240	.009	.233	.680
1.280	.009	.224	.689
1.320	.009	.216	.699
1.360	.008	.207	.707
1.400	.008	.203	.716
1.440	.008	.199	.724
1.480	.008	.190	.732
1.520	.007	.186	.740
1.560	.007	.182	.747
1.600	.007	.178	.755
1.640	.007	.169	.762
1.680	.007	.165	.769
1.720	.007	.161	.776
1.760	.006	.156	.782
1.800	.006	.148	.788
1.840	.006	.144	.795
1.880	.006	.140	.800
1.920	.006	.135	.806
1.960	.005		.812

THETA

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P

THETA	E	ET	F
4.000	•001	•034	•962
4.040	•001	•033	•964
4.080	•001	•033	•965
4.120	•001	•032	•965
4.160	•001	•031	•967
4.200	•001	•030	•969
4.240	•001	•030	•970
4.280	•001	•029	•971
4.320	•001	•028	•972
4.360	•001	•027	•973
4.400	•001	•027	•975
4.440	•001	•026	•976
4.480	•001	•025	•977
4.520	•001	•025	•978
4.560	•001	•024	•979
4.600	•001	•023	•980
4.640	•001	•022	•981
4.680	•001	•021	•982
4.720	•001	•021	•983
4.760	•001	•020	•984
4.800	•001	•019	•985
4.840	•001	•019	•986
4.880	•001	•018	•987
4.920	•001	•017	•988
4.960	•001	•016	•989
5.000	•001	•015	•990
5.040	•001	•014	•991
5.080	•001	•013	•991
5.120	•001	•012	•993
5.160	•001	•011	•994
5.200	•001	•011	•994
5.240	•001	•010	•995
5.280	•001	•010	•996
5.320	•001	•010	•996
5.360	•001	•010	•997
5.400	•001	•010	•997
5.440	•000	•010	•998
5.480	•000	•010	•998
5.520	•000	•009	•998
5.560	•000	•009	•999
5.600	•000	•008	•999
5.640	•000	•008	•999
5.680	•000	•007	•999
5.720	•000	•007	•999
5.760	•000	•006	•999
5.800	•000	•006	•999
5.840	•000	•005	•999
5.880	•000	•005	•999
5.920	•000	•005	•999
5.960	•000	•005	1.000
			1.000
			1.000

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FTN 4.5+410

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PAGE 1

PROGRAM FLBTS (INPUT,TAPE5=INPUT,OUTPUT,TAPE6=OUTPUT)

PROGRAM TO CALCULATE THE MAGNITUDE OF FLOW BETWEEN TWO CSTRS

C
C DEFINITIONS
C CA(N)=THE MEASURED DYE CONCENTRATION OF OUTER CSTR,AT TIME =N*dt
C AFTER PULSE DYE INJECTION,IN PPB
C VT=HYDRAULIC LOADING,IN LIT./MIN.
C VA=VOLUME OF THE OUTER CSTR,IN LIT.
C CTR(N)=CALCULATED DYE CONCENTRATION
C AFTER PULSE DYE INJECTION,IN PPB
C DT=THE SAMPLING INTERVAL,IN MIN.
C AB=THE CALCULATED FLOW RATE BETWEEN TWO CSTRS ARE CALCULATED N TIMES.
C DE=THE THEORETICAL DETENTION TIME OF OUTER CSTR IF THERE WERE
C NO INNER CSTR,OUTER CSTR DYE CONCENTRATIONS INSERTED

DIMENSION CAV(151),DYELO(151),DYINDI(151),CBI(151)

DIMENSION CBAY(151),SCBAV(151),ABFLR(151),THETA(151),SCAV(151)

DIMENSION DYIN(151),CAV(151),TIME(151)

READ(5,388) DYIN,DYCON,DET,L

388 FORMAT(F10.4,DE10.2,F10.2,I10)

READ(5,1) DT,VIN,VA,VBTOTA

1 FORMAT(F10.2,I10,I3F10.2)

2 READ(5,2){CA(I),I=1,L}

FORMAT(F5.1)

51 WRITE(6,997)////////////////

997 FORMAT(20X,1H INPUT DATA,//)

995 FORMAT(10X,1H INPUT DATA,//)

995 WRITE(6,101) #SINGLE STAGE OPERATION#,10X)

111 FORMAT(10X,VT#TEMPERATURE = 15 DEG. C#,10X)

306 FORMAT(10X,VT#FLOW RATE =#,F6.2,#LIT./MIN.#,10X)

305 FORMAT(10X,VT#VOLUME OF THE REACTOR =#,F7.1,#LIT.#,10X)

158 FORMAT(10X,VT#VOLUME OF THE OUTER CSTR =#,F6.2,#LIT.#,10X)

159 FORMAT(10X,VT#VOLUME OF THE INNER CSTR =#,F6.2,#LIT.#,10X)

102 FORMAT(6,102) DT =#,F5.2,#MIN#,10X)

308 FORMAT(10X,VT#THE AMOUNT OF DYE INJECTEN =#,F7.4,#LIT.#,10X)

3.09 FORMAT(6,308) DYCON

FORMAT(10X,VT#CONCENTRATION OF THE DYE ADDED =#,F10.2,#PPB#,//)

C
C CALCULATION OF THE TOTAL AMOUNT OF DYE WHICH EXIST IN OUTER CSTR
C DO 3 I=1,L
C CAV(I)=CA(I)*VA
C 3 CONCINUTATION OF THE AVERAGE DYE CONCENTRATION IN OUTER CSTR

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TS

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M=L-1
DO 4 I=1,M
K=I+1
CAV(I)=0.5*(CA(I)+CA(K))
CONTINUE

C CALCULATION OF THE AMOUNT OF DYE WHICH HAS LEFT THE REACTOR
SDYEL0(1)=0,0

DC 5 I=2,L
K=I-1
DYELC(I)=CAV(K)*DT*VT
SDYEL0(I)=SDYEL0(K)+DYEL0(I)

5 CONTINUE

C CALCULATION OF THE TOTAL AMOUNT OF DYE CONTAINED IN INNER CSTR.
DYINU(I)=0.0

DO 6 I=2,L
DYINTD(I)=CAV(I)-CAV(1)-SDYEL0(I)

6 CONTINUE

CBT(I)=DYINTD(I)/VB
7 CONTINUE

C CALCULATION OF THE AVERAGE DYE CONCENTRATION IN INNER CSTR.
DO 8 I=2,L
K=I-1
CAV(K)=(CBT(I)+CBT(K))*0.5
CONTINUE

C CALCULATION OF THE SUMS OF AVERAGE DYE CONCENTRATIONS
SCBAV(I)=CAV(I)
SCAV(I)=CAV(1)

DO 9 I=2,M
K=I-1
SCBAV(I)=SCBAV(K)+CAV(I)
SCAV(I)=SCAV(K)+CAV(1)

9 CONTINUE

C CALCULATION OF THE FLOW RATE BETWEEN INNER AND OUTER CSTRS
IK=N-1
DO 21 I=1,IK
K=I+1
ARFLR(I)=DYINDI(K)/(DT*(SCAV(I)-SCBAV(I)))
21 CONTINUE

C CALCULATION OF TIME VALUES
TIME(1)=0.0
DO 20 I=2,L
KL=I-1
TIME(I)=TIME(KL)+DT

20 TIME(I)=TIME(I)+DT

C CALCULATION OF THE THETA VALUES
THETA(1)=0.0
DO 10 I=2,L
THETA(I)=TIME(I)/OE!

10 THETA(I)=TIME(I)/OE!

LBTCS 73/73 TS /7

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C PRINT OUT STATEMENTS

```
      WRITE(6,959)
959   FORMAT(2'0X',#CALCULATED VALUES#,10X,///)
      WRITE(6,14) SHTHETA,6X,SHABFLR,4X,3HCRT,9X,#CA#,///)
14    FORMAT(10X,5HSHETA,6X,5HABFLR,4X,3HCRT,9X,#CA#,//)
      DO 55 I=1,L
      IF(I.EQ.10) GO TO 99
      IF(I.EQ.28) GO TO 99
      IF(I.EQ.46) GO TO 99
      IF(I.EQ.62) GO TO 99
      IF(I.EQ.82) GO TO 99
      IF(I.EQ.100) GO TO 99
      IF(I.EQ.140) GO TO 99
      GO TO 333
      WRITE(6,97) /////
97    FORMAT(6,13)
      WRITE(6,13) SHTHETA(1),SHABFLR(4X,3HCRT,9X,#CA#,//)
113   FORMAT(6,11)
113   WRITE(6,11) (THETA(I),CBT(I),CA(I))
      WRITE(6,2,7X,6X#FF6.2,7X,F2.2,/)
11    FORMAT(10X,F6.2,7X,6X#FF6.2,7X,F2.2,/)
      IF(I.GT.N) GO TO 54
      WRITE(6,19) (ABFLR(I))
19    FORMAT(10X,9X,F6.2,7X)
      GO TO 55
      WRITE(6,56)
54    FORMAT(10X,/)
      CONTINUE
55    END
```

--ENTER Y POINTS--
3B INPUT 43B OUTPUT 38 TAPESE

--EXTEND NAL S--
INPCI. INPCR. OUTCI. Q8NTRY.

--STATEMENT LABELS--
3638 .2 F 3668 .3 ID 0B
08 .6 ID 0B 0B
08 .10 ID 0B 0B
5148 .29 ID 0B 510B
3518 .55 D 353B .11 1D
3258 .159 D 445B .56 FF
4278 .309 FF 436B .11 1D
4218 .995 F 460B .305 F
4618 .333 F 402B .997 F

--ID--
0B .8 ID 0B
0B .14 ID 0B
4748 FFFFFF
370B 501B
503B 414B
414B 357B
357B .306 F
.306 F
.333 F
.997 F

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INPUT DATA

SINGLE STAGE OPERATION
TEMPERATURE = 15 DEG. C
FLOW RATE = 4.00 LIT./MIN.
VOLUME OF THE REACTOR = 100.0 LIT.
VOLUME OF THE OUTER CSTR = 40.00 LIT.
VOLUME OF THE INNER CSTR = 60.00 LIT.
DT = 1.00 MIN
THE AMOUNT OF DYE INJECTED = 0.010 LIT.
CONCENTRATION OF THE DYE ADDED = 23800.00 PPB

CALCULATED VALUES

THETA	ABFLR	CBT	CA
0.00		0.00	570.00
.10	13.84	97.80	376.00
.20	11.94	128.73	296.00
.30	10.53	138.40	254.00
.40	11.15	160.67	198.00
.50	9.90	149.57	195.00
.60	9.60	150.57	175.00
.70	9.35	149.40	160.00
.80	9.05	145.73	150.00
	8.76		

THETA	ABFLR	CBT	CA
.90		141.33	142.00
1.00	8.73	140.27	130.00
1.10	8.65	137.20	122.00
1.20	8.77	136.07	112.00
1.30	8.85	133.50	105.00
1.40	8.91	130.00	100.00
1.50	8.95	126.13	96.00
1.60	9.02	122.53	92.00
1.70	9.12	119.20	88.00
1.80	9.26	116.13	84.00
1.90	9.51	114.03	79.00
2.00	9.87	112.27	74.00
2.10	10.03	108.03	73.00
2.20	10.26	104.57	71.00
2.30	10.58	101.93	68.00
2.40	10.98	99.50	65.00
2.50	11.37	96.57	63.00
2.60	11.81	93.77	61.00
	12.31		

?

