Characteristics of Drug Poisonings Presenting to the Emergency Department: An Electronic Medical Record Database Analysis

Matthew Bell PharmD, ACPR Candidate 01/10/2021

Co-investigators: Dr. Anne Holbrook, Christine Wallace, Dr. Erich Hanel and Dr. Kaitlynn Rigg

Submitted to the Pharmacy Residency Advisory Committee in fulfillment of the requirements for the Certificate of the Ontario Hospital Residency Program (2020-2021)

Acknowledgements

I would like to personally thank Dr. Anne Holbrook for her continued support and contributions to the design, analysis and writing of this project. Without her ongoing dedication this project would not have been possible. I would also like to thank Dr. Gary Foster for his contributions towards the data organization and analysis, Sonia Shiels and Juan Gil for their contributions with the data collection, Dr. Erich Hanel, Dr. Kaitlynn Rigg and Cathy Burger for their contributions to the study design. Lastly, I would like to thank Christine Wallace for her ongoing support of this project who provided the needed guidance throughout the entire process.

Table of	[•] Contents:
----------	------------------------

1.0 Abstract
2.0 Introduction
2.1 Background2
2.2 Study Purpose and Objectives3
3.0 Methods
3.1 Design3
3.2 Inclusion and Exclusion Criteria3
3.3 Data Collection4
3.4 Data Organization6
3.5 Statistical Analysis
3.6 Geographical Heat Maps7
4.0 Results
4.1 Incidence
4.2 Demographics9
4.3 Drugs Involved
4.4 Hospital Outcomes and Resource Utilization12
5.0 Discussion
5.1 Study Strengths
5.2 Study Limitations16
5.3 Implications17
6.0 Conclusion17
Appendix A: Definitions21
Appendix B: Coding Algorithm
Appendix C: ICD-10-CA Codes23
Appendix D: Variables44
Appendix E: Data Collection48
Appendix E: Data Collection48 Appendix F: Data Organization49

1.0 Abstract

<u>Rationale</u>: Drug poisonings are a frequent diagnosis in the Emergency Department (ED), requiring patient management from multiple services. Although there is considerable research detailing the fatal opioid overdose epidemic, little is known about drug poisonings as a whole. Our objective was to describe the drug poisonings seen in the ED at St. Joseph's Hospital Hamilton (SJHH), a large academic urban hospital.

<u>Methods</u>: This study was a retrospective, descriptive study using data abstracted from Dovetale – Epic[®], an electronic medical record for calendar years 2018 through 2020. Patients were identified by ICD-10 drug poisoning codes and data were collected on demographics, drugs involved, hospital management and outcomes to the end of the acute admission. Data were stratified by the intent of drug poisoning.

<u>Results</u>: In total, 2983 drug poisoning visits were included, patient mean age 38.3 years (SD 16.2), 54.7% female, yielding an overall incidence rate of 16.0 drug poisonings/1000 ED visits (8.1 intentional, 6.6 non-intentional and 1.3 unknown). The intentional drug poisoning cohort was younger (mean 36.1 +/-15.7 versus 41.0 +/- 16.6 years) with a higher proportion of females (67.1% versus 42.5%) than the non-intentional cohort. The most prevalent drugs for intentional drug poisonings were antidepressants (26.9%), benzodiazepines (24.9%) and acetaminophen (21.9%) compared to opioids (any opioid, 46.9% including fentanyl, 15.9%, heroin, 11.5%, other opioids, 15.0%) for non-intentional. An antidote was ordered for 26.7% of patients; most commonly N-acetylcysteine for intentional (7.9%) and naloxone for non-intentional (17.1%). The rate of return visit to the ED with a subsequent drug poisoning was 25.9% within a mean follow up of 18.4 months. Mental Health Services (Psychiatric Emergency referral or consult) and Addictions Services consults were ordered in 33.5% and 6.5% of patients respectively. Only 716 (24.0%) of patients were admitted for inpatient care from the ED with an accompanying acute in-hospital mortality rate of 1.0%. The mean length of stay for the initial ED stay and acute hospitalization was 2.2 days (SD 5.8).

<u>Conclusion</u>: This study illustrates a high rate of hospital utilization due to drug poisoning associated with several drugs and suggesting a significant rate of poor outcome, resource utilization and recidivism.

2.0 Introduction

2.1 Background

Drug Poisoning Epidemiology

Drug poisonings are a frequent diagnosis in the United States representing 0.4-2% of all ED visits^{1,2}. The term poisoning is frequently associated with an act of malicious intent, however in medical terminology it is an overarching term that can also be used to describe a drug overdose, an accidental ingestion, or intentional self-harm³. According to the Canadian Coding Standards (CCS), a poisoning is defined as a substance taken incorrectly that results in harm. This definition can include both intentional, taking a drug or substance with the purpose of self-harm, or non-intentional, accidently taking a drug/substance or too much of said drug/substance⁴. These definitions exclude acute intoxications, inebriations, and adverse drug reactions (Appendix A).

Across Canada in 2014, there were 13,438 hospitalizations for a self-inflicted injury, of which 11,564 (86.1%) were a poisoning⁵. In 2012, poisonings accounted for 23.3% of suicides in Canada. Ontario Poison Control (OPC) reported 52,414 calls to the OPC Centre in 2019. Of these, 40% were for a patient in the ED or admitted to a healthcare facility, 1.5% had life threatening symptoms, and 0.1% resulted in death⁶. Locally, the City of Hamilton's Public Health Service provides weekly summary reports on ED visits for drug misuse by incorporating triage data into an epidemiologic surveillance informatics system. At the current rate of ED visits in Hamilton, it is estimated there will be 4732 visits for drug misuse and 1924 visits for overdose in 2021. Hamilton rates non-intentional drug overdose in the top 3 of the most burdensome health outcomes in the city⁷.

Drugs Involved

In the OPC report, 6 of the top 10 classes of medications responsible for consults with them were pain medications (acetaminophen, acetylsalicylic acid, etc.), anxiolytics and sleeping pills, antidepressants, antihistamines, cough and cold and cardiovascular medications⁶. A retrospective chart review completed in the EDs of Montreal characterizing 369 patients who attempted suicide between 2009 and 2010, found similar medications were used in overdose attempts⁸. A retrospective review of hospital discharge abstracts in British Columbia using drug poisoning ICD-10 codes found antiepileptics, sedatives, hypnotics, psychotropics and non-opioid analgesic to be the most common causes of hospitalization from intentional drug poisonings versus narcotics and psychedelics for non-intentional drug poisonings⁹. Opioid poisonings are a particular concern Canada-wide, given their frequency and burden of avoidable accidental death and morbidity. A recent report by the Government of Canada notes an 89% increase in opioid-related deaths in 2020 compared to 2019; of the approximately 17 deaths per day, the majority were non-intentional (96%) and involved fentanyl (82%)¹⁰.

Management of Drug Poisonings

A drug poisoning can be difficult to diagnose, as patients often present with unexplained symptoms complicated by altered mental status and the lack of reliable information. Once diagnosed, and in those who present immediately following a drug poisoning, techniques to decrease the absorption (e.g. decontamination via administration of charcoal) or to counteract the damaging effect of the poison (e.g. antidotes) are utilized. The best practices from the Institute for Safe Medication Practices (ISMP) for targeted medications in acute hospitals recommends that antidotes, reversal agents and rescue agents be readily available along with order sets and protocols to support expeditious emergency administration¹¹. OPC provides guidelines on the recommended antidotes to have stocked in the acute care settings¹². This list contains treatments such as n-acetylcysteine (NAC), which prevents serious hepatotoxicity following acetaminophen overdoses, and naloxone to reverse the respiratory depression of opioid overdose^{13,14}. The NICE self-harm guidelines for the acute management and prevention of recurrence recommends self-harm patients in the ED receive psychosocial assessments, be considered for gastrointestinal decontamination and/or activated charcoal if they present early, have appropriate samples collected (e.g. blood) for analysis and providing clinicians should have access to a poisoning centre for further contact¹⁵. Following initial presentation, patients should be assessed for their risk of repetition of self-harm and underlying mental health disorder to determine if referral, discharge, or admission is appropriate.

St. Joseph's Healthcare Hamilton (SJHH)'s Charlton campus is a 600-bed teaching hospital in downtown Hamilton with an ED that sees 66,000 visits per year and includes the emergency psychiatry specialty services for the region¹⁶. Beginning in December 2017, SJHH transitioned to the electronic medical record (EMR) system, Dovetale, powered by Epic[®].

2.2 Study Purpose and Objectives

A literature review of the MEDLINE database back to 1996 did not reveal any study characterizing drug poisonings presenting to an Ontario hospital. Our objective in this study was to characterize drug poisonings seen initially in the ED at SJHH, a large academic urban hospital.

Primary Objective:

• To describe the incidence of all acute drug poisonings presenting to the ED of St. Joseph's Healthcare Hamilton

Secondary Objectives:

- To characterize the demographics (sex, age, location), comorbidities and intention of patients with a drug poisoning and the drugs involved.
- To describe the management of these drug poisonings e.g., antidotes, psychiatric assessments, and laboratory drug testing, etc.
- To describe the in-hospital outcomes (mortality, intensive care admissions, length of stay and readmission/recidivism)
- To detail the impact of the Covid-19 pandemic on the incidence of drug poisonings

3.0 Methods

3.1 Design

The study was as a descriptive, retrospective case series involving data analysis of the SJHH EMR system, Dovetale - Epic[®].

3.2 Inclusion and Exclusion Criteria

Inclusion Criteria

• Patients 18 years or older seen in the SJHH ED or Urgent Care (UC) diagnosed with a poisoning that was the result of a drug, biologic agent, medication, or a combination of such, between January 1st, 2018, and December 31st, 2020.

Exclusion Criteria:

- Poisoning caused by a non-drug substance
- Patients diagnosed with an adverse drug reaction, an acute intoxication/inebriation or drug poisoning was a result of a medical error
- Drug poisoning occurred in hospital

3.3 Data Collection

We received ethics approval for this study from the Hamilton Integrated Research Ethics Board (Project #12680-C). The Health Data and Information Team searched Dovetale-Epic[®] EMRs of patients registered at triage and seen by a physician in our ED or UC centre from January 1st 2018 to December 31st 2020. From this search, all medical records of ED/UC visits for a drug poisoning were identified if they had at least one ICD-10 poisoning and 'external cause' code assigned to their visit (Figure 1). These codes are assigned to a patient's visit by the Health Data and Information Team if a drug poisoning was a current diagnosis and was treated or contributed to the use of hospital resources. External cause codes are assigned to visits where a diagnosis is made during such visit that is the result of an environmental event or circumstance^{3,17}. In relation to drug poisonings, self-harm is the environmental event/circumstance which can then be described and coded as intentional, non-intentional or unknown intent. For drug poisonings, external cause codes are assigned with a drug poisoning code that provides context to the drug involved in the drug poisoning event. Examples of the diagnostic codes included are shown in Table 1.

The ICD-10 drug poisoning codes are grouped into 15 drug classes which describe the individual drugs or families involved in the drug poisoning (Appendix C-Table 2). The external cause codes which describe the intent of the drug poisoning are categorized into 3 classes (Appendix C-Tables 1)¹⁷. The Canadian Coding Standards provides guidance on how to apply the ICD-10 codes to improve reproducibility and accuracy of documenting poisonings (Appendix B-Figure 1)³. A similar search was done by the Health Data and Information Team to identify all medical records of patients admitted to hospital to an acute medical unit through the ED with a drug poisoning diagnosis. An acute medical unit excludes mental health units, therefore patients admitted to an acute mental health unit from the ED will have only their ED data included in this study.

ICD-10 Code Class	Subclass of Codes	Example
External Cause Codes	Nonintentional (X40-X44)	X42- Non-intentional poisoning by
(3 subclasses of codes) ^a		Narcotics and psychodysleptics
	Intentional (X60-X64)	X62- Intentional Self-Poisoning by
		Narcotics and psychodysleptics
	Unknown Intent (Y10-Y14)	Y12- Unknown Intent of Poisoning
		by Narcotics and psychodysleptics
Drug Poisoning Codes	Nonopioid analgesics, antipyretics	T39.1- 4-aminophenol derivatives
(15 different subclasses of codes) ^a	and antirheumatics (T39)	
	Narcotics and psychodysleptics	T40.4- Fentanyl
	(T40)	
	Psychotropic drugs, not classified	T43.0- tricyclic and tetracyclic
	elsewhere (T43)	antidepressants

Table 1: Examples of codes utilized to identify the drug poisoning population in the ED

^a For a full listing of codes refer to Appendix C: Tables 1 and 2.

Identification of Study Population

Patients diagnosed at SJHH with a drug poisoning as documented in their ED discharge abstracts were identified using ICD-10-CA coding (Appendix C- Tables 1&2). The initial report generated contained additional information with the coding, including the demographics, providers involved in care, discharge location, length of stay, and special care admissions (Appendix E, Table 7).

ICD-10-CA Codes included in each poisoning chart as per the CCS by CIHI

Poisoning Code: Drug, Medication, or biologic substance cause of poisoning	Manifestation Code: The resulting signs, symptoms and disease states	External Cause Code: Intentional, accidental or unknown intention of poisoning	Occurrence: Location of poisoning

Additional Data Collection

From the cohort identified in the previous step, additional data was then collected from these patients' electronic health records. This data included ordered antidotes, referrals, and laboratory drug levels (Appendix E, Table 7).

Review of Collected Information

The two generated reports were merged into a single document with data organized by patient medical record number. The exclusion criteria were then applied utilizing ICD-10 coding (Appendix C, Tables 3-6) and patient demographics.



Deidentification of Data

A unique five-digit numeric code (i.e. study ID) was assigned to each patient in place of their medical record number. If a patient had multiple ED visits on the same day, they were assigned the same study ID (i.e. counted as one unique poisoning incident). If a patient had multiple ED visits on different days, they were assigned the same study ID with an additional alphanumeric character making a 6-digit study ID to identify re-visits. Each visit with a diagnosis of a poisoning, including re-visits, were counted in the analysis as unique poisoning incidents. The document linking the study ID with the medical record number was stored on the secure hospital server.

From the cohort identified by the Health Data and Information Team, additional data were retrieved from two main parts of the EMR: CIHI and non-CIHI data (Appendix E: Data Collection). CIHI data was coded data by the Health Data and Information Team for patients' visits and included the patient demographics, comorbidities, intent of drug poisoning, drugs involved, most responsible physician (MRP) and in-hospital outcomes. We utilized the MRP as a surrogate to understand the services responsible for the care of the patients diagnosed with a drug poisoning during their ED visit and/or admission to an acute medical unit. In all visits where the MRP was a psychiatrist, data collection was limited to the ED. In-hospital outcomes were collected from discharge dispositions and are presented separately for patients discharged from the ED versus an inpatient medical unit.

Non-CIHI data was collected from the patient's medical record during the selected visit; this included mental health, addictions, pharmacy and social work consults, orders for medications on the recommended antidote list, and selected lab orders for drug screens and levels (Appendix D, List 2-4). The collected data were from the National Ambulatory Care Reporting System (NACRS) and Discharge Abstract Database (DAD) which include admissions to ED and acute medical units respectively. Data were not collected from Ontario Mental Health Reporting System (OMHRS), which holds psychiatric unit admissions.

3.4 Data Organization

To prepare the collected data for descriptive analysis, the qualitative data such as the comorbidities, drug poisoning codes and discharge locations were organized into groupings as detailed in Appendix F -Tables 1-8. Drug poisoning codes containing all different possible drug causes were organized into 32 unique groupings based on clinical significance or if the drug was estimated to be involved in greater then 3% of the drug poisonings (Appendix F -Table 2). Using the external cause codes (Appendix F, Table 1), visits identified were organized into non-intentional, intentional, and unknown intent drug poisonings for the analysis and description. If a patient was admitted to an acute medical unit and visits existed for both the ED and inpatient unit, the inpatient codes were given preference over the ED codes. This decision was made with the assumption that there would be more information available to improve the accuracy of the diagnosis and therefore the assigned codes. This applied to all collected data except labs, consults and antidotes, where no preference was given to inpatient versus ED data and both sets of data were used conjointly.

3.5 Statistical Analysis

The incidence rate of acute drug poisonings leading to ED visits was described per 1000 ED visits. To calculate the incidence rate, the number of ED visits for the specified cohort was divided by the total number of patients to register at triage and see a physician in the ED from January 1st, 2018 to December 31st, 2020 (186673 ED visits) and then multiplied by 1000 to give the incidence rate per 1000 ED visits. This rate was further described in relation to the intent of the drug poisoning, opioids, and the date the Covid-19 pandemic was declared (January 1st, 2018 to March 10th, 2020 and March 11th, 2020 to December 31st, 2020, inclusive). For the rates before and after the pandemic was declared, the denominator was reflective of the number of ED visits during these two time periods (141035 and 45638 ED visits, respectively).

Descriptive analysis was completed to describe the demographics, incidence, drug causes, services involved and resource utilization for non-intentional, intentional, and unknown intent drug poisonings. The quantitative outcome measures (continuous data) were described using mean and a respective standard deviation. The qualitative outcome measures (categorical data), which were grouped together as described above, were presented with incidence and respective percentage. The data were analyzed per number of visits. Patients with multiple visits for a drug poisoning would be represented multiple times in the data analysis. For the demographics, the age, number of female patients, and visits where the patient was diagnosed with a selected comorbidity were summed and then divided by the total number of ED visits within that selective cohort (i.e. total drug poisonings, intentional, non-intentional and unknown intent). This mean value was utilized to present the mean data such as average age in our population. If the value was to be presented as a percentage, such as the percent of population that was female, this value was then multiplied by 100. A similar approach was used in the statistical analysis for admission to special care units, most responsible provider (MRP), selected consults, labs, antidotes, and disposition. This approach was also utilized for the drugs involved in the acute drug poisonings, therefore the percentage calculated reflects how many drug poisoning visits involved such drug class. In doing so, it allowed for a single drug poisoning event to involve multiple contributing drugs.