THETA	ABFLR	CBT	C4
2.70		91.14°	59.00
2.80	12.76	87.87	58.00
2.90	13.64	86.80	54.00
3.00	-I	83.90	53.00
3.10		81.07	52.00
3.20		78.30	51.00
3.30		77.00	48.00
3.40		74.50	47.00
3.50		72.77	45.00
3.60		70.47	44.00
3.70		68.23	43.00
3.80		66.77	41.00
3.90		64.73	40.00
4.00		62.77	39.00
4.10		61.57	37.00
4.20		59.80	36.00
4.30		58.10	35.00
4.40		57.17	33.00

THETA	ABFLR	CBT	CA
4.50		55.67	32.00
4.60		54.23	31.00
4.70		52.87	30.00
4.80		51.57	29.00
4.90		50.33	28.00
5.00		49.17	27.00
5.10		48.07	26.00
5.20		47.03	25.00
5.30		46.07	24.00
5.40		45.17	23.00
5.50		43.98	22.50
5.60		42.97	21.80
5.70		42.08	21.00
5.80		41.17	20.30
5.90		40.03	20.00
6.00		39.39	19.00

INPUT DATA

SINGLE STAGE OPERATION
TEMPERATURE = 15 DEG. C
FLOW RATE = 2.50 LIT./MIN.
VOLUME OF THE REACTOR = 100.0LIT.
VOLUME OF THE OUTER CSTR = 40.00LIT.
VOLUME OF THE INNER CSTR = 60.00LIT.
DT = 2.00MIN
THE AMOUNT OF DYE INJECTED = .0090LIT.
CONCENTRATION OF THE DYE ADDED = 23800.00PPB

CALCULATED VALUES

THETA	ABFLR	CBT	CA
0.00		0.00	540.00
.13	10.92	128.54	295.00
.25	9.76	153.54	225.00
.38	8.88	156.04	195.00
.50	8.29	153.96	175.00
.63	7.95	151.42	158.00
.75	7.86	149.58	142.00
.88	7.85	146.25	130.00
1.00	7.93	142.50	120.00
	8.05		

THETA	ABFLR	CBT	CA
1.13		138.17	112.00
1.25	8.23	133.79	105.00
1.38	8.53	130.00	98.00
1.50	8.78	124.67	94.00
1.63	9.22	121.08	88.00
1.75	9.68	116.58	84.00
1.88	10.26	112.42	80.00
2.00	-I	108.58	76.00
2.13		104.38	73.00
2.25		101.13	69.00
2.38		97.50	66.00
2.50		93.42	64.00
2.63		89.50	62.00
2.75		85.75	60.00
2.88		80.75	60.00
3.00		77.17	58.00
3.13		80.48	46.50
3.25		78.02	44.50

THETA	ABFLR	CBT	CA
3.38		75.38	43.00
3.50		73.21	41.00
3.63		70.85	39.50
3.75		68.27	38.50
3.88		66.13	37.00
4.00		64.10	35.50
4.13		62.21	34.00
4.25		60.44	32.50
4.38		58.44	31.50
4.50		56.88	30.00
4.63		55.44	28.50
4.75		54.13	27.00
4.88		52.58	26.00
5.00		51.48	24.50
5.13		50.15	23.50
5.25		49.04	22.30
5.38		47.75	21.50
5.50		46.66	20.50

THETA	ABFLR	CBT	CA
5.63		45.66	19.50
5.75		44.57	18.75
5.88		43.54	18.00
6.00		42.53	17.30
6.13		41.52	16.70
6.25		40.62	16.00
6.38		39.78	15.30
6.50		38.79	14.90
6.63		38.08	14.15
6.75		37.01	14.00
6.88		36.27	13.40
7.00		35.43	13.00
7.13		34.49	12.80
7.25		33.78	12.30
7.38		32.97	12.00
7.50		32.18	11.70

THETA	ABFLR	CBT	CA	-125-
7.63		31.56	11.20	
7.75		30.84	10.90	
7.88		30.14	10.60	
8.00		29.54	10.20	
8.13		28.83	10.00	
8.25		28.14	9.80	
8.38		27.47	9.60	
8.50		26.88	9.30	
8.63		26.32	9.00	
8.75		25.78	8.70	
8.88		25.27	8.40	
9.00		24.71	8.20	
9.13		24.17	8.00	
9.25		23.64	7.80	
9.38		23.13	7.60	
9.50		22.71	7.30	
9.63		22.32	7.00	
9.75		21.84	6.85	
9.88		21.59	6.40	

THETA	ABFLR	CBT	CA
10.13	20.89	5.90	
10.25	20.61	5.60	
10.38	20.29	5.40	
10.50	19.98	5.20	
10.63	19.69	5.00	
10.75	19.41	4.80	
10.88	19.15	4.60	
11.00	18.91	4.40	
11.13	18.69	4.20	
11.25	18.48	4.00	
11.38	18.15	4.00	
11.50	17.81	4.00	
11.63	17.48	4.00	
11.75	17.15	4.00	
11.88	16.81	4.00	
12.00	16.48	4.00	
12.13	16.15	4.00	
12.25	15.81	4.00	

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PROGRAM DEDBIS(INPUT, TAPES=INPUT, OUTPUT, TAPE6=OUTPUT)

X=THE TIME PASSED AFTER PULSE DYE INJECTION, IN MINUTE
Y=THE MEASURED DYE CONCENTRATION AT TIME T, THE DYE CONCENTRATION
AT TIME JUST AFTER PULSE DYE INJECTION
 ΔTH_1^2 =THE HYDRAULIC PULSE LOADING AND INTERFLOW RATES RESPECTIVELY,
INTERMINUTES PER MINUTE OBSERVATIONS

NOB=NUMBER OF PARAMETERS
NP=NUMBER OF PARAMETERS
MIT²,EPS1,EP2,FLAM,FNU=SOME PREASSIGNED VALUES IN UWHAUS SEARCH
DIMENSION X(17),TH(2),DIFF(2),SIGNS(2),SCRAT(100)
COMMON X(17)

EXTERNAL FUN

WRITEx(6,800)

WRITEx(6,801)

WRITEx(6,802)

WRITEx(6,803)

WRITEx(6,804)

WRITEx(6,805)

WRITEx(6,806)

WRITEx(6,807)

WRITEx(6,808)

WRITEx(6,809)

WRITEx(6,810)

WRITEx(6,811)

WRITEx(6,812)

WRITEx(6,813)

WRITEx(6,814)

WRITEx(6,815)

WRITEx(6,816)

WRITEx(6,817)

WRITEx(6,818)

WRITEx(6,819)

WRITEx(6,820)

WRITEx(6,821)

WRITEx(6,822)

WRITEx(6,823)

WRITEx(6,824)

WRITEx(6,825)

WRITEx(6,826)

WRITEx(6,827)

WRITEx(6,828)

WRITEx(6,829)

WRITEx(6,830)

WRITEx(6,831)

WRITEx(6,832)

WRITEx(6,833)

WRITEx(6,834)

WRITEx(6,835)

WRITEx(6,836)

WRITEx(6,837)

WRITEx(6,838)

WRITEx(6,839)

WRITEx(6,840)

WRITEx(6,841)

WRITEx(6,842)

WRITEx(6,843)

WRITEx(6,844)

WRITEx(6,845)

WRITEx(6,846)

CNC

800 CALCULATION OF INTERFLOW BETWEEN INNER AND OUTER CSTR BY FITTING THE IMPULSE DYE INJECTI-#1//
801 FORHAT(1,10,X,X) MUL A SUBROUTINE HAS BEEN EMPLOYED TO FIT THE #2//
802 FORHAT(1,10,X,X) ON RESULTS TO THE DEORIC AND BISCHOFF S FOR-#3//
803 FORHAT(1,10,X,X) DATA TO THE FORMULA #4//
804 FORHAT(1,10,X,X) #5//
805 FORHAT(1,10,X,X) HYDRAULIC LOADING=2.5 LITER PER MINUTE #6//
806 READ(5,1) NO 8 2 NP, MIT, EPS1, EPS2
1 FORHAT(3,1) F LAM, FNU
11 READ(5,2) X(1), I=1,4)
READ(5,2) (Y(I), I=5,8)
READ(5,2) (X(I), I=9,12)
READ(5,2) (X(I), I=13,16)
X(17)=3.20
READ(5,2) (Y(I), I=1,4)
READ(5,2) (Y(I), I=5,8)
READ(5,2) (Y(I), I=9,12)
READ(5,2) (Y(I), I=13,16)
2 FORHAT(5,3) (TH(I), I=1,2)
3 FORHAT(2F10⁻³)
DO 16 I=1, NP
DIFF(I)=C.01
SIGNS(I)=0.0
CALL UWHAUS(TFLMI, FUN, NOB, Y, NP, TH, DIFF, SIGNS, EPS1, EPS2, MIT, FLAM, F
1 NU, SCRAT)

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```

SUBROUTINE FUN(NPROB, TH, F, NOB, NP)
DIMENSION TH(2), F(17)
COMMON X(17)
VA=1.0
VB=6.0
C=(TH(1)/VA)+(TH(2)/VA)*(1.+VA/VB)
R=SQR((0.*25*(C*2.)-(TH(1)*TH(2))/(VA*VB)))
SR1=-0.5*(C)-R
SR2=-0.5*(C)+R
RK=(1./(2.*P))*((SR2+TH(2))/VB)
RL=(1./(2.*P))*((SR1+TH(2))/VB)
DO 2 I=1,NOB
 2 CONTINUE
F(I)=RK*EXP(+SR2*X(I))-RL*EXP(+SR1*X(I))
RETURN
END

```

--COMMON BLOCKS--

//

--EXTERNAL--

```

EXP.      SQRT.

```

--STATEMENT LABELS--

0B

```

--VARIABLE MAP--
          EXP.      R      64B      17      579 B.E.F.
          FUN.     -AU    0B      NOB      AU      0B
          NPROB   -IRAU  -    RK      SQR.      72B B.E.F.
          -        -      -    SQR.      SF2      74B
          -        -      -    V A      V X      67B
          -        -      -    -        -      0B. /-

```

IT LENGTH 17 STATEMENTS 19 SYMBOLS
 E USED MODEL 73 COMPILATION .021 SECONDS

17

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SUBROUTINE UWHAUS(NPROB, MODEL, NOB, Y, NP, TH, DIFF, SIGNS, EPS1, EPS2,
1 MIT, FNU, SCRAT(1))
1 DIMENSION SCRAT(1)

TA=1
TB=TA+NP
TC=TB+NP
TD=TC+NP
TE=TD+NP
TF=TE+NP
TG=TF+NOB
TH=IG+NOB
IJ=IH + NP * NOB
IJ = CALL HAUSS8(NPROB, MODEL(IH), YNP, TH, DIFF, SIGNS, EPS1, EPS2, MIT
1 FNU, SCRAT(IA), SCRAT(IF), SCRAT(IH), SCRAT(IG), SCRAT(IJ),
2 SCRAT(IF), SCRAT(IH), SCRAT(IG), SCRAT(IJ), SCRAT(IH), SCRAT(II),
3 SCRAT(IJ))
3 RETURN
END

--EXTERNAL S--

.US59

--VARIABLE MAP--

EPS1	R.A.	0.8
FNU	R.A.	0.8
HAUSS8	R.I.	1.37 B
IH	I.I.	1.34 B
IF	I.I.	1.36 B
II	I.I.	1.31 B
MODEL	I.A.	1.41 B
NP	I.A.	1.03 B
SCRAT	R.A.	0.8
TH	R.A.	0.8
Y	R.A.	C3

LENGTH 20 STATEMENTS
USED MODEL 73 COMPILED 114 SECONDS

26 SYMBOLS

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SUBROUTINE HAUSSE(NPROB, MODEL, NBO, Y₁, TBF, R, A, D, DELZ), TH, DIFZ, SIGNS, EP1S, EP2S,

1NIT, FLAX, FNU, FORTRAN VERSION

H. J. WERTZ ADAPTED FOR THE CDC 6400 (J. F. MACREGOR)