To calculate the percent of acute drug poisoning visits to the ED that were for a subsequent drug poisoning during our study period (referred to as a re-visit), the number of visits for patients in each cohort that had multiple visits were summed. The number of individual patients with multiple visits was then subtracted from the summed visits before being divided by the total number of visits. To present as a percentage this number was then multiplied by 100. For the mean number of visits per patient, the total ED visits for each cohort were divided by the number of unique patients represented in that respective cohort. Patients may be represented in multiple cohorts: 2211 patients in all drug poisonings, 1120 patients in intentional poisoning, 1038 patients in non-intentional and 227 in the unknown intent poisoning cohort. To determine the average follow up time during our study period, the number of days between the ED visit for an acute drug poisoning and the end of our study period (December 31st, 2020) was calculated for each visit. This value was then summed and divided by the total number of ED visits in the respective cohort to give the mean follow up time for each cohort.

For the length of stay, a visit was assigned a minimum value of 1 day if the patient presented to the ED. For each subsequent day a patient was in the ED or admitted to an acute medical unit, the additional days were then tallied. The length of stay for each visit was then the sum of their days in the ED and on a medical unit. The length of stay of all visits within each cohort was then summed and divided by the total number of respective visits to present the mean length of stay.

3.6 Geographical Heat Maps

Forward sortation areas (FSAs) are the first three characters of Canadian postal codes; these were collected for each patient visit and used to create geographical heat maps. These heat maps are designed to highlight the relative prevalence of city ward residence or origin of the patients and suggest where further health interventions may be helpful. Once stratified for the intention of the drug poisoning, the number of the drug poisonings occurring in each FSA was entered into Tableau Public[®] software to create the maps. This data represents the total number of drug poisonings occurring in each FSA for the study period: it does not account for the population of said FSA or rate of drug poisonings. This data was analyzed in two ways: 1) per poisoning visit and 2) per patient. The per patient analysis was completed to account for patients who had multiple drug poisoning visits in the study period, therefore their FSA would be included only for the first drug poisoning visit at our ED. Homeless patients were excluded from the dataset if they did not have an FSA assigned to their visit. Patient's living in a shelter had the FSA of the shelter used in the analysis if assigned to their visit.

4.0 Results

Our study identified 3704 possible drug poisoning visits presenting to SJHH ED (Figure 2). After accounting for duplicate visit entries defined as two visits for the same patient/date/drug, there were 3089 unique drug poisoning visits. A total of 106 visits were excluded; patient's age less than 18 years (66 visits), drug poisoning was a result of a medical error (8 visits), and the drug poisoning occurred in the hospital (32 visits). A total of 2983 unique drug poisoning visits.

Figure 2: ED visits identifed with a diagnosed drug poisoning from January 1st, 2018 to December 31st, 2020



4.1 Incidence

Between January 1st 2018 and December 31st 2020, 2983 drug poisoning visits occurred where a patient was diagnosed with a drug poisoning at our ED, resulting in a rate of 16.0/1000 ED visits. Most poisonings were determined to be intentional (50.5%) with an incidence rate of 8.1/1000 ED visits followed by non-intentional (41.4%) with an incidence rate of 6.6/1000 ED visits and unknown intent (8.1%) with an incidence rate of 1.3/1000 ED visits. After the Covid-19 pandemic was declared, an increase in the rate of all drug poisonings was observed as compared with the time period prior to the pandemic (16.8 versus 15.7) as seen in Table 2. The rate of opioid poisoning diagnosis/1000 ED visits also increased during this time (5.1 versus 4.3).

	All	Intentional	Non-Intentional	Unknown Intent	All Opioids
Pre-COVID (Jan 2018- March 11 th , 2020)	15.73	7.91	6.25	1.57	4.33
Post-COVID (March 12 th , 2020-December 31 st , 2020)	16.76	8.55	7.78	0.44	5.13
Total Study Period	15.97	8.06	6.62	1.30	4.53

Table 2: Rate of Drug Poisonings per 1000 ED Visits from January 1st 2018 to December 31st 2020

4.2 Demographics

The patients' mean age was 38.3 years (SD 16.2 years) and 1632 (54.7%) were female. The intentional drug poisoning cohort was younger (36.2 versus 41.0 years) with a higher proportion of females (67.1% versus 42.5%) compared to the non-intentional drug poisoning cohort. A higher percentage of patients diagnosed with a non-intentional drug poisoning were homeless (10.8%) and had a chronic health condition (12.5%) compared to the patients with an intentional drug poisoning diagnosis, where more patients had a diagnosed mental health disorder (47.4%). Many patients (25.9%) had multiple ED visits with a drug poisoning diagnosis during the study period with a mean follow up of 18.4 months. During the 3-year study period, a patient with an intentional drug poisoning ED visits compared with 1.3 poisoning ED visits for patients with a non-intentional drug poisoning diagnosis. Patient characteristics according to intention of drug poisoning are shown in Table 3.

Table 3: Characteristics of Patients Presenting to the ED with a Drug Poisoning from January 1st 2018 toDecember 31st 2020

	All Poisonings (N=2983)	Intentional (N= 1505, 50.45%)	Non- Intentional (N=1236, 41.43%)	Unknown Intent (N=242, 8.11%)
Demographics				
Age, years, mean (SD)	38.28 (16.15)	36.15 (15.74)	41.02 (16.57)	37.45 (14.01)
Female (%)	1632 (54.71)	1010 (67.11)	525 (42.48)	97 (40.08)
Mental Health Disorder ^a (%)	866 (29.03)	714 (47.44)	121 (9.79)	31 (12.81)
Addiction Disorder ^a (%)	400 (13.41)	210 (13.95)	164 (13.27)	26 (10.74)
Selective Chronic Health Condition ^a (%)	278 (9.32)	102 (6.78)	154 (12.46)	22 (9.09)
Homeless (%)	211 (7.07)	61 (4.05)	133 (10.76)	17 (7.02)
Recidivism				
# Patients to Revisit the ED ^b (%)	772 (25.88)	440 (29.24)	277 (22.41)	55 (22.73)
Mean Number of ED Visits/ Poisoning Patient ^b (SD)	1.35 (4.29)	1.34 (7.83)	1.19 (3.03)	1.07 (3.01)

^o Disorders defined as addiction, mental health or selected chronic health disorders (Chronic pain, HIV, cancer, CVD, diabetes, COPD, Dementia, Kidney Disease and Liver cirrhosis) are listed in Appendix F; Table 13- Comorbidities

^b This includes only visits to the ED with a Drug poisoning diagnosis

Geographical heat maps were created from the collected FSAs. As seen in Figure 3, the L8L forward sortation area which encapsulates the western industrial area of downtown Hamilton was the most common area for a patient diagnosed with a drug poisoning to reside. Patients with ED visits with a diagnosed non-intentional drug poisoning were highly localized in this FSA (133 visits) and its surroundings: L8P (102 visits), L8N (89 visits) and L8H (77 visits). The intentional drug poisoning cohort was more evenly dispersed across Hamilton led by patients residing in the following FSAs: L9C (120 visits), L8L (117 visits), L8H (103 visits) and L8P (102 visits). When looking at the number of patients without an FSA, indicating possible homelessness, a similar result was seen compared to a formal homelessness diagnosis (Table 3); more patients diagnosed with a non-intentional drug poisoning having no FSA (13.6%) versus patients with an intentional drug poisoning diagnosis (2.7%). Similar results were seen when the data were analyzed per patient (Appendix G Table 5 and Figure 1).



Figure 3: Geographical Heat Maps using Forward Sortation Areas of Patients Diagnosed with a(n) (A) Drug Poisonings, (B) Intentional Drug Poisoning and (C) Non-Intentional Drug Poisoning analyzed per visit

4.3 Drugs Involved

Details on the drugs involved in poisonings are shown in Table 4. The most commonly cited drugs were opioids, benzodiazepines, antidepressants, acetaminophen, antiepileptics and antipsychotics. For intentional drug poisonings: antidepressants (26.9%), benzodiazepines (24.9%) and acetaminophen (21.9%) were the most prevalent whereas non-intentional drug poisonings were dominated by opioids (46.9%), primarily fentanyl (15.9%), heroin (11.5%), and other opioids (15.0%). Drug poisonings with more than one drug involved were seen in 38.9% of intentional drug poisonings compared to 20.0% of non-intentional. Overall, there was a mean of 1.5 drugs involved per drug poisoning event.