```
      DIMENSION TH(NQ), DIFZ(NQ), SIGNS(NQ), Y(NQ)
      DIMENSION Q(NQ), P(NQ), PHI(NQ), TB(NQ)
      DIMENSION F(NBO)  R(NBO)
      DIMENSION A(NQ,NQ) D(NQ,NQ)
      DIMENSION TH(1)  DIFZ(1)  SIGNS(1), DELZ(NBO,NQ)
      1 PHI(1), TB(1)  R(1), Y(1), P(1), E(1),
      ACOS(X) = ATAN(SQRT(1.0/X**2 - 1.0))
      NP = NQ
      NPROB = NPROB
      EPS1 = EP1S
      EPS2 = EP2S
      NPSQ = NP * NP
      NSCRAC = 5 * NP + NPSQ + 2 * NOB + NP * NOB
      WRITE(06,100) NPROB, NOB, NP, NSCRAC
      WRITE(06,101) NP, TH, TEMP, THEP)
      CALL F(GAS$60(1)
      CALL F(GAS$60(2)
      CALL F(GAS$60(3)
      IF(XEN0(NP-1)*0.98, 99, 16
      IF(FNU-1, 99, 16
      15 COYTINUE
      16 COYTINUE
      17 TEMP = ABS(DIFZ(I))
      18 IF(AHIN1(1.0-TEMP), ABS(TH(I))) 99, 99, 19
      19 COYTINUE
      GAT = FLAX
      NIT = 1
      LAOS(FPS1) 0, 5, 70, 70
      5 70 SSSQ = 0
      EPS1 = 0
      CALL MODEL(NPROB, TH, F, NOB, NP)
      DO 90 I=1, NOB
      R(I) = Y(I) - F(I)
      90 SSSQ = SSSQ + P(I) * F(I)
      WRITE(06,1003) SSSQ
      C
      C 100 GA = GA / FNU
      WRITE(06,1004) NIT
      101 JS = 1 - NOB
      DO 130 J=1, IP
      TEP = TH(J)
      P(J) = DIFZ(J) * TH(J)
      TH(J) = TH(J) + P(J)
      Q(J) = JS + NOB
      JS = JS + NOB
      CALL MODEL(NPROB, TH, DELZ(JS), NOB, NP)
      C
      BEGIN ITERATION
```

= HAUSS9

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IJ = JS-1
DO 120 I=1,J+1
IJ = DELZ(I,J) = DELZ(I,J) - F(I)
120 Q(J) = Q(J)/P(J)

Q = XT*2 (STEEPEST DESCENT)

C 130 TH(J) = TEMP
TF(LAOS) 131, 131, 414
131 DO 131 J=1,1
SUM = 0
KJ = NOB*(J-1)
KIJ = NOB*(I-1)
DO 160 K = 1, NOB

KIJ = KIJ + 1
SUM = SUM + DELZ(KI) * DELZ(KJ)
TEMP = SUM/(P(I)*P(J))
IJ = J + NP*(I-1)
DO 170 I = TEMP
IJ = I + NP*(J-1)
DE(IJ) = SQRT(D(IJ))

COV = 153 T = 1, NP
DO 153 J = 1, T
IJ = IJ + NP
A(IJ) = D(IJ) / E(I-1) * E(J)
IJ = J + NP*(I-1)
A(IJ) = A(IJ)

C 153

A = SCALED MOMENT MATRIX

I = 155 I = NP
DO 155 I=1,NP
P(I) = Q(I)/E(I)
PHI(I) = P(I)
II = NP + 1 + II
A(II) = A(II) + GA

C I=1
CALL MATIN(A, NP, P, I, DET)

P/E = CORRECTION VECTOR

C STEP=1.0
SUM1=0.
SUM2=0.

SUM3=0.
DO 231 I=1,NP
SUM1 = P(I)*PHI(I) + SUM1
SUM2 = P(I)*P(I) + SUM2
SUM3 = PHI(I)*PHI(I) + SUM3
PHI(I) = P(I)/SQRT(SUM2*SUM3)
TEMP = MIN1(TEMP, 1.0)
TEMP = 57.295*ACOS((TEMP)
WEITER(06, 10L1) DET, TEMP
170 DO 220 I = 1, NP

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P(I) = PHI(I) * STEP / E(I)
TBI(I) = TH(I) + P(I)

220 COYTTRUE
WRTF(06,7000)
FOR I=1,6 TEST POINT PARAMETER VALUES )
WRTF(06,2005) (TB(I),I=1,NP)
DO I=2,1 T=1, NP
IF(SIGN(I)=1) 221, NP
IF(SIGN(I)=1.0, TH(I)) + SSQ(GN(1.0,TS(I))) 663, 221, 221
COYTTRUE=0
SCAL230 N0EL(NPROB, TB, F, NOB, NP)
DO I=1,NOB
R(I)=Y(I)-F(I)
SUMB=SUMB+R(I)*R(I)
WATTE(05,1043)*SUMB
WTF(SUMB-(1.0+EPS1)*SSQ) 662, 663
WTF(AMY4*(TEMP-30.0, GA)) 665, 664
STEP=STEP/2.0
INTCNT=INTCNT+1
WTF(INTCNT-36) 170, 2700
664 GAT=GA*FHU
INTCANT=INTCNT+1
WTF(INTCNT-36) 665, 2700, 2700
665 WTI(F(06,1007)
DO I=69,I=1,NP
TH(I)=TB(I)
CALTE(95,1040,GA,SUMB)
WTF(EPS1) 229,229,229
229 DO I=40,T=1, NP
OF(ABS(P(I)/(1.0-E-20+ABS(TH(I)))-EPS2)) 240, 240, 241
225 WTF(EPS1) 270,270,265
240 COYTTRUE
WRTF(06,1009) EPS2
265 TF(A3S(SUMB-SS0)-EPS1*SS0) 256, 266, 270
266 WRTF(761010) EPS1
270 SST=SUMB
NTT=NTT+1
WTF(NTT-NTT) 100, 100, 280
2710 FOR IAT(06,2710) *** THE SUM OF SQUARES CANNOT BE REDUCED TO THE SUM
10F SQUARES AT THE END OF THE LAST ITERATION - ITERATING STOPS */
C
C 280 WRITE(05,1011) (F(I), I=1,NOB)
WRITE(06,1012) (R(I), I=1,NOB)
WRITE(05,2001) (R(I), I=1,NOB).
WRTF(GAS60(4,NOB,TEMP,0)
CALL GAS60(4,NOB,TEMP,0)
FOR KAT(//16H XBRINE-X MATIX)
SSQ=SUMB

```

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```
10FF=NOB-NP
10FITE(06,1015)
10 CALL MAT(D,NP,P,I,DET)
00 7692 I=1 NP
00 I = I + NP*(I-1)
10 F(I) = SQRT(D(I))
7692 D(I) = 1 NP
10 I = 1 NP
10 I = I + NP*(I-2) - 1
10 J = I + NP*(I-2)
10 J = I, NP
10 A(I,J) = D(I,J) / (E(I)*E(J))
10 A(I,J) = A(J,I)
10 CALL GASS6(3,10,341,TEMP,A)
10 F(I,DF) = 341
10 SDEV = SSQ/FLOAT(IDF)
10 WRITE(06,14) SDEV, IDF
10 SDEV = SQRT(SDEV)
DO 391 I=1, NP
P(I)=TH(I)+2.0*E(I)*SDEV
TBY(I)=TH(I)-2.0*E(I)*SDEV
341 WRITE(06,1039)
CALL GASS6(2, NP, TB, P, TEMP)
LAOS = 1
LAOT0 = 101
DO 415 K = 1, NOB
414 DO 420 I=1, NP
DO 420 J=1, NP
ITSUB = K+NOR*(I-1)
DESUB1 = DELZ(K+ISUB)
DEBUG1 = DELZ(K+ NOR*(I-1))
C
ITSUB = K+NOR*(J-1)
DESUB2 = DELZ(K+ISUB)
C
IJ = I + NP*(J-1)
DESUB3 = O(IJ)/D(IJ)*(DIFZ(I)*TH(I)*DIFZ(J)*TH(J))
420 TEXP = TEMP + DEBUG1 * DEBUG2 * SQR(TEMP) * SDEV
RE(K)=F(K)+TENP
F(K)=F(K)-TEHP
NRITE(06,1008)
415
415 I=1, NOB, 10
ITF=IE+1
ITF=(NOB-IE) L30,435,435
435 I=NOB
435 WRITE(06,2001) (R(J),J=1,IE)
425 WRITE(06,2006) (F(J),J=1,IE)
410 WRITE(06,1033) NPPOB
RETUN
99 WRITE(06,1034)
410 GO TO 410
10000 FORMAT(3SH1NON-LINEAR ESTIMATION PROBLEM NUMBER 13// 15
14H OBSERVATIONS, 15, 11H PARAMETERS 14, 17H SCRATCH REQUIRED)
```

Haus 59 73/73 TS

FTN 4.5+4.10

03/23/76 16 $\frac{1}{2}$:22.024 PAGE 5

```

ATAN: GASS60 MATIN MODEL OUTCI. OUTCR. SQRT.
      . . .
      --EXTERNAL S-- TAPE63

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73/73 TS

FTN 4.5+610

03/23/76 16:02.24

PAGE 1

SUBROUTINE MATIN(A, NVAR, B, NVAR, NB, DET)

DIMENSION A(NVAR, 1), B(NVAR, 1)

PIVOTH = A(1, 1)

DET = 1.0

DO 550 ICOL = 1, NVAR

PIVOTH = A(ICOL, ICOL)

PIVOTH = A(IN1(PIVOT, PIVOTH))

DET = PIVOT * DET

C DIVIDE PIVOT ROW BY PIVOT ELEMENT

A(ICOL, ICOL) = 1.0
PIVOT = ANAX((PIVOT, 1)*PIVOT)

DO 350 L = 1, NVAR

350 IF (L .NE. ICOL) A(ICOL, L) * PIVOT

372 DO 370 L = 1, NB

370 B(ICOL, L) = B(ICOL, L) * PIVOT

C REDUCE NON-PIVOT ROWS

C 371 DO 550 L = 1, NVAR

551 IF (L .NE. ICOL) 551, 550, 551

551 A(L, ICOL) = 0.

DO 450 L = 1, NVAR

450 A(L, L) = A(L, L) - A(ICOL, L)*T

552 DO 550 L = 1, NB

550 B(L, L) = B(L, L) - B(ICOL, L)*T

550 RETURN

END

--STATEMENT LABEL MAP--

*370 ID 0B :371 0 628
*560 ID 0B :550 0 1329

46B
65B

--VARIABLE LABEL MAP--

0B INTRINSIC VAR-DIM
151B
1403 ENTRY
1403
150B

AMAX1 RA CB
B ICOL TU 154B
L1 TAU 153B
NB PIVOT R 147B
T R 152B

1

PAGE 1

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FTN 4.5+410

SUBROUTINE GASS60(IITYPE,NQ,A,B,C)

DIMENSION A(NQ),B(NQ),C(NQ,NQ)

NP=NQ

NP=NP/10

LOW=1

LUP=0

TFC(NP)=15,20,30

REFURN

LUP=NP

IF(LOW=LUP) 30,30,15

30 WRITE(06,590) (J,J=LOW,LUP)

40 WRITE(06,600) (A(J),J=LOW,LUP)

50 WRITE(06,600) (S(J),J=LOW,LUP)

60 WRITE(06,600) (B(J),J=LOW,LUP)

70 IF(IITYPE.EQ.4) GO TO 70

80 DO 90 I=LOW,LUP

90 WRITE(06,720) I,(C(J,I),J=LOW,I)

100 GO TO 71

110 LOW=LOW+1

120 WRITE(06,720) I,(C(I,J),J=LOW,I)

130 CONTINUE

140 LUP=LUP+1

150 LOW2=LOW-NP

160 WRITE(06,4) 96,96,100

170 IF(I=LOW2.NP) 97

180 DO 95 I=LOW2,NP

190 WRITE(06,720) I,(C(J,I),J=LOW,LUP)

200 GO TO 100

210 WRITE(06,721) NP

220 LOW=LOW+10

230 LUP=LUP+10

240 NR=N2-1

250 GO TO 10

260 FORHAT(1/10,9112)

270 FORHAT(40E12.4)

280 FORHAT(1HG2,13,1X,F7.4,9F12.4)

290 FORHAT(1HG,13,1X,E10.4,9E12.4)

300 CONTINUE

310 RETURN

320 END

--EXTERNAL S--
GOTCR.. OUTCI.. OUTCR.. TAPE6--

PLEQ.8M	20474	L2	SL-SYS10	11/10/75	COMPASS	3.	75115
CLSF.SQ	20536	131	SL-SYS10	11/10/75	COMPASS	3.	75115
/CLSV.FD/	20607	127	SL-SYS10	11/10/75	COMPASS	3.	75115
/GET.FD/	20676	123	SL-SYS10	11/10/75	COMPASS	3.	75115
/GET.RT/	21021	17	SL-SYS10	11/10/75	COMPASS	3.	75115
GET.SQ	21030	11	SL-SYS10	11/10/75	COMPASS	3.	75115
Z.SQ	21043	1002	SL-SYS10	11/10/75	COMPASS	3.	75115
FSU.SQ	22043	101	SL-SYS10	11/10/75	COMPASS	3.	75115
F.SQ	22144	104	SL-SYS10	11/10/75	COMPASS	3.	75115
	22250	21					

1.624 CP SECONDS

343628 CM. STORAGE USED

7 TABLE MOVES

CALCULATION OF INTERFLOW BETWEEN INNER AND

OUTER CSTR BY FITTING THE IMPULSE DYE INJECTION

ON RESULTS TO THE OEDRIC AND BISCHOFF'S FOR-

MULA

UWHAUS SUBROUTINE HAS BEEN EMPLOYED TO FIT THE
DATA TO THE FORMULA

HYDRAULIC LOADING=2.5 LITER PER MINUTE

NON-LINEAR ESTIMATION, PROBLEM NUMBER***.

117 OBSERVATIONS, 2 PARAMETERS

82 SCRATCH REQUIRED

INITIAL PARAMETER VALUES

$\cdot 2500^{\frac{1}{2}} \cdot 01$ $\cdot 1400^{\frac{2}{2}} \cdot 02$

PROPORTIONS USED IN CALCULATING DIFFERENCE QUOTIENTS

$\cdot 1900^{\frac{1}{2}} \cdot 01$ $\cdot 1000^{\frac{2}{2}} \cdot 01$

INITIAL SUM OF SQUARES = $\cdot 1115 \cdot 01$

DETERMINANT = $\cdot 9841 \cdot 00$ ANGLE IN SCALED COORD. NO. $2.94^{\frac{1}{2}}$ DEGREES

TEST POINT PARAMETER VALUES
 $\cdot 2688^{\frac{1}{2}} \cdot 01$ $\cdot 1052^{\frac{2}{2}} \cdot 02$

TEST POINT SUM OF SQUARES = $\cdot 5723 \cdot 02$

PARAMETER VALUES VIA REGRESSION

$\cdot 2688^{\frac{1}{2}} \cdot 01$ $\cdot 1052^{\frac{2}{2}} \cdot 02$

AMDOA = $\cdot 1000 \cdot 02$

SUM OF SQUARES AFTER REGRESSION = $\cdot 5723121 \cdot 01$

↗

DETERMINANT = $\cdot 9931 \cdot 00$ ANGLE IN SCALED COORD. NO. $2.57^{\frac{1}{2}}$ DEGREES

TEST POINT PARAMETER VALUES
 $\cdot 2664^{\frac{1}{2}} \cdot 01$ $\cdot 1041^{\frac{2}{2}} \cdot 02$

TEST POINT SUM OF SQUARES = $\cdot 5696 \cdot 02$

PARAMETER VALUES VIA REGRESSION

$\cdot 2664^{\frac{1}{2}} \cdot 01$ $\cdot 1041^{\frac{2}{2}} \cdot 02$

AMDOA = $\cdot 1000 \cdot 03$

SUM OF SQUARES AFTER REGRESSION = $\cdot 5695983 \cdot 01$

138-

DETERMINANT = .9934E+00 ANGLE IN SCALED COORD. = 4.66 DEGREES

TEST POINT PARAMETER VALUES
.2663E+01 .1038E+02

TEST POINT SUM OF SQUARES = .5695E-02

PARAMETER VALUES VIA REGRESSION

.2663E+01 .1038E+02

LAMBDA = .100E-04

SUM OF SQUARES AFTER REGRESSION = .5694891E-02

DETERMINANT = .9935E+00 ANGLE IN SCALED COORD. = 4.22 DEGREES

TEST POINT PARAMETER VALUES
.2663E+01 .1038E+02

TEST POINT SUM OF SQUARES = .5695E-02

PARAMETER VALUES VIA REGRESSION

.2663E+01 .1038E+02

LAMBDA = .100E-05

SUM OF SQUARES AFTER REGRESSION = .5694764E-02

DETERMINANT = .9935E+00 ANGLE IN SCALED COORD. = 4.32 DEGREES

TEST POINT PARAMETER VALUES
.2663E+01 .1037E+02

TEST POINT SUM OF SQUARES = .5695E-02

PARAMETER VALUES VIA REGRESSION

.2663E+01 .1037E+02

LAMBDA = .100E-06

SUM OF SQUARES AFTER REGRESSION = .5694740E-0

DETERMINANT = .9936E+00 ANGLE IN SCALED C0020, NO. 6
TEST POINT PARAMETER VALUES

.2663E+01 .1037E+02

TEST POINT SUM OF SQUARES = .5595E-02

PARAMETER VALUES VIA REGRESSION

.2663E+01 .1037E+02

LAMBDA = .100E-07

SUM OF SQUARES AFTER REGRESSION = .5694735E-0
ITERATION STOPS - RELATIVE CHANGE IN SUM OF SQUARES LESS THAN .1000E-05

FINAL FUNCTION VALUES

• 10000E+01	• 5738E+00	- • 3999E+00	• 3242E+00	• 2869E+00	• 2648E+00
• 2490E+00	• 2359E+00	- • 2243E+00	• 2134E+00	• 1757E+00	• 1594E+00
• 2933E+00	• 1936E+00	• 1844E+00	• 1757E+00	• 1674E+00	• 1594E+00
• 1519E+00					

RESIDUALS

• 7105E-14	- • 2775E-01	• 1606E-01	• 3677E-01	• 3710E-01	• 2719E-01
• 1302E-01	• 4083E-02	- • 2252E-02	- • 6442E-02		
• 9264E-02	- • 1261E-01	- • 1643E-01	- • 1369E-01	- • 1237E-01	- • 1164E-01
- • 1189E-01					

XPRIME-X MATRIX

	1	2
1	• 3258E-01	
2	- • 3472E-03	• 5737E-03

CORRELATION MATRIX

	1	2
1	1.0000	
2	• 0803	1.0000

VARIANCE OF RESIDUALS = • 3796E-03, 15 DEGREES OF FREEDOM

INDIVIDUAL CONFIDENCE LIMITS FOR EACH PARAMETER (ON LINEAR HYPOTHESIS)

1	• 2885E+01	• 1201E+02
2	• 2447E+01	• 8742E+02

APPROXIMATE CONFIDENCE LIMITS FOR EACH FUNCTION VALUE

• 1000E+01	• 6042E+00	• 4215E+00	• 3361E+00	• 2955E+00	• 2741E+00
• 1000E+01	• 5633E+00	• 3783E+00	• 3124E+00	• 2793E+00	• 2556E+00
• 2591E+00	• 2466E+00	• 2352E+00	• 2245E+00	• 2024E+00	
• 2389E+00	• 2253E+00	• 2133E+00			