Drug Class	All Poisonings	Intentional	Non-Intentional	Unknown Intent
Prescription Opioids Excluding Fentanyl	174 (5.83)	97 (6.45)	70 (5.66)	7 (2.89)
Fentanyl	238 (7.98)	26 (1.73)	197 (15.94)	15 (6.20)
Heroin	182 (6.10)	10 (0.66)	142 (11.49)	30 (12.40)
Other Opioids ^a	259 (8.68)	39 (2.59)	185 (14.97)	35 (14.46)
Cocaine	105 (3.52)	30 (1.99)	58 (4.69)	17 (7.02)
Other Psychostimulants ^b	163 (5.46)	51 (3.39)	95 (7.69)	17 (7.02)
Cannabis	61 (2.04)	11 (0.73)	43 (3.48)	7 (2.89)
Psychedelics ^c	87 (2.92)	9 (0.60)	57 (4.61)	21 (8.68)
Acetaminophen	436 (14.62)	329 (21.86)	93 (7.52)	14 (5.79)
Salicylates	33 (1.11)	26 (1.73)	7 (0.57)	0 (0)
NSAIDs	149 (4.09)	122 (8.11)	26 (2.10)	1 (0.41)
Tricyclic Antidepressants	47 (1.58)	41 (2.72)	5 (0.40)	1 (0.41)
Other Antidepressants	449 (15.05)	364 (24.19)	74 (5.99)	11 (4.55)
Benzodiazepines	515 (17.26)	375 (24.92)	112 (9.06)	28 (11.57)
Antiepileptics ^d	337 (11.30)	253 (16.81)	75 (6.07)	9 (3.72)
Antipsychotics	273 (9.15)	210 (13.95)	54 (4.37)	9 (3.72)
Antiallergic and antiemetics ^e	152 (5.10)	118 (7.84)	29 (2.35)	5 (2.07)
Poisonings with more than 1 drug involved	868 (29.10)	585 (38.87)	247 (19.98)	36 (14.88)
Mean number of Drugs involved per poisoning episode (SD)	1.50 (0.97)	1.71 (1.13)	1.31 (0.73)	1.18 (0.50)

Table 4: Classes of Drugs Involved in Diagnosed Drug Poisoning ED Visits from January 1st 2018 to December 31st2020

^a Other opioids include drugs such as tramadol, buprenorphine, pentazocine and paracodinc

^b Other Psychostimulants with Abuse Potential include drugs such as dextroamphetamine, methylphenidate and caffeine

^c Psychedelics include drugs such as Lysergic acid diethylamide, mescaline, and psilocine

^d Antiepileptics includes drugs such as carbamazepine, phenytoin and valproic acid

^e Antiallergic and antiemetics include drugs such as diphenhydramine, dimenhydrinate and cetirizine

4.4 Hospital Outcomes and Resource Utilization

In our data set, 716 (24.0%) patients were admitted for inpatient acute care services from the ED. There were up to 375 patients (12.6%) who were transferred to another acute inpatient care location, which may include our own inpatient mental health services that are not reflected in the 24% of patients admitted. More patients with an intentional drug poisoning diagnosis were transferred to another acute care facility or service, 296 (19.7%),

compared to patients with a non-intentional drug poisoning, 59 (4.8%). In all drug poisonings, 251 (8.4%) patients required an intensive level of care: 104 (3.5%) were admitted to the Intensive Care Unit (ICU), 2 (0.1%) were admitted to a Medical Step-Down Unit (MSDU) and 145 (4.9%) received combined care in each of the ICU and MSDU. In patients diagnosed with an intentional drug poisoning, Psychiatry was responsible for 744 (49.4%) patients and General Internal Medicine for 350 (23.3%) patients, whereas in the majority of patients with a non-intentional drug poisoning diagnosis, Emergency Medicine remained the responsible service (864 patients, 69.9%).

During the acute inpatient non-Mental Health admission, a small percentage received formal consults from Mental Health (4.9%), Addictions (6.5%), Social Work (1.7%) and/or Pharmacy (1.6%). An antidote was ordered from OPC's Recommended Antidote List in 797 (26.7%) patients: N-acetylcysteine was the most prominent in intentional drug poisonings, 119 (7.9%) patients versus naloxone in non-intentional, 211 (17.1%) patients, as seen in Table 5.

During a patient's ED visit and/or admission to an acute medical unit, 31 (1.0%) patients died: 19 (1.5%) with a non-intentional drug poisoning, 7 (0.5%) with an intentional drug poisoning and 5 (2.1%) with an unknown intent of drug poisoning. The mortality rate was higher in patients admitted to a medical unit (3.4%) compared to the patients seen only in the ED (0.3%). The majority of patients, 2332 (78.2%), were discharged home from the ED or their acute medical unit. Overall, more patients with a non-intentional drug poisoning left against medical advice, 111 (9.0%), versus the intentional drug poisoning cohort, 41 (2.7%).

Table 5: Drug Poisoning Characteristics –Outcomes and Resources

	All Poisonings	Intentional	Non-Intentional	Unknown
Resource Utilization				
# Admitted to an acute medical unit (%)	716 (24.00)	414 (27.51)	262 (21.20)	40 (16.53)
Mean Length of Stay (SD), Days ^a	2.24 (5.83)	2.18 (5.31)	2.36 (6.31)	2.03 (6.41)
# Admitted to SCU ^b (%)	251 (8.41)	146 (9.70)	90 (7.28)	15 (6.20)
Most Responsible Medical Service (%)				
General Internal Medicine	629 (21.09)	350 (23.26)	243 (19.66)	36 (14.88)
Psychiatry	853 (28.60)	744 (49.44)	82 (6.63)	27 (11.16)
Critical Care	126 (4.22)	73 (4.85)	44 (3.56)	9 (3.72)
Surgery	4 (0.13)	1 (0.07)	3 (0.24)	0 (0)
Emergency	1371 (45.96)	337 (22.39)	864 (69.90)	170 (70.25)
Medical specialties involved in patient's care (SD)	2.57 (1.53)	2.99 (1.43)	2.15 (1.51)	2.17 (1.50)
Selected Consult Ordered ^g (%)	388 (13.01)	237 (15.75)	121 (9.79)	30 (12.40)
Mental Health ^c	147 (4.93)	128 (8.50)	14 (1.13)	5 (2.07)
Social Work	52 (1.74)	24 (1.59)	22 (1.78)	6 (2.48)
Addictions	193 (6.47)	96 (6.38)	78 (6.31)	19 (7.85)
Pharmacy	47 (1.58)	24 (1.59)	16 (1.29)	7 (2.89)
Selected Antidotes Ordered ^d (%)	797 (26.72)	409 (27.18)	327 (26.46)	61 (25.21)
Activated Charcoal	67 (2.25)	62 (4.11)	3 (0.24)	2 (0.83)
N-Acetylcysteine	150 (5.03)	119 (7.91)	29 (2.35)	2 (0.83)
Naloxone	342 (11.46)	92 (6.11)	211 (17.07)	39 (16.12)
Selected Blood/Urine Drug Testing ^h (%)	1116 (37.41)	644 (42.79)	368 (29.77)	104 (42.98)
Urine Drug Screen	276 (9.25)	148 (9.83)	99 (8.01)	29 (11.98)
Acetaminophen	1055 (35.37)	629 (41.79)	330 (26.70)	96 (39.67)
Salicylate	1046 (35.07)	629 (41.79)	322 (26.05)	95 (39.26)
Ethanol	1050 (35.20)	628 (41.73)	327 (26.46)	95 (39.26)
Other Drug Blood Concentration	111 (3.72)	66 (4.39)	32 (2.59)	13 (5.37)
Disposition From ED Without Medical Adn	nission (%)			
In Hospital Mortality	7 (0.31)	0	4 (0.41)	3 (1.49)
Left Against Medical Advice	142 (6.26)	30 (2.75)	91 (9.34)	21 (10.40)
Transfer to Another Acute Facility or a Speciality Service ^e	225 (9.93)	175 (16.04)	41 (4.21)	9 (4.46)
Discharged Home	1853 (81.74)	868 (79.56)	819 (84.09)	166 (82.18)
Admission to a non-acute centre ^f	40 (1.76)	18 (1.65)	19 (1.95)	3 (1.49)
Disposition From ED/INP with Medical Adr	mission (%)			
In Hospital Mortality	24 (3.35)	7 (1.69)	15 (5.73)	2 (5.00)
Left Against Medical Advice	40 (5.59)	11 (2.66)	20 (7.63)	9 (22.50)
Transfer to Another Acute Facility or a Speciality Service ^e	150 (20.95)	121 (29.23)	18 (6.86)	11 (27.50)
Discharged Home	479 (66.90)	261 (63.04)	200 (76.34)	18 (45.00)
Admission to a non-acute centre ^f	23 (3.21)	14 (3.38)	9 (3.44)	0

* For each drug poisoning encounter a patient may have more than 1 selected consult ordered. If multiple of the same consults were ordered for a patient they were included only once in the analysis.