1.0773 CP SECONDS

34423B CM STORAGE USED

7 TABLE MOVES

CALCULATION OF INTERFLOW BETWEEN INNER AND

OUTER CSTR BY FITTING THE IMPULSE DYE INJECTI-

ON RESULTS TO THE DEDRIC AND BISCHOFF'S FOR-

MULA

UWHÄUS SUBROUTINE HAS BEEN EMPLOYED TO FIT THE

DATA TO THE FORMULA

HYDRAULIC LOADING=4.00 LITER PER MINUTE

NON-LINEAR ESTIMATION, PROBLEM NUMBER**#

21 OBSERVATIONS, 2 PARAMETERS 98 SCRATCH REQUIRED

INITIAL PARAMETER VALUES

$\begin{matrix} 1 \\ 2 \end{matrix}$
•4000E+01 •1400E+02

PROPORTIONS USED IN CALCULATING DIFFERENCE QUOTIENTS

$\begin{matrix} 1 \\ 2 \end{math>$
•1000E-01 •1000E-01

INITIAL SUM OF SQUARES = •1346E-01

DETERMINANT = •1001E+01 ANGLE IN SCALED COORD. = 1.49 DEGREES

TEST POINT PARAMETER VALUES

•4001E+01 •1128E+02

ST POINT SUM OF SQUARES = •9095E-02

PARAMETER VALUES VIA REGRESSION

$\begin{matrix} 1 \\ 2 \end{math>$
•4001E+01 •1128E+02

LAMBDA = •100E-02

SUM OF SQUARES AFTER REGRESSION = •9094594E-02

DETERMINANT = •9983E+00 ANGLE IN SCALED COORD. = 2.22 DEGREES

TEST POINT PARAMETER VALUES

•3992E+01

•1107E+02

ST POINT SUM OF SQUARES = •9047E-02

PARAMETER VALUES VIA REGRESSION

$\begin{matrix} 1 \\ 2 \end{math>$
•3992E+01 •1107E+02

LAMBDA = •100E-03

SUM OF SQUARES AFTER REGRESSION = •9046588E-02

DETERMINANT = .9974E+00
TEST POINT PARAMETER VALUES
.3991E+01 .1103E+02

TEST POINT SUM OF SQUARES = .9044E-02

PARAMETER VALUES VIA REGRESSION

.3991E+01 .1103E+02²

LAMBDA = .100E-04

SUM OF SQUARES AFTER REGRESSION = .9044100E-02

DETERMINANT = .9973E+00.
TEST POINT PARAMETER VALUES
.3991E+01 .1102E+02

TEST POINT SUM OF SQUARES = .9044E-02

PARAMETER VALUES VIA REGRESSION

.3991E+01 .1102E+02²

LAMBDA = .100E-05

SUM OF SQUARES AFTER REGRESSION = .9043873E-02

DETERMINANT = .9973E+00
TEST POINT PARAMETER VALUES
.3991E+01 .1102E+02

TEST POINT SUM OF SQUARES = .9044E-02

PARAMETER VALUES VIA REGRESSION

.3991E+01 .1102E+02²

DETERMINANT = .9973E+00
ANGLE IN SCALED COORD. NO. 2.84 DEGREES

TEST POINT PARAMETER VALUES
.3991E+01 .1102E+02

TEST POINT SUM OF SQUARES = .9044E-02

DETERMINANT = .9973E+00
ANGLE IN SCALED COORD. NO. 2.86 DEGREES

TEST POINT PARAMETER VALUES
.3991E+01 .1102E+02

TEST POINT SUM OF SQUARES = .9044E-02

DETERMINANT = .9973E+00
ANGLE IN SCALED COORD. NO. 2.88 DEGREES

TEST POINT PARAMETER VALUES
.3991E+01 .1102E+02

TEST POINT SUM OF SQUARES = .9044E-02

-144-

MBDA = .100E+06

SUM OF SQUARES AFTER REGRESSION = .9043838E-02

TERMINANT = .9972E+00 ANGLE IN SCALED COORD. NO. 2.88⁶ DEGREES

ST POINT PARAMETER VALUES

.3991E+01 .1102E+02

ST POINT SUM OF SQUARES = .9044E-02

PARAMETER VALUES VIA REGRESSION

.3991E+01 .1102E+02

MBDA = .100E+07 SUM OF SQUARES AFTER REGRESSION = .9043831E-02

ERATION STOPS - RELATIVE CHANGE IN SUM OF SQUARES LESS THAN .1000E-05

FINAL FUNCTION VALUES

.1000E+01	.7057E+00	.5276E+00	.4183E+00	.3500E+00	.3060E+00
.2765E+00	.2558E+00	.2404E+00	.2282E+00	.1868E+00	.1802E+00
.2180E+00	.2092E+00	.2012E+00	.1938E+00		
.1739E+00	.1679E+00	.1621E+00	.1565E+00		
.1511E+00					

RESIDUALS

.7105E-14	.4674E-01	.8580E-02	.2669E-01	.2962E-02	.3603E-01
.3047E-01	.2418E-01	.2261E-01	.2080E-01		
.9971E-02	.4844E-02	.5154E-02	.7150E-02	.1181E-01	.1222E-01
.1293E-01	.1389E-01	.1508E-01	.1849E-01		
.2910E-01					

XPRIME-X MATRIX

1	2
1 .1954E-01	
2 .2113E-03	.8293E-03

CORRELATION MATRIX

1	2
1 1.0000	
2 -.0525	1.0000

VARIANCE OF RESIDUALS = .4760E-03, 19 DEGREES OF FREEDOM

INDIVIDUAL CONFIDENCE LIMITS FOR EACH PARAMETER (ON LINEAR HYPOTHESIS)

1	2
.4303E+01	.1254E+02
.3678E+01	.9502E+01

\$J03,660,031LINR,3,1000,300,, DOUG BEECROFT
\$SCHED, SCR=5, CORE=52
\$FTNU(S,X,L,P)

PROGRAM LINRG

C DOUG BEECROFT, ENVIRONMENTAL PROTECTION SERVICE, SEPTEMBER 1972 LI
C THIS PROGRAM USES LEAST SQUARES TO ESTIMATE THE REGRESSION LI
C COEFFICIENTS FOR ANY LINEAR FUNCTION INVOLVING LESS THAN TEN LI
C INDEPENDENT VARIABLES. OUTPUT INCLUDES MEANS AND STANDARD LI
C DEVIATIONS FOR ALL VARIOELES, THE CORRELATION MATRIX, LI
C REGRESSION COEFFICIENTS, AN ANALYSIS OF VARIANCE TABLE, THE LI
C SQUARE OF THE MULTIPLE CCRELATION COEFFICIENT, AND A TABLE OF LI

C RESIDUALS. THE REQUIRED DATA CARDS ARE LISTED BELOW.

-148-

C A. COL. 1-8 PROBLM, AN EIGHT CHARACTER ALPHANUMERIC
C CODE TO BE USED IN IDENTIFYING THE
C PROBLEM.

C COL. 9-11 NPTS, THE NUMBER OF OBSERVATIONS. NPTS
C MUST BE LESS THAN 251.

C COL. 12-13 NVAR, THE TOTAL NUMBER OF VARIABLES (THE
C NUMBER OF INDEPENDENT VARIABLES + 1).
C NVAR MUST BE LESS THAN 11.

C COL. 14-15 NDEPNO, THE NUMBER OF THE DEPENDENT
C VARIABLE, THAT IS, THE COLUMN OF THE INPUT
C DATA MATRIX CONTAINING THE VALUES OF THE
C DEPENDENT VARIABLE.

C B. COL. 1-72 VARIABLE FORMAT CARD WITH WHICH THE INPUT
C DATA IS TO BE READ. SEE THE BMD MANUAL,
C PAGES 22-28, FOR A MORE COMPLETE
C DESCRIPTION OF THIS TYPE OF CARD.

C C. THE INPUT DATA. EACH CARD CONTAINS ONE OBSERVATION OF
C EACH VARIABLE. THE GENERAL FORM OF THE
C INPUT DATA IS THE SAME AS FOR THE BMD
C PROGRAMS (BMD MANUAL, PAGES 11-12).

C D. EOF CARD (STANDARD C.G.I.W. GREY-STRIPE ECF)

C NOTE -- CARDS A THROLGH C MAY BE REPEATED AS OFTEN AS DESIRED.

COMMON CORMAT,NINDEP
REAL MEANS(10),COEFF(9)
DIMENSION DATMAT(250,10),CATUSE(250,10),RESID(250),VARFMT(9)
DIMENSION CORMAT(10,10),S(10),SPARE(9),IVAR(10),YFIT(250)
EQUIVALENCE (DATUSE(1),YFIT(1)), (DATUSE(251),RESID(1))
EQUIVALENCE (VARFMT,DATLSE)

C READ CONTROL INFORMATION

10 READ 99, PROBLM,NPTS,NVAR,NCEPNO,VARFMT
99 FORMAT (A8,I3,2I2/9A8)
IF (IFEEOF(60).EQ.-1) STOP
DO 100 I=1,NPTS
100 READ (60,VARFMT) (DATMAT(I,J),J=1,NVAR)
NINDEP=NVAR-1

C COMPUTE MEANS OF ALL VARIABLES, CENTRE DATA

DO 200 I=1,NVAR
MEANS(I)=0.0
DO 300 J=1,NPTS
300 MEANS(I)=MEANS(I)+DATMAT(J,I)
MEANS(I)=MEANS(I)/NPTS
DO 200 J=1,NPTS
200 DATUSE(J,I)=DATMAT(J,I)-MEANS(I)

C COMPUTE SUM OF SQUARES OF DEVIATIONS FROM MEAN FOR VARIABLES

DO 500 I=1,NVAR
S(I)=J.0

```
    DO 600 J=1,NPTS  
600 S(I)=S(I)+DATUSE(J,I)**2  
    SAVE=SQRT(S(I))  
    DO 500 J=1,NPTS  
500 DATUSE(J,I)=DATUSE(J,I)/SAVE
```

-149-

```
C  
C      PRINT CONTROL INFORMATION, MEANS, AND STANDARD DEVIATIONS  
C
```

```
PRINT 44, PROBLM,NPTS,NVAR,NDEPND  
44 FORMAT (27H1 MULTIPLE LINEAR REGRESSION  
1     /13H0 PROBLEM CODE,A23  
2     /23H NUMBER OF OBSERVATIONS,10X,I3  
3     /20H NUMBER OF VARIABLES,I16  
4     /29H DEPENDENT VARIABLE IS NUMBER,I7/  
5     /30H0 MEANS AND STANDARD DEVIATIONS  
6 //11X,42H VARIABLE NO.   MEAN   STANDARD DEVIATION/  
DO 1030 I=1,NVAR  
    SAVE=SQRT(S(I)/(NPTS-1))  
1030 PRINT 22,I,MEANS(I),SAVE  
    22 FORMAT (16X,I2,2F15.5)
```

```
C  
C      COMPUTE AND PRINT CORRELATION MATRIX  
C
```

```
DO 700 I=1,NVAR  
DO 700 J=1,NVAR  
    CORHAT(I,J)=0.0  
DO 700 K=1,NPTS  
700 CORMAT(I,J)=CORMAT(I,J)+DATUSE(K,I)*DATUSE(K,J)  
PRINT 88,(I,I=1,NVAR)  
88 FORMAT (/19H0 CORRELATION MATRIX//9X,10(7X,I2))  
DO 800 I=1,NVAR  
800 PRINT 77,I,(CORMAT(I,K),K=1,NVAR)  
77 FORMAT (8X,I1,3X,13F9.5)
```

```
C  
C      TRANSFER DEPENDENT VARIABLE TO LAST ROW AND LAST COLUMN  
C
```

```
DO 18 I=1,NVAR  
18 IVAR(I)=I  
IF (NDEPND.EQ.NVAR) GO TO 32  
DO 17 I=1,NVAR  
    SWAP=CORMAT(I,NVAR)  
    CORMAT(I,NVAR)=CORMAT(I,NDEPND)  
17 CORMAT(I,NDEPND)=SWAP  
DO 16 I=1,NVAR  
    SWAP=CORMAT(NVAR,I)  
    CORMAT(NVAR,I)=CORMAT(NDEPND,I)  
16 CORMAT(NDEPND,I)=SWAP  
    IVAR(NDEPND)=NVAR  
    IVAR(NVAR)=NDEPND.
```

```
C  
C      COMPUTE PRODUCT OF TRANSPOSE MATRIX OF X VALUES WITH Y VALUES  
C
```

```
32 DO 900 I=1,NINDEP  
    SPARE(I)=0.0  
    L=IVAR(I)  
    DO 900 K=1,NPTS  
900 SPARE(I)=SPARE(I)+DATUSE(K,L)*DATUSE(K,NDEPND)
```

```
C  
C      COMPUTE INVERSE OF CORRELATION MATRIX  
C
```

```
CALL MATINV
```

C COMPUTE AND PRINT REGRESSION COEFFICIENTS

-150-

C PRINT 66
66 FORMAT (//24H REGRESSION COEFFICIENTS
1 //11X,27H VARIABLE NC. COEFFICIENT)
CONST=MEANS(NDEPND)
DO 1000 I=1,NINDEP
COEFF(I)=0.0
L=IVAR(I)
DO 400 K=1,NINDEP
400 COEFF(I)=COEFF(I)+CORMAT(I,K)*SPARE(K)
COEFF(I)=COEFF(I)*SQRT(S(NDEPND)/S(L))
CONST=CONST-COEFF(I)*MEANS(L)
1000 PRINT 55, L, COEFF(I)
55 FORMAT (14X,I2,E21.5)
PRINT 14, CONST
14 FORMAT (/11X,38H CONSTANT TERM IN PREDICTION EQUATION =,F12.5)

C
C PRINT ANALYSIS OF VARIANCE AND CORRELATION COEFFICIENT
C

REGSS=0.0
DO 1040 I=1,NPTS
YFIT(I)=CONST
DO 1050 J=1,NINDEP
L=IVAR(J)
1050 YFIT(I)=YFIT(I)+COEFF(J)*BATMAT(I,L)
RESID(L)=DATMAT(L,NDEPND)-YFIT(I)
1040 REGSS=REGSS+YFIT(I)*BATMAT(I,NDEPND)
TOTAVG=NPTS*MEANS(NDEPNC)**2
TOTSS=TOTAVG+S(NDEPND)
RESSS=TOTSS-REGSS
REGSS=S(NDEPND)-RESSS
REGMS=REGSS/NINDEP
I=NPTS-1
J=NPTS-NVAR
VARYNC=RESSS/J
F=REGMS/VARYNC
CORRCO=REGSS/S(NDEPND)
PRINT 1107,TOTSS,NPTS,TOTAVG,S(NINDEPND),I,REGSS,NINDEP,REGMS,F,
1 RESSS,J,VARYNC,CORRCO
1107 FORMAT (21H ANALYSIS OF VARIANCE /1H,12X,71HSOURCE OF VARIATION
1SUM OF SQUARES DEGREES OF FREEDOM MEAN SQUARE,10X,1HF
2 /1H,12X,19HTOTAL (UNCORRECTED),F16.4,I15/1H ,27X,4HMEAN,
3 F16.4,14X,1H/1H ,14X,17HTOTAL (CORRECTED),F16.4,I15/1H ,21X
4 13HREGRESSION,F16.4,I15,F21.4,F15.4/1H ,23X,8HRESIDUAL,F16.4
5 I15,F21.4/33H CORRELATION COEFFICIENT (R**2) =,F9.5)

C
C PRINT TABLE OF RESIDUALS.
C

PRINT 12
12 FORMAT (19H1TABLE OF RESIDUALS
1 /1H,4X,29HOBSERVATION NO. OBSERVED Y,8X,8HFITTED Y,8
2 8HRESIDUAL/)
DO 1060 I=1,NPTS
1060 PRINT 13, I,DATMAT(I,NDEPND),YFIT(I),RESID(I)
13 FORMAT (10X,I3,10X,F11.5,6X,F11.5,5X,F10.5)
GO TO 10
END
SUBROUTINE MATINV
C
C MATRIX INVERSION
C

```

COMMON A,N
DIMENSION A(10,10),INDEX(10,3)
EQUIVALENCE (IROW,JROW),(ICOLUMN,JCOLUMN),(AMAX,T,SWAP)
DO 60 J=1,N
60 INDEX(J,3)=0

```

```

C
C      SEARCH FOR PIVOT ELEMENT
C

```

```

DO 550 I=1,N
AMAX=0.0
DO 105 J=1,N
IF (INDEX(J,3).EQ.1) GO TO 105
DO 100 K=1,N
IF (INDEX(K,3)-1) 80,100,715
80 IF (AMAX.GE.ABS(A(J,K))) GO TO 100
IROW=J
ICOLUMN=K
AMAX=ABS(A(J,K))
100 CONTINUE
105 CONTINUE
INDEX(ICOLUMN,3)=INDEX(ICOLUMN,3)+1
INDEX(I,1)=IROW
INDEX(I,2)=ICOLUMN
IF (IROW.EQ.ICOLUMN) GO TO 310

```

```

C
C      INTERCHANGE ROWS
C

```

```

DO 200 L=1,N
SWAP=A(IROW,L)
A(IROW,L)=A(ICOLUMN,L)
200 A(ICOLUMN,L)=SWAP

```

```

C
C      REDUCE NON-PIVOT ROWS
C

```

```

310 PIVOT=A(ICOLUMN,ICOLUMN)
A(ICOLUMN,ICOLUMN)=1.0
DO 350 L=1,N
350 A(ICOLUMN,L)=A(ICOLUMN,L)/PIVOT
DO 550 L1=1,N
IF (L1.EQ.ICOLUMN) GO TO 550
T=A(L1,ICOLUMN)
A(L1,ICOLUMN)=0.0
DO 450 L=1,N
450 A(L1,L)=A(L1,L)-A(ICOLUMN,L)*T
550 CONTINUE

```

```

C
C      INTERCHANGE COLUMNS
C

```

```

DO 710 I=1,N
L=N+1-I
IF (INDEX(L,1).EQ.INDEX(L,2)) GO TO 710
JROW=INDEX(L,1)
JCOLUMN=INDEX(L,2)
DO 705 K=1,N
SWAP=A(K,JROW)
A(K,JROW)=A(K,JCOLUMN)
705 A(K,JCOLUMN)=SWAP
710 CONTINUE
DO 730 K=1,N
IF (INDEX(K,3).NE.1) GO TO 715

```

C
C PRINT ERROR MESSAGE
C

-152-

715 PRINT 99

99 FORMAT (61H0INVERSE OF MATRIX CANNOT BE COMPUTED -- EXECUTION TERM
1INATED)

STOP

END

IN(2.3)/MASTER INTEGER WORD SIZE = 1 , * OPTION IS CFF , C OPTION IS OFF

PROGRAM CON95

CON9501

C DOUG BEECROFT, ENVIRONMENTAL PROTECTION SERVICE, OCTOBER 1972 CON9501

C THIS PROGRAM COMPUTES THE 95 PERCENT CONFIDENCE LIMITS FOR ANY CON9501
C LINEAR REGRESSION EQUATION. THE NECESSARY DATA CARDS ARE CON9501
C LISTED BELOW. CON9501

C A. COL. 1-8 PROBLM, AN EIGHT CHARACTER ALPHANUMERIC CON9501
C CODE USED TO IDENTIFY THE PROBLEM. CON9501

C COL. 9-11 NPTS, THE NUMBER OF OBSERVATIONS USED IN CON9501
C COMPUTING THE REGRESSION COEFFICIENTS. CON9501
C NPTS MUST BE LESS THAN 501. CON9501

C COL. 12-13 NVAR, THE TOTAL NUMBER OF VARIABLES CON9501
C INVOLVED IN THE REGRESSION (THE NUMBER OF CON9501
C INDEPENDENT VARIABLES + 1). NVAR MUST BE CON9501
C LESS THAN 11. CON9501

C COL. 14-16 NLIM, THE NUMBER OF CONFIDENCE LIMITS TO CON9501
C BE COMPUTED. CON9501

C COL. 17-30 VARYNC, THE RESIDUAL MEAN SQUARE OR CON9501
C VARIANCE ABOUT THE REGRESSION. THIS CON9501
C STATISTIC IS GIVEN IN THE ANALYSIS OF CON9501
C VARIANCE TABLE COMPUTED BY LINRG OR SPRG. CON9501
C THE NUMBER MUST CONTAIN A DECIMAL POINT. CON9501

C B. THE COMPUTED REGRESSION COEFFICIENTS. EACH CON9501
C COEFFICIENT, BEGINNING WITH THE CONSTANT CON9501
C TERM AND FOLLOWED BY THE VARIABLE CON9501
C COEFFICIENTS, IN ORDER, MUST OCCUPY 15 CON9501
C COLUMNS OF THE CARD. IF NVAR IS GREATER CON9501
C THAN 5, TWO CARDS WILL BE REQUIRED. CON9501

C C. COL. 1-72 VARIABLE FORMAT CARD TO BE USED IN READING CON9501
C THE INPUT DATA. SEE THE BMD MANUAL, PAGES CON9501
C 22-28 FOR A DESCRIPTION OF THIS TYPE OF CON9501
C CARD. ONLY THE INDEPENDENT VARIABLE CON9501
C VALUES NEED BE READ IN. CON9501

C D. THE INPUT DATA. THE RAW DATA FROM WHICH THE CON9501
C REGRESSION COEFFICIENTS WERE CALCULATED. CON9501
C THE OBSERVED VALUES OF THE DEPENDENT CON9501
C VARIABLE MUST NOT BE INCLUDED. VALUES CON9501
C OF THE OTHER VARIABLES MUST APPEAR IN THE CON9501
C SAME ORDER AS THEIR RESPECTIVE CON9501
C COEFFICIENTS IN CARD B. CON9501

C E. COL. 1-72 VARIABLE FORMAT CARD TO BE USED IN READING CON9501
C THE POINTS AT WHICH CONFIDENCE LIMITS ARE CON9501

-15-

D4N(2.3)/MASTER INTEGER WORD SIZE = 1 , * OPTION IS CFF , C OPTION IS OFF

```

C          COMPUTE COVARIANCE MATRIX           CON95
C          CALL MATINV                         CON95
C          READ IN THE CO-ORDINATES OF THE POINT AND COMPUTE THE FITTED Y CON95
C          READ &S, VAREFMT                   CON95
      DO 70 J=1,NLIM                         CON95
      POINT(1)=1.0                           CON95
      READ (60,VAREFMT) (POINT(I),I=2,NVAR)   CON95
      YFIT=COEFF(1)                          CON95
      DO 200 I=2,NVAR                        CON95
      200 YFIT=YFIT+COEFF(I)*POINT(I)        CON95
C          COMPUTE PRODUCT OF THE POINT AND THE COVARIANCE MATRIX CON95
C          CONFID=3.0                         CON95
      DO 800 I=1,NVAR                        CON95
      SPARE=3.0                            CON95
      DO 900 K=1,NVAR                        CON95
      900 SPARE=SPARE+POINT(K)*COVINV(K,I)   CON95
      800 CCNFID=CONFID+SPARE*PCINT(I)       CON95
C          CHOOSE APPROPRIATE T VALUE AND COMPUTE CONFIDENCE LIMIT CON95
C          IJK=NPTS-NVAR                     CON95
      IF (IJK.GT.30) GO TO 10                CON95
      T=TVALUES(IJK)                         CON95
      GO TO 20                                CON95
      10 IF (IJK.GT.35) GO TO 30              CON95
      T=TVALUES(30)                           CON95
      GO TO 20                                CON95
      30 IF (IJK.GT.50) GO TO 40              CON95
      T=TVALUES(31)                           CON95
      GO TO 20                                CON95
      40 IF (IJK.GT.90) GO TO 50              CON95
      T=TVALUES(32)                           CON95
      GO TO 20                                CON95
      50 T=TVALUES(33)                         CON95
      20 CONFID=SQRT (CONFID*VARYNC)+T       CON95
      UF=YFIT+CONFID                         CON95
      DOWN=YFIT-CONFID                       CON95
      70 PRINT 65, J,DOWN,YFIT,UF             CON95
      66 FORMAT (I15,F18.5,F11.5,F14.5)     CON95
      GO TO 65                                CON95
      END .                                 CON95
  