5.0 Discussion

This is the first study in Ontario describing a large cohort of patients with drug poisonings presenting to an urban academic hospital ED. The number of drug poisonings was larger than expected compared to that estimated by City of Hamilton's Public Health Report⁷. This may be explained by the large number of intentional drug poisonings seen in our cohort that the public health system is not designed to capture. In contrast to a review of Ontario and Alberta discharge abstracts from 2010-2018, which showed that ED visits for non-intentional drug poisonings were nearly twice as common as intentional, we found a higher incidence of intentional versus non-intentional drug poisonings¹⁷. Our results may be biased towards intentional drug poisonings as our centre is the regional mental health facility which includes the emergency psychiatry specialty services for the area. The demographics described in the study aligned with that reported in the other studies completed in British Columbia, United States, and Europe with a younger and predominately female population diagnosed with an intentional drug poisoning and an older, predominately male population diagnosed with a non-intentional drug poisoning^{9,17-20}.

The geographical heat maps identified the western industrial area of downtown as the regions of Hamilton with the highest incidence of drug poisonings. These FSAs do not reflect the highest population density within Hamilton, as those belong to the FSAs immediately southwest of this area²¹. Therefore, our maps indicate these areas with the high incidence of drug poisonings are not only due to the population size but additional factors. Studies have demonstrated that individuals with a lower socioeconomic status are at greater risk of drug poisoning mortality²². Cross-referencing our maps to maps assessing poverty in Hamilton, the areas with a high incidence of acute drug poisoning also have some of the highest poverty rates²³. This is reflected in current Canadian guidance for opioid poisoning patients recommending not only the involvement of medications such as methadone but also non-medical "wrap around" services in an effort to target the social factors that also play a large role in drug poisonings such as housing, education, and employment²⁴.

In the care of the acute drug poisoning patients, our study demonstrated a higher rate of leaving against medical advice, mortality and lower admission rate compared to a similar study conducted in the United States¹⁹. The study completed in the United States, differed from our study as it did not include inpatient admissions and only reported ED data. If comparing only ED mortality rate, then similar rates were seen (0.3 versus 0.1%)¹⁹. The mortality rates reported in the study completed in the United States and ours are much lower (0.3 and 3.4% respectively) than that reported in drug poisonings presenting to EDs in British Columbia (11.6%)⁹. To explain the overall low admission rate in our study (24.0%) compared to that of the United States study (41.2%), it can be hypothesized that either our population was less sick, there is a lower threshold for admission in the United States, or there was a difference in the study populations (e.g. a greater number of intentional poisonings in our study leading to more admissions to psychiatric units which is was not captured/reflected in our data)¹⁹.

The frequent involvement of antidepressants, antipsychotics, and antiepileptics (e.g. valproic acid which is prescribed for various mental health conditions), along with the high incidence of underlying mental health disorders in the intentional poisoning cohort as described in this study and others, points to a need for a larger focus on interventions, both pharmacologic and nonpharmacological, for the prevention of self-harm in this population^{8,9}. In the acute care setting, there is increased urgency as studies suggest 15-25% of patients within this cohort will attempt a subsequent intentional poisoning within one year of their previous attempt²⁵. A Cochrane review found cognitive-behavioural-based psychotherapy (CBT) compared to treatment as usual lead

^a Only includes acute inpatient at our centre; does not include if admitted to another facility or admitted to mental health

^b SCU (Special care units); intensive care unit, medical step-down unit and/or surgical step-down unit

^c Not a full representation of mental health services involvement as psychiatry was also the most responsible physician in many cases as described in Table 3.

^d Medications listed on Ontario Poison Control's recommended antidote list for acute care facilities (Appendix D / List 1).

^e This includes Inpatient care, including specialty services that may be within SJHH (inpatient rehab, inpatient psychiatry and inpatient chronic/complex continuing care), military medical facilities and subacute care where this occurs within acute care hospitals.

^f This includes long-term care homes (24-hour nursing), mental health and/or addiction treatment centres and hospice/palliative care facilities.

to fewer patients repeating self-harm incidents²⁶. A meta-analysis found no benefit in repeat self-harm incidents comparing hospital admission to no admission in all self-harm patients, but a benefit was seen in the intentional self-harm cohort²⁷. Our study demonstrated this high recidivism risk in the intentional cohort, with 29% of patients diagnosed with an intentional poisoning re-visiting the ED during our study period with a subsequent drug poisoning diagnosis.

Patients diagnosed with a non-intentional drug poisoning did not demonstrate as high of a recidivism risk as the intentional cohort, however almost 9% did leave against medical advice (AMA). This rate is higher than that reported in the United States ED, with a leaving AMA rate of 1.2%¹⁹. As opioids were the leading drug class resulting in non-intentional poisonings, 46.9%, this high rate may be in part to the undermanaging of patients' withdrawal symptoms and ensuring patients feel comfortable in the hospital setting. This result may be contributed to by the underutilization of addictions consults in the non-intentional poisoning cohort (6.3% of patients received a formal addiction consult). Our results reflect that there was an unexpected small number of consults ordered which likely underestimates allied health involvement in the care of acute drug poisoning patients. This may be attributed to consults not formerly ordered through the EMR system ie. verbal consults to addictions services in the ED or the expectation of a pharmacist to review home medications. Different strategies need to be investigated to prevent patients from leaving AMA, such as managing patient's withdrawal symptoms in the ED, with and without addictions services, to provide the necessary time to implement other harm reduction strategies.

Optimizing the research into harm reduction strategies for the non-intentional cohort is critical given the predominance of opioids in this population, the high incidence of fentanyl involvement, and the known increasing opioid related mortality in Canada^{28,29}. As demonstrated in our study, the incidence of acute drug poisonings and in particular poisonings as the result of opioids, has increased during the pandemic putting an emphasis on the need for interventions to curb opioid epidemic-related harm. In a study in the United States, only 16.6% of patients received follow up treatment for their opioid use disorder within 90 days of a nonfatal opioid poisoning³⁰. Initiation of opioid maintenance treatment in the ED in patients with opioid use disorder increased patients' involvement in addiction treatment at 30 days compared to outpatient referrals and brief interventions alone³¹. Further studies should investigate the impact of initiating opioid maintenance treatment on the risk of patients leaving AMA and repeating opioid poisonings. One harm reduction strategy in the ED is the distribution of naloxone kits. These are provided by the Public Health Unit to distribute to patients in the ED; across Hamilton from 2018-2019 the Public Health unit and associated sites including the SJHH ED distributed over 13,000 naloxone kits which reportedly lead to 1699 opioid overdoses being reversed³².

5.1 Study Strengths

Our study is the first to report on drug poisoning characteristics presenting to an Ontario ED. This study was completed at a large academic urban hospital that contains the regional psychiatric emergency services allowing for a substantial collection of data on patients diagnosed with intentional and non-intentional drug poisonings. Compared to previous studies that looked at drug poisonings using ICD-10 codes, this study also utilized additional data from the EMR to report on recidivism and the in-patient management of drug poisonings^{8,9}. This data represents the most recent drug poisoning trends in Hamilton and may translate to other urban areas across Ontario. By capturing multiple years of drug poisonings up until December 2020, this study incorporates drug poisoning data both before and after the Covid-19 pandemic.

5.2 Study Limitations

In this study, primary record review of patient EMRs was not completed to verify the accuracy of the assigned ICD-10 drug poisoning codes and other data. However, the method applied uses 'gold standard' CIHI coding procedures and diagnostic extraction methodology and has been shown to have a high degree of accuracy in

algorithms that utilize poisoning ICD-10 codes to identify prescription opioid–related deaths in Canada when compared to coroners' data³³. Also, determining the intent of a drug poisoning can be difficult as demonstrated by the 242 drug poisoning events with an unknown intent. This may limit the validity of these designations. Finally, our results on in hospital outcomes and resource utilization is a truncated data set due to capturing data from only ED visits and admissions to acute care units for drug poisoning and therefore missing data from admissions to acute mental health and other psychiatric units. This led to an underrepresentation on the number of admissions, involvement of psychiatric services and the overall resource utilization involved with acute drug poisonings.

5.3 Implications

Suggested by the high number of acute drug poisonings and relatively low mortality rate, our current practices are effective for the acute management of drug poisonings. Although, not uncommon to our facility, the high recidivism rate and number of patients leaving against medical advice in the intentional and non-intentional cohort respectively suggests an area of improvement in the time after the acute management. Policies and funding should look to target increasing the psychiatric and addictions teams' presence in the ED. Order sets can be designed for the optimal management of patient's withdrawal symptoms to prevent patients leaving before receiving their full spectrum of care. Little is known on what can be done to prevent self-harm events from reoccurring. Questions remain such as what is the optimal way to initiate opioid maintenance treatment in the ED in patients following an opioid poisoning or do intentional self-harm patients benefit from prolonged admissions to psychiatric units? These questions need to be answered before more firm policies can be implemented to improve our care of this large, vulnerable population.

6.0 Conclusion

Our analysis of a large cohort of patients presenting with drug poisoning at an urban ED with emergency Psychiatry services, suggests that drug poisoning is common, caused by multiple drugs, associated with a significant rate of poor outcomes, high resource utilization and has a high rate of recidivism. These results will be useful to inform healthcare provider education, hospital policy, and planning. In addition, research is needed to decrease drug poisoning rates and find cost-effective management strategies to optimize patient outcomes.

References

- Substance Abuse and Mental Health Services Administration (2011). Drug Abuse Warning Network: National estimates of drug-related emergency department visits. DAWN Series D-30, DHHS Publication No. (SMA) 08-4339. Rockville, MD.
- Rui P, Kang K. (2017). National Hospital Ambulatory Medical Care Survey: 2017 Emergency Department Summary Tables. National Center for Health Statistics. Available from:https://www.cdc.gov/nchs/data/nhamcs/web tables/2017 ed web tables-508.pdf.
- 3. Canadian Institute for Health Information. (2018). *Canadian Coding Standards for Version 2018 ICD-10-CA and CCI*. Ottawa, ON.
- 4. Canadian Institute for Health Information. (2018). *Bulletin: Opioid Overdose Coding Direction.* Ottawa, ON: CIHI.
- 5. Skinner, R., McFaull, S., Draca, J., Frechette, M., Kaur, J., Pearson, C., & Thompson, W. (2016). Suicide and self-inflicted injury hospitalizations in Canada (1979 to 2014/15). *Health promotion and chronic disease prevention in Canada : research, policy and practice, 36*(11), 243–251.
- 6. Thompson, M. (2009). *Ontario Poison Centre, Annual Report 2009* (pp. 1-18, Rep.). Toronto, ON: Ontario Poison Control.
- 7. City of Hamilton. (2017). Hamilton Opioid Information System Emergency Department Visits and Hospital Admissions. Retrieved September 09, 2020, from https://www.hamilton.ca/public-health/reporting/hamilton-opioid-information-system-emergency-department-visits-and-hospital.
- 8. Mikhail, A., Tanoli, O., Légaré, G., Dubé, P. A., Habel, Y., Lesage, A., Low, N., Lamarre, S., Singh, S., & Rahme, E. (2019). Over the counter Drugs and other substances Used in Attempted Suicide Presented to Emergency Departments in Montreal, Canada. *Crisis*, *40*(3), 166–175.
- 9. Jiang, A., Smith, J., Rajabali, F., Zheng, A., Purssell, R., and Pike, I. (2018). Patterns in poisoning hospitalizations and deaths in British Columbia, 2008 to 2013. BC Medical Journal, 60 (10); 495-502.
- 10. Special Advisory Committee on the Epidemic of Opioid Overdoses. Opioid and Stimulant-related Harms in Canada. Ottawa: Public Health Agency of Canada; June 2021. https://health-infobase.canada. ca/substance-related-harms/opioids-stimulants
- 11. Institute for Safe Medication Practices (2020). *ISMP Targeted Medication Safety Best Practices for Hospitals*. https://www.ismp.org/guidelines/best-practices-hospitals_
- 12. Ontario Poison Control (2014). Guidelines for Stocking Emergency Antidotes; www.ontariopoisoncentre.ca/pdf/63065-56235%20web%20OPC_Antidote.pdf
- 13. Heard K. J. (2008). Acetylcysteine for acetaminophen poisoning. *The New England journal of medicine*, *359*(3), 285–292. https://doi.org/10.1056/NEJMct0708278
- 14. Smilkstein, M. J., Knapp, G. L., Kulig, K. W., & Rumack, B. H. (1988). Efficacy of oral N-acetylcysteine in the treatment of acetaminophen overdose. Analysis of the national multicenter study (1976 to 1985). *The New England journal of medicine*, *319*(24), 1557–1562.
- 15. National Institute For Health and Care Excellence (2003). Self-harm in over 8s: short-term management and prevention of recurrence. Clinical guideline [CG16]. Updated 2019.
- 16. St Joseph's Healthcare Hamilton. (2014). Emergency Department, Charlton Campus. Retrieved September 09, 2020 from: www.stjoes.ca/hospital-services/emergency-services/emergency-department,-charlton-campus.
- 17. Jiang A, Belton, kL, & Fuselli P (2020). Evidence Summary on the Prevention of Poisoning in Canada. Parachute: Toronto, ON.
- 18. Callanan, V. J., & Davis, M. S. (2012). Gender differences in suicide methods. *Social psychiatry and psychiatric epidemiology*, *47*(6), 857–869.
- 19. Xiang, Y., Zhao, W., Xiang, H., & Smith, G. A. (2012). ED visits for drug-related poisoning in the United States, 2007. *The American Journal of Emergency Medicine*, *30*(2), 293–301.

- 20. European Monitoring Centre for Drugs and Drug Addiction (2016), *Hospital emergency presentations and acute drug toxicity in Europe: update from the Euro-DEN Plus research group and the EMCDDA*, Publications Office of the European Union, Luxembourg.
- 21. City of Hamilton (2017). City of Hamilton Ward Boundaries 2001-2016 Statistics Canada Information. https://map.hamilton.ca/static/pdfs/wardmaps/AllWards_Statistics.pdf.
- 22. Ye, X., Sutherland, J., Henry, B., Tyndall, M., & Kendall, P. (2018). At-a-glance Impact of drug overdoserelated deaths on life expectancy at birth in British Columbia. Aperçu - Impact des décès par surdose de drogue sur l'espérance de vie à la naissance en Colombie-Britannique. *Health promotion and chronic disease prevention in Canada : research, policy and practice, 38*(6), 248–251.
- 23. Community Profiles- Hamilton. (2009). The Social Planning and Research Council of Hamilton. https://www.sprc.hamilton.on.ca/wp-content/uploads/2008/11/Community-Profile-Hamilton-November-2009.pdf
- 24. Government of Canada. Background document: public consultation on strengthening Canada's approach to substance use issues. Published September 2018. Accessed April 14, 2021. https://www.canada.ca/content/dam/ hc-sc/documents/services/substance-use/canadian-drugssubstances-strategy/strengthening-canadaapproach-substance-use-issue/strengthening-canadaapproach-substance-use-issue.pdf
- 25. Owens, D., Horrocks, J., & House, A. (2002). Fatal and non-fatal repetition of self-harm. Systematic review. *The British journal of psychiatry : the journal of mental science*, *181*, 193–199.
- Hawton, K., Witt, K., Salisbury, T., Arensman, E., Gunnell, D., Hazell, P., Townsend, E. and van Heeringen, K. Psychosocial interventions for self-harm in adults. Cochrane Database of Systematic Reviews: 5 (2016).
- 27. Carroll, R., Metcalfe, C., & Gunnell, D. (2014). Hospital management of self-harm patients and risk of repetition: systematic review and meta-analysis. *Journal of affective disorders*, *168*, 476–483.
- 28. Gomes, T., Khuu, W., Martins, D., Tadrous, M., Mamdani, M., and Paterson, J. (2018). Contributions of prescribed and non-prescribed opioids to opioid related deaths: population-based cohort study in Ontario, Canada. *BMJ*; 362 :k3207.
- 29. Gomes T, Kitchen SA, Murray R. (2021). Measuring the Burden of Opioid-Related Mortality in Ontario, Canada, During the COVID-19 Pandemic. *JAMA Open*;4(5):e2112865.
- Kilaru, A. S., Xiong, A., Lowenstein, M., Meisel, Z. F., Perrone, J., Khatri, U., Mitra, N., & Delgado, M. K. (2020). Incidence of Treatment for Opioid Use Disorder Following Nonfatal Overdose in Commercially Insured Patients. *JAMA network open*, *3*(5), e205852.
- D'Onofrio, G., O'Connor, P. G., Pantalon, M. V., Chawarski, M. C., Busch, S. H., Owens, P. H., Bernstein, S. L., & Fiellin, D. A. (2015). Emergency department-initiated buprenorphine/naloxone treatment for opioid dependence: a randomized clinical trial. *JAMA*, *313*(16), 1636–1644.
- City of Hamilton (2019). Hamilton Drug Strategy- 2019 Report to the Community. www.hamilton.ca/sites/default/files/media/browser/2020-02-18/hamilton-drug-strategy-report-tocommunity-2019.pdf.
- 33. Gladstone, E., Smolina, K., Morgan, S. G., Fernandes, K. A., Martins, D., & Gomes, T. (2016). Sensitivity and specificity of administrative mortality data for identifying prescription opioid-related deaths. *CMAJ* : *Canadian Medical Association journal = journal de l'Association medicale canadienne*, 188(4), E67–E72
- 34. Canadian Health Institute of Health Information. (2015). International Statistical Classification of Disease and Related Health Problems. Tenth Revision. World Health Organization revised by CIHI.
- 35. World Health Organization. (1971). International Drug Monitoring: The Role of National Centres. Report of WHO Meeting. Geneva, Switzerland: World Health Organization Technical Report Series, 498.
- 36. Tournier, M., Molimard, M., Cougnard, A., Abouelfath, A., Fourrier, A., & Verdoux, H. (2005). Psychiatric disorders and their comorbidity in subjects with parasuicide by intentional drug overdose: prevalence and gender differences. *Psychiatry research*, *136*(2-3), 93–100.