```

AN(2.3)/MASTER INTEGER WORD SIZE = 1 , * OPTION IS CFF , C OPTION IS OFF

```

SUBROUTINE MATINV                                                    CON951
C
C     MATRIX INVERSION                                            CON951
C
COMMON A,N                                                        CON951
DIMENSION A(10,10),INDEX(10,3)                                CON951
EQUIVALENCE (IROW,JROW),(ICOLUMN,JCOLUMN),(AMAX,T,SWAP)    CON951
DO 60 J=1,N                                                    CON951
60 INDEX(J,3)=0                                                CON951
C
C     SEARCH FOR PIVOT ELEMENT                                CON951
C
DO 550 I=1,N                                                    CON951
AMAX=0.0                                                        CON951
DO 105 J=1,N                                                    CON951
IF (INDEX(J,3).EQ.1) GO TO 105                            CON951
DO 100 K=1,N                                                    CON951
IF (INDEX(K,3).EQ.1) 80,100,715                            CON951
80 IF (AMAX.GE.ABS(A(J,K))) GO TO 100                    CON951
IROW=J                                                            CON951
ICOLUMN=K                                                        CON951
AMAX=ABS(A(J,K))                                            CON951
100 CONTINUE                                                    CON951
105 CONTINUE                                                    CON951
INDEX(ICOLUMN,3)=INDEX(ICOLUMN,3)+1                        CON951
INDEX(I,1)=IROW                                                CON951
INDEX(I,2)=ICOLUMN                                            CON951
IF (IROW.EQ.ICOLUMN) GO TO 310                            CON951
C
C     INTERCHANGE ROWS                                        CON951
C
DO 250 L=1,N                                                    CON951
SWAP=A(IROW,L)                                                CON951
A(IROW,L)=A(ICOLUMN,L)                                    CON951
200 A(ICOLUMN,L)=SWAP                                        CON951
C
C     REDUCE NON-PIVOT ROWS                                CON951
C
310 PIVOT=A(ICOLUMN,ICOLUMN)                                CON951
A(ICOLUMN,ICOLUMN)=1.0                                        CON951
DO 350 L=1,N                                                    CON951
350 A(ICOLUMN,L)=A(ICOLUMN,L)/PIVOT                        CON951
DO 550 L1=1,N                                                CON951
IF (L1.EQ.ICOLUMN) GO TO 550
T=A(L1,ICOLUMN)
A(L1,ICOLUMN)=1.0                                            CON951
DO 450 L=1,N                                                    CON951
450 A(L1,L)=A(L1,L)-A(ICOLUMN,L)*T                        CON951
550 CONTINUE                                                    CON952
C
C     INTERCHANGE COLUMNS                                    CON952
C

```

PAN(2.3)/MASTER INTEGER WORD SIZE = 1 , * OPTION IS OFF , 0 OPTION IS OFF

```
DC 710 I=1,N                                                                    CON352
L=N+1-1                                                                    CON952
IF (INDEX(L,1).EQ.INDEX(L,2)) GO TO 710                            CON1952
JROW=INDEX(L,1)                                                        CON352
JCOLUMN=INDEX(L,2)                                                    CON1952
DC 705 K=1,N                                                            CON352
SWAP=A(K,JROW)                                                        CON952
A(K,JROW)=A(K,JCOLUMN)                                            CON952
705 A(K,JCOLUMN)=SWAP                                                CON952
710 CCONTINUE                                                            CON1952
DC 710 K=1,N                                                            CON952
IF (INDEX(K,3).NE.1) GO TO 715                                    CON352
730 CCONTINUE                                                            CON952
RETURN                                                                    CON1952
C                                                                            CON952
C PRINT ERROR MESSAGE                                                    CON352
C                                                                            CON952
715 PRINT 99                                                            CON1952
99 FORMAT (6IHOINVERSE OF MATRIX CANNOT BE COMPUTED -- EXECUTION TERMCON952
11HATEU)
STOP                                                                    CON352
END                                                                        CON952
```

SASI FORTRAN DIAGNOSTIC RESULTS FOR MATINV

NO ERRORS

APPENDIX 3

In this appendix, the following are included:

Table 3-1: Calculation of reaction rate constants for hydraulic loading of 2.5 l/min, assuming zero order kinetics.

Table 3-2: Calculation of reaction rate constants for hydraulic loading of 4.0 l/min, assuming zero order kinetics (Analytical results of parallel and series operation are included).

Table 3-3: Same as table 1, but for first order kinetics.

Table 3-4: Same as table 2, but for first order kinetics.

Zero Order Kinetics

In order to determine K and SB values of equation 17 and 18 (page 50,53), n was taken to be zero.

$$v S_o - K A_A - v_B S_A + v_B S_B - v S_A = 0 \quad (1)$$

$$v_B S_A - K A_B - v_B S_B = 0 \quad (2)$$

Equation (1) and (2) yields:

$$K = \frac{v(S_o - S_A)}{A}$$

$$S_B = S_A - \frac{K A_B}{v_B}$$

The calculations are shown on Table 3-1 and 3-2 of this appendix for hydraulic loadings of 2.5 l/min and 4.0 l/min respectively.

First Order Kinetics

(a) For hydraulic loading of 4 l/min (Table 3-4)

Formula 17 and 18 yields:

$$K = 2.45 \frac{S_A}{S_B} - 2.45 \quad (1)$$

$$S_B^2 - (0.956 S_A - 0.285 S_o) S_B - 0.329 S_A^2 = 0$$

by inserting:

$$v = 4 \text{ l/min}$$

$$V_B = 14 \text{ l/min}$$

$$A_B = 5.7 \text{ m}^2$$

$$A_A = 1.9 \text{ m}^2$$

$$n = 1$$

(b) For hydraulic loading of 2.5 l/min (Table 3-3)

Formula 17 and 18 gives

$$K = 2.10 \frac{s_A}{s_B} = 2.10$$

$$s_B^2 - (0.879 s_A - 0.208 s_O) s_B - 0.329 s_A^2 = 0$$

TABLE 3-1

date	Run No.	Temp. °C	Hydraulic Loading (L/min)	S_O ppm	S_A ppm	$S - S_A$ ppm	$K = \frac{(S_O - S_A)}{A} V$ mg/m ² -min	$K \frac{AB}{VB}$	$S_B = S_A - \frac{K}{V_B}$ (ppm)	A_B mg/m ² -hr	K mg/m ² -hr	lnK
7/9/75	18	5	2.5	13.6	10.25	3.35	1.10	0.52	9.73	66.11	4.18	
8/9/75	18	5	2.5	25.20	17.10	8.10	2.66	1.26	15.84	159.60	5.07	
2/10/75		10	2.5	17.00	12.30	4.70	1.54	0.73	11.57	92.40	4.53	
3/10/75		10	2.5	18.00	12.90	5.10	1.67	0.79	12.11	100.20	4.61	
1/9/75		15	2.5	13.75	4.75	9.00	2.96	1.27	3.48	177.60	5.17	
2/9/75		15	2.5	13.30	4.70	8.60	2.82	1.33	3.37	169.20	5.13	
0/10/75		20	2.5	30.00	6.90	23.10	7.59	3.60	3.30	455.40	6.12	
1/10/75		20	2.5	29.00	13.00	16.00	5.26	2.49	.10.51	315.60	5.75	
5/9/75		25	2.5	32.00	6.20	25.80	8.48	4.02	2.18	528.80	6.23	
8/9/75		25	2.5	22.90	3.85	19.05	6.26	2.97	0.88	375.60	5.92	
6/9/75		25	2.5	29.00	2.80	26.20	16.78	---	0.00	1007.36	6.91	

TABLE 3-2

Date	Run No.	Temp. °C	Hydraulic Loading 1/min	S_o (ppm)	S_A (ppm)	$S_o - S_A$ (ppm)	$K = \frac{(S_o - S_A) V}{A}$ mg/m ² -min	$\frac{A_B}{K_{V_B}}$	$S_B = S_A - K \frac{A_B}{V_B}$	$\frac{A_B}{mg/m^2 \cdot hr}$	lnK
17/9/75		5	4	13.60	12.65	0.95	0.49	0.19	12.46	30.00	3.40
18/9/75		5	4	26.20	23.60	2.60	1.36	0.55	23.05	82.10	4.40
21/9/75		5	4	15.35	11.45	3.90	2.05	0.83	10.62	123.15	4.81
22/10/75		5	4	17.00	15.30	1.70	0.89	0.36	14.94	53.68	3.98
10/7/75		5	4	14.10	10.40	3.70	1.94	0.78	9.62	116.84	4.76
22/10/75		10	4	17.00	14.90	2.10	1.10	0.44	14.46	66.31	4.19
23/10/75		10	4	18.00	15.00	3.00	1.57	0.63	14.37	94.73	4.55
17/3/75		10	4	17.00	9.40	7.60	3.99	1.62	7.78	239.99	5.48
3/7/75		15	4	19.00	14.40	4.60	2.42	0.98	13.42	145.26	4.97
10/10/75		20	4	30.00	10.00	20.00	10.52	4.28	5.72	631.57	6.44
3/10/75		25	4	30.00	7.65	22.35	11.76	4.78	2.87	705.94	6.55
16/8/75		25	4	40.00	16.7	23.30	12.26	4.99	11.71	735.75	6.60
17/8/75		25	4	54.00	42.00	12.00	6.31	2.56	39.44	378.94	5.93

TABLE 3-3

Date	S_O ppm	0.208 S_O	S_A ppm	0.879 S_A	$0.879 S_A^-$	$0.329 S_A^2$	$a S_B^{2-b} S_B - c = 0$	$\sqrt{b^2 - 4ac}$
17/9/75	13.60	2.82	10.25	9.00	6.18	34.56	$S_B^{2-6.18} S_B - 34.56$	13.28
18/9/75	25.20	5.24	17.10	15.03	9.79	96.20	$S_B^{2-9.79} S_B - 96.20$	21.93
22/10/75	17.00	3.53	12.30	10.86	7.33	49.77	$S_B^{2-7.33} S_B - 49.77$	15.89
23/10/75	18.00	3.74	12.90	11.33	7.59	54.74	$S_B^{2-7.59} S_B - 54.74$	16.63
11/9/75	13.75	2.86	4.75	4.17	1.31	7.42	$S_B^{2-1.31} S_B - 7.42$	5.60
12/9/75	13.30	2.76	4.70	4.13	1.37	7.26	$S_B^{2-1.37} S_B - 7.26$	5.56
10/10/75	30.00	6.24	6.90	6.06	-0.18	15.66	$S_B^{2+0.18} S_B - 15.66$	7.91
11/10/75	29.00	6.03	13.00	11.42	5.39	55.60	$S_B^{2-5.39} S_B - 55.60$	15.85
25/9/75	32.00	6.65	6.20	5.44	-1.21	12.64	$S_B^{2+1.21} S_B - 12.64$	7.21
28/9/75	22.90	4.76	3.85	3.38	-1.38	4.87	$S_B^{2+1.38} S_B - 4.87$	4.62
26/9/75	29.00	6.03	2.80	2.46	-3.57	2.57	$S_B^{2+3.57} S_B - 2.57$	4.79

TABLE 3-3 (Cont'd)

$-b + \sqrt{b^2 - 4ac}$	$-b - \sqrt{b^2 - 4ac}$	S_B	$K = 2.10 \frac{SA}{SB} 2.10$	lnk
19.66	-7.10	9.73	0.112	-2.18
31.72	-12.14	15.86	0.165	-1.80
23.22	-8.56	11.61	0.124	-2.08
24.22	-9.04	12.11	0.138	-1.98
6.91	-4.29	3.45	0.79	-0.23
6.93	-4.19	3.46	0.76	-0.27
7.73	-8.09	3.86	1.68	-0.51
21.24	-10.46	10.62	0.47	-0.75
6.00	-8.41	3.00	2.23	-0.80
3.24	-6.00	1.62	2.89	+1.06
+1.22	-8.36	0.61	7.53	2.01

TABLE 3-4

Date	S_O (ppm)	$0.285 S_O$ (ppm)	S_A , (ppm)	$0.956 S_A$ (ppm)	$0.956 S_A - 0.285 S_O$ (ppm)	$0.329 S_A^2$	$a S_B^2 - b S_B - c = 0$	$\sqrt{b^2 - 4ac}$
17/9/75	13.60	3.87	12.65	12.09	8.22	52.64	$S_B^2 - 8.22 S_B - 52.64 = 0$	16.67
18/9/75	26.20	7.46	23.60	22.56	15.10	183.23	$S_B^2 - 15.10 S_B - 183.23 = 0$	30.99
21/9/75	15.35	4.37	11.45	10.94	6.57	43.13	$S_B^2 - 6.57 S_B - 43.13 = 0$	14.68
22/9/75	17.00	4.84	14.90	14.24	9.40	73.04	$S_B^2 - 9.40 S_B - 73.04 = 0$	19.50
10/10/75	14.10	4.01	10.40	9.94	5.93	35.58	$S_B^2 - 5.93 S_B - 35.38 = 0$	13.32
22/10/75	17.00	4.84	15.30	14.62	9.78	77.01	$S_B^2 - 9.78 S_B - 77.01 = 0$	20.09
23/10/75	18.00	5.13	15.00	14.34	9.21	74.02	$S_B^2 - 9.21 S_B - 74.02 = 0$	19.51
17/11/75	17.00	4.84	9.40	8.98	4.14	29.07	$S_B^2 - 4.14 S_B - 29.07 = 0$	11.55
3/11/75	19.00	5.41	14.40	13.76	8.35	68.22	$S_B^2 - 8.35 S_B - 68.22 = 0$	18.50
10/10/75	30.00	8.55	10.00	9.56	1.01	32.90	$S_B^2 - 1.01 S_B - 32.90 = 0$	11.51
3/10/75	30.00	8.55	7.65	7.31	-1.24	19.253	$S_B^2 - 1.24 S_B - 19.35 = 0$	8.86
16/11/75	40.00	11.40	23.30	22.27	10.87	178.61	$S_B^2 - 10.87 S_B - 178.61 = 0$	28.85
17/11/75	54.00	15.39	42.00	40.15	24.76	580.35	$S_B^2 - 24.76 S_B - 580.75 = 0$	54.15

TABLE 3-4 (Cont'd)

$-b + \sqrt{b^2 - 4ac}$	$-b - \sqrt{b^2 - 4ac}$	S_B (ppm)	$K = 2.45 \frac{S_A}{S_B} - 2.45$	Link
24.89	-8.45	12.44	0.041	-3.18
46.09	-15.89	23.04	0.059	-2.83
21.25	-8.11	10.62	0.191	-1.65
28.90	-10.10	14.45	0.076	-2.57
19.25	-7.39	9.62	0.195	-1.63
29.87	-10.31	14.93	0.060	-2.81
28.72	-10.30	14.36	0.109	-2.21
15.69	7.41	7.84	0.468	-0.75
26.85	9.95	13.42	0.178	-1.72
12.52	-10.50	6.26	1.463	0.38
7.62	-10.10	3.81	2.470	0.90
39.72	-17.98	19.86	0.431	-0.84
78.91	-29.39	39.46	0.157	-1.85

APPENDIX 4

ANALYSIS OF VARIANCE

The analysis of variance results for the linearized Arrhenius models were obtained by using computer program (Appendix 2). The mean square pure errors were calculated using repeats. F test was applied to check lack of fit for 95% confidence level.

ANOVA NO. 1 (For hydraulic loading of 2.5 l/min) (Zero order kinetics)

Linearized Arrhenius model:

$$\ln K = -\frac{E}{R} \left(\frac{1}{T} - \frac{1}{T_0} \right) + \ln A_0 e^{-E/RT_0}$$

$$\ln K = -7792 \left(\frac{1}{T} - \frac{1}{T_0} \right) + 5.35$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Square	F
Total (uncorrected)	330.2041	11		
Mean	323.2488	1		
Total (corrected)	6.9553			
Regression	5.4942	1	5.4942	33.84
Residual	1.4611	9	0.1623	
Pure Error	0.9614	6	0.1603	
Lack of Fit	0.4996	3	0.1663	1.03

Lack of fit = 1 < 4.75
 Pure Error

Pure Error Estimates:

$$P_E^2 = \sum_{i=1}^m \sum_{j=1}^K (y_{ij} - \bar{y}_i)^2 = 2x(0.44)^2 + (0.04)^2 x 2 + (0.02)^2 x 2 + (0.185)^2 x 2 \\ + (0.12)^2 + (0.43)^2 + (0.55)^2 = 0.9614$$

where P_E^2 = pure error square,

\bar{y}_i = mean of observations at ith condition,

y_{ij} = jth observation at ith condition,

K = number of observation at each condition, and

m = number of conditions studied.

ANOVA NO. 2 (For hydraulic loading = 4 l/min, zero order kinetics)

Linearized Arrhenius Model:

$$\ln K = -8753 \left(\frac{1}{T} - \frac{1}{T_0} \right) + 5.35$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Square	F
Total (uncorrected)	348.6450	13		
Mean	335.6864	1		
Total (corrected)	12.9586	12		
Regression	9.8983	1	9.8983	35.57
Residual	3.0603	11	0.2782	
Pure Error	2.5544	8	0.3193	
Lack of Fit	0.5059	3	0.1686	0.53

Pure Error Estimate:

$$\begin{aligned} P_E^2 &= (0.49)^2 + (0.87)^2 + (0.13)^2 + (0.54)^2 + (0.29)^2 + (0.74)^2 \\ &\quad + (0.55)^2 + (0.19)^2 + (0.19)^2 + (0.43)^2 + (0.24)^2 = 2.5544 \end{aligned}$$

$$\frac{\text{Lack of Fit}}{\text{Pure Error}} = \frac{0.1686}{0.3193} = 0.528 < F_{0.95}(3,8) = 4.0662$$

ANOVA NO. 3 (For combined data of 2.5 l/min and 4.0 l/min operation and zero order kinetics)

Linearized Arrhenius Model:

$$\ln K = -8357 \quad \left(\frac{1}{T} - \frac{1}{T_0} \right) + 5.34$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Squares	F
Total (uncorrected)	678.8491	24		
Mean	658.2490	1		
Total (corrected)	20.6001	23		
Regression	16.0295	1		35.57
Residual	4.5706	22		
Pure Error	1.8717	14	0.1336	
Lack of Fit	2.6989	8	0.3373	2.52

Pure Error Estimate:

$$P_E^2_C = \frac{P_E^2_{2.5} \times v_{2.5} + P_E^2_{4.0} \times v_{4.0}}{v_{2.5} + v_{4.0}} = \frac{6 \times 0.96145 + 8 \times 2.5544}{6 + 8}$$

$$= 1.8717$$

where: $P_{E_C}^2$ = Pure error square of combined data

$P_{E_{2.5}}^2$ = Pure error square of data for 2.5 l/min hydraulic loading

$P_{E_{4.0}}^2$ = Pure error square of data for 4.0 l/min hydraulic loading

$v_{2.5}$ = degree of freedom 2.5 l/min hydraulic loading data

$v_{4.0}$ = degree of freedom 4.0 l/min hydraulic loading data

$$\frac{\text{Lack of Fit}}{\text{Pure Error}} = \frac{0.3373}{0.1336} = 2.52 < F_{0.95}(8,14) = 2.70$$

Therefore, statistically it is possible to use same kinetic expression for 2.5 l/min and 4.0 l/min hydraulic loading for zero order kinetics assumption.

ANOVA 4 Hydraulic loading = 4.0 l/min and first order kinetic assumption

Linearized Arrhenius Model:

$$\ln K = -8063 \left(\frac{1}{T} - \frac{1}{T_0} \right) + (-1.34)$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Square	F
Total (uncorrected)	51.5043	13		
Mean	33.1841	1		
Total (corrected)	18.3202	12		
Regression	8.3984	1	8.3984	9.31
Residual	9.9218	11	0.9020	
Pure Error	7.9832	8	0.9979	
Lack of Fit	1.9386	3	0.6462	0.64

Pure Error Estimate:

$$\begin{aligned} p_E^2 &= (0.80)^2 + (0.458)^2 + (0.72)^2 + (0.19)^2 + (0.74)^2 + (0.88)^2 \\ &\quad + (0.28)^2 + (1.16)^2 + (1.49)^2 + (0.24)^2 + (1.25)^2 = 7.9832 \end{aligned}$$

ANOVA NO. 5 Hydraulic loading = 2.5 l/min and first order kinetic assumption

Linearized Arrhenius Equation

$$\ln K = -13857 \left(\frac{1}{T} - \frac{1}{T_0} \right) - 0.57.$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Square	F
Total (uncorrected)	22.9571	11		
Mean	2.2113	1		
Total (corrected)	20.7458	10		
Regression	17.3758	1	17.3758	46.40
Residual	3.3699	9	0.3744	
Pure Error	1.6779	6	0.2796	
Lack of Fit	1.6919	3	0.5639	2.01

Pure Error Estimation:

$$P_E^2 = (0.19)^2 \times 2 + (0.05)^2 \times 2 + (0.026)^2 \times 2 + (0.625)^2 \times 2 + \\ (0.72)^2 \times 2 + (0.49)^2 \times 2 + (0.23)^2 \times 2 = 1.6779$$

ANOVA NO. 6 (For combined data of 2.5 l/min and 4.0 l/min operation hydraulic loading and first order kinetics)

Linearized Arrhenius Equation:

$$\ln K = -11214 \left(\frac{1}{T} - \frac{1}{T_0} \right) - 0.92606$$

Source of Variation	Sum of Squares	Degrees of Freedom	Mean Square	F
Total (uncorrected)	74.4614	24		
Mean	27.5247	1		
Total (corrected)	46.9367	23		
Regression	28.8633	1	28.8633	35.1342
Residual	18.0734	22	0.8215	
Pure Error	5.2809	14	0.3772	
Lack of Fit	12.7925	8	1.5990	

Pure Error Estimate

$$P_{E_C}^2 = \frac{(7.9832) \times 8 + 1.6779 \times 6}{14} = 5.2809$$

$$\frac{\text{Lack of Fit}}{\text{Pure Error}} = \frac{1.5990}{0.3772} = 4.23 > F_{0.95}(8, 14) \approx 2.70$$