- 37. Lin, C., Yen, T. H., Juang, Y. Y., Lin, J. L., & Lee, S. H. (2014). Psychiatric comorbidity and its impact on mortality in patients who attempted suicide by paraquat poisoning during 2000-2010. *PloS one*, *9*(11), e112160.
- 38. Boscarino, J. A., Kirchner, H. L., Pitcavage, J. M., Nadipelli, V. R., Ronquest, N. A., Fitzpatrick, M. H., & Han, J. J. (2016). Factors associated with opioid overdose: a 10-year retrospective study of patients in a large integrated health care system. *Substance abuse and rehabilitation*, *7*, 131–141.

Appendices

Appendix A: Definitions

Definitions as per CIHI and WHO^{3,34,35}:

<u>Drug Poisoning</u>- When a substance (drug, medicament, or biological agent) is taken incorrectly and results in harm. A poisoning can be described as a drug overdose, accidental ingestion, or intentional self-harm.

<u>Incorrect use</u>- Wrong drug or dosage given or taken, self-prescribed drug taken in combination with a prescribed drug or taken not as recommended or taking any drug with alcohol.

Intentional poisoning- Purposely self-inflicted or a suicide attempt.

<u>Non-Intentional poisoning</u>- A drug was taken accidentally, too much of a drug was accidentally taken, or the wrong drug was given or taken.

<u>Adverse drug reaction</u>- A substance (drug, medicament, or biological agent) taken correctly as prescribed (correct dose, for therapeutic or prophylactic use) that results in a reaction. This includes allergic reactions, drug interactions or an accumulative effect leading to toxicity.

<u>Acute intoxication or inebriation</u>- Alteration in a person's level of consciousness, cognition, perception, affect or behaviour that resolves with time due to a psychoactive substance.

<u>Acute Medical Unit-</u>Medical units with acute care services where abstracts are created upon discharge for the DAD database. In particular these exclude admissions to mental health units as they are reported to the Ontario Mental Health Reporting System (OMHRS).

Appendix B: Coding Algorithm

This study utilized ICD-10-CA coding for its population identification and parts of the data collection.

- At SJHH, like all other hospitals, ICD-10-CA codes are applied for each admission to patients' discharge summaries in accordance with the Canadian Coding Standards and CIHI standard procedures (Figure 1)³.
- Each drug poisoning is coded with the drug(s) (Appendix C- Table 2), the manifestation of the drug poisoning, the intent of the drug poisoning (Appendix C- Table 1) and the location it occurred.





Appendix C: ICD-10-CA Codes

The following tables define the ICD-10-CA codes with their corresponding descriptions³⁴.

Table 1: ICD-10 Drug Poisoning Codes for external causes of morbidity and mortality by drug or substance

 (Accidental=X40-44, Intentional=X60-X64, and unknown intent=Y10-Y14)

Accidental Poisoning by and exposure to noxious substances			
Medication Class	Code	Examples	
Nonopioid analgesics,	X40	4-aminophenol derivatives	
antipyretics and antirheumatics		Nonsteroidal anti-inflammatory drugs	
		[NSAID]	
		Pyrazolone derivatives	
		Salicylate	
Antiepileptic, sedative-hypnotic,	X41	Antidepressants	
antiparkinsonism and		Barbiturates	
psychotropic drugs, not		Hydantoin derivatives	
elsewhere classified		Iminostilbenes	
		Methaqualone compounds	
		Neuroleptics	
		Psychostimulants	
		Succinimides and oxazolidinediones	
		Tranquillizer	
Narcotics and psychodysleptics	X42	Cannabis (derivatives)	
[hallucinogens], not elsewhere		Cocaine	
classified		Codeine	
		Heroin	
		Lysergide [LSD]	
		Mescaline	
		Methadone	
		Morphine	
		Opium (alkaloids)	
Drugs acting on the autonomic	X43	Parasympatholytics [anticholinergics and	
nervous system		antimuscarinics] and spasmolytics	
		Parasympathomimetics [cholinergics]	
		Sympatholytics [antiadrenergics]	
		Sympathomimetics [adrenergics]	
Other and unspecified drugs,	X44	Agents primarily acting on smooth and	
medicaments and biological		skeletal muscles and the respiratory system	
substance		Anaesthetics (general)(local)	
		Drugs affecting the:	
		-cardiovascular system	
		 gastrointestinal system 	
		Hormones and synthetic substitutes	
		Systemic and haematological agents	
		Systemic antibiotics and other anti-	
		infectives	
		Therapeutic gases	
		Topical preparations	

		Vaccines	
		Water-balance agents and drugs affecting	
		mineral and uric acid metabolism	
Intentional Self-Poisoning by and exposure to noxious substances			
Nonopioid analgesics,	X60	4-aminophenol derivatives	
antipyretics and antirheumatics		Nonsteroidal anti-inflammatory drugs	
		[NSAID]	
		Pyrazolines derivatives	
		Salicylate	
Antiepileptic, sedative-hypnotic,	X61	Antidepressants	
antiparkinsonism and		Barbiturates	
psychotropic drugs, not		Hydantoin derivatives	
elsewhere classified		Iminostilbenes	
		Methagualone compounds	
		Neuroleptics	
		Psychostimulants	
		Succinimides and oxazolidinediones	
		Tranguillizer	
Narcotics and psychodysleptics	X62	Cannabis (derivatives)	
[hallucinogens], not elsewhere		Cocaine	
classified		Codeine	
		Heroin	
		Lysergide [LSD]	
		Mescaline	
		Methadone	
		Morphine	
		Opium (alkaloids)	
Drugs acting on the autonomic	X63	Parasympatholytics [anticholinergics and	
nervous system		antimuscarinics] and spasmolytics	
		Parasympathomimetics [cholinergics]	
		Sympatholytics [antiadrenergics]	
		Sympathomimetics [adrenergics]	
Other and unspecified drugs,	X64	Agents primarily acting on smooth and	
medicaments and biological		skeletal muscles and the respiratory system	
substance		Anaesthetics (general)(local)	
		Drugs affecting the:	
		-cardiovascular system	
		- gastrointestinal system	
		Hormones and synthetic substitutes	
		Systemic and haematological agents	
		Systemic antibiotics and other anti-	
		infectives	
		Therapeutic gases	
		Topical preparations	
		Vaccines	
		Water-balance agents and drugs affecting	
		mineral and uric acid metabolism	
Unknown Intent	of Poisoning by and ex	posure to noxious substances	
	Y10	4-aminophenol derivatives	

Nonopioid analgesics,		Nonsteroidal anti-inflammatory drugs
antipyretics and antirheumatics		[NSAID]
		Pyrazolines derivatives
		Salicylate
Antiepileptic, sedative-hypnotic,	Y11	Antidepressants
antiparkinsonism and		Barbiturates
psychotropic drugs, not		Hydantoin derivatives
elsewhere classified		Iminostilbenes
		Methaqualone compounds
		Neuroleptics
		Psychostimulants
		Succinimides and oxazolidinediones
		Tranquillizer
Narcotics and psychodysleptics	Y12	Cannabis (derivatives)
[hallucinogens], not elsewhere		Cocaine
classified		Codeine
		Heroin
		Lysergide [LSD]
		Mescaline
		Methadone
		Morphine
		Opium (alkaloids)
Drugs acting on the autonomic	Y13	Parasympatholytics [anticholinergics and
nervous system		antimuscarinics] and spasmolytics
		Parasympathomimetics [cholinergics]
		Sympatholytics [antiadrenergics]
		Sympathomimetics [adrenergics]
Other and unspecified drugs,	Y14	Agents primarily acting on smooth and
medicaments and biological		skeletal muscles and the respiratory system
substance		Anaesthetics (general)(local)
		Drugs affecting the:
		-cardiovascular system
		- gastrointestinal system
		Hormones and synthetic substitutes
		Systemic and haematological agents
		Systemic antibiotics and other anti-
		infectives
		Therapeutic gases
		Topical preparations
		Vaccines
		Water-balance agents and drugs affecting
		mineral and uric acid metabolism

Table 2: ICD-10 Poisoning Codes by drugs, medicaments, and biological Substances (T36-T50)

Drug Class (code)	Code	Family Medication Class
Systemic Antibiotics (T36)	T36.0	Penicillins
	T36.1	Cephalosporins and other beta-lactam
		antibiotics
	T36.2	Chloramphenicol group
	T36.3	Macrolides
	T36.4	Tetracyclines
	T36.5	Aminoglycosides
	T36.6	Rifamycins
	T36.7	Antifungal antibiotics, systemically used
	T36.8	Other systemic antibiotics
	T36.9	Systemic antibiotic, unspecified
Other systemic anti-infectives	Т37.0	Sulfonamides
and anti-parasitics (T37)	T37.1	Antimycobacterial drugs
	T37.2	Antimalarials and drugs acting on other blood
		protozoa
	Т37.3	Other antiprotozoal drugs
	Т37.4	Anthelminthics
	T37.5	Antiviral drugs
	Т37.8	Other specified systemic anti-infectives and
		antiparasitics
	Т37.9	Systemic anti-infectives and antiparasitics,
		unspecified
Hormones and their synthetic	T38.0	Glucocorticoids and synthetic analogues
substitutes and antagonists, not	T38.1	Thyroid hormones and substitutes
elsewhere classified (T38)	T38.2	Thyroid hormones and substitutes
	T38.3	Thyroid hormones and substitutes
	T38.4	Oral contraceptives
	T38.5	Other estrogens and progestogens
	T38.6	Antigonadotropins, antiestrogens and
		antiandrogens not elsewhere classified
	T38.7	Androgens and anabolic congeners
	Т38.8	Other and unspecified hormones and their
		synthetic substitutes
	Т38.9	Other and unspecified hormone antagonists
Nonopioid analgesics,	Т39.0	Salicylates
antipyretics and antirheumatics	T39.1	4-Aminophenol derivatives
(T39)	Т39.2	Pyrazoline derivatives
	Т39.3	Other Non-steroidal anti-inflammatory drugs
		(NSAIDs)
	T39.4	Antirheumatics, not elsewhere classified
	Т39.8	Other nonopioid analgesics and antipyretics not
		elsewhere classified
	Т39.9	Nonopioid analgesics, antipyretics and
		antirheumatics, unspecified
Narcotics and pyschodysleptics	T40.0	Opium
(T40)	T40.1	Heroin

	T40.2	Other opioids
		-codeine
		-morphine
	T40.3	Methadone
	T40.4	Other synthetic narcotics
	T40.5	Cocaine
	T40.6	Other and unspecified narcotics
	T40.7	Cannabis (derivatives)
	T40.8	Lysergide (LSD)
	T40.9	Other and unspecified pyschodysleptics
		(hallucinogens) (
		-Mescaline
		-Psilocin
		-Psilocybin
Anaesthetics and therapeutic	T41.0	Inhaled anesthetics
gases (T41)	T41.1	Intravenous anesthetics
	T41.2	Other and unspecified general anesthetics
	T41.3	Local anesthetics
	T41.4	Anesthetics, unspecified
	T41.5	Therapeutic gases
		-carbon dioxide
		-oxygen
Antiepileptic, sedative-hypnotic	T42.0	Hydantoin derivatives
and antiparkinsonism drugs	T42.1	Iminostilbenes
		-carbamazepine
	T42.2	Succinimides and oxazolidinediones
	T42.3	Barbiturates
	T42.4	Benzodiazepines
	T42.5	Mixed antiepileptics not classified elsewhere
	T42.6	Other antiepileptic and sedative-hypnotic drugs
		-valproic acid
		-methaqualone
	T42.7	Antiepileptic and sedative hypnotic drugs,
		unspecified
	T42.8	Antiparkinsonism drugs and other central
		muscle tone depressants
		-amantadine
Psychotropic drugs, not	T43.0	Tricyclic and tetracyclic antidepressants
classified elsewhere (T43)	T43.1	Monoamine-oxidase inhibitor antidepressants
	T43.2	Other and unspecified antidepressants
	T43.3	Phenothiazine antipsychotics and neuroleptics
	T43.4	Butyrophenone and thioxanthene neuroleptics
	T43.5	Other and unspecified antipsychotics and
		neuroleptics
	T43.6	Psychostimulants with abuse potential
	T43.8	Other psychotropic drugs, not elsewhere
		classified
	43.9	Psychotropic drug, unspecified
	T44.0	Anticholinesterase agents

Drugs primarily affecting the	T44.1	Other parasympathomimetics (cholinergics)
autonomic nervous system	T44.2	Ganglionic blocking drugs, not elsewhere
(T44)		classified
	T44.3	Other parasympatholytics and spasmolytics not
		elsewhere classified
		-Papaverine
	T44.4	Predominately alpha-adrenoreceptor agonists,
		not elsewhere classified
	T44.5	Predominately beta-adrenoreceptor agonists,
		not elsewhere classified
	T44.6	Alpha-adrenoreceptor antagonists, not
		elsewhere classified
		-ergot alkaloids
	T44.7	Beta-adrenoreceptor antagonists, not elsewhere
		classified
	T44.8	Centrally acting and adrenergic-neuron-blocking
		agents, not elsewhere classified
	T44.9	Other and unspecified drugs primarily affecting
		the autonomic nervous system
		-drug stimulating both alpha and beta
		adrenoreceptors
Primarily systemic and	T45.0	Antiallergic and antiemetic drugs
hematological agents, not	T45.1	Antineoplastic and immunosuppressive drugs
elsewhere classified (145)		-antineoplastic antibiotics
	745.0	
	145.2	Vitamins not elsewhere classified
	145.3	Enzymes, not elsewhere classified
	145.4	Iron and its compounds
	145.5	Anticoagulants
	145.6	Fibrinolysis-affecting drugs
	145.7	Anticoagulant antagonists, vitamin k and other coagulants
	T45.8	Other primarily systemic and haematological
		agents
		-liver preparations and other antianaemic
		agents
		-Natural blood products
		-Plasma substitute
	T45.9	Primarily systemic and haematological agents,
		unspecified
Agents affecting the	T46.0	Cardiac stimulant glycosides and drugs of similar
cardiovascular system (T46)		action
	T46.1	Calcium channel blockers
	46.2	Other antidysrhythmic drugs not elsewhere
		classified
	T46.3	Coronary vasodilators, not elsewhere classified
		-dipyridamole
	T46.4	Angiotensin converting enzyme inhibitors

	T46.5	Other antihypertensive drugs, not classified
		elsewhere
		-clonidine
	T46.6	Antihyperlipidaemic and antiarteriosclerotic
		drugs
	T46.7	Peripheral vasodilators
	T46.8	Antivaricose drugs, including sclerosing agents
	T46.9	Other and unspecified agents primarily affecting
		the cardiovascular system
Primarily affecting the	T47.0	Histamine H ₂ Receptor Antagonists
gastrointestinal system (T47)	T47.1	Other antacids and anti-gastric secretion drugs
	T47.2	Stimulant laxatives
	T47.3	Saline and osmotic laxatives
	T47.4	Other laxatives
	T47.5	Digestants
	T47.6	Antidiarrheal drugs
	T47.7	Emetics
	T47.8	Other agents primarily affecting the
		gastrointestinal system
	T47.9	Agent primarily affecting the gastrointestinal
	-	systemic, unspecified
Primarily acting on smooth and	T48.0	Oxytocic drugs
skeletal muscles and the	T48.1	Skeletal muscle relaxants [neuromuscular
respiratory system (T48)		blocking agents]
	T48.2	Other and unspecified agents primarily acting on
		muscles
	T48.3	Antitussives
	T48.4	Expectorants
	T48.5	Anti-common cold drugs
	T48.6	Antiasthmatics. not elsewhere classified
		-Salbutamol
	T48.7	Other and unspecified agents primarily acting on
		the respiratory system
Topical agents primarily	T49.0	Local antifungal, anti-infective and anti-
affecting skin and mucous		inflammatory drugs, not elsewhere classified
membranes and by	T49.1	Antipruritics
Ophthalmological,	T49.2	Local astringents and local detergents
otorhinolaryngological and	T49.3	Emollients, demulcents and protectants
dental drugs (T49)	T49.4	Keratolytics, keratoplasties and other hair
		treatment drugs and preparations
	T49.5	Ophthalmological drugs and preparations
		-eye anti-infectives
	T49.6	Otorhinolaryngological drugs and preparations
		-ear, nose and throat anti-infectives
	T49.7	Dental drugs, topically applied
	T49.8	Other topical agents
		-spermicides
	T49.9	Topical agent, unspecified
	T50.0	Mineralocorticoids and their antagonists

Diuretics and other unspecified	T50.1	Loop diuretics
drugs, medicaments and	T50.2	Carbonic-anhydrase inhibitors,
biological substances (T50)		benzothiadiazides and other diuretics
		-acetazolamide
	T50.3	Electrolytic, caloric and water balance agents
		-oral rehydration salts
	T50.4	Drugs affecting uric acid metabolism
	T50.5	Appetite depressants
	T50.6	Antidotes and chelating agents, not classified
		elsewhere
		-Alcohol deterrents
	T50.7	Analeptics and opioid receptor antagonists
	T50.8	Diagnostic agents
	T50.9	Other unspecified drugs, medicaments and
		biological substances
		-acidifying agents
		-alkalizing agents
		-immunoglobulins
		-immunologicals
		-lipotropic drugs
		-parathyroid hormones and derivatives

 Table 3: ICD-10 Codes for toxic effects of non-medical substances (T51-T65)

Drug Class (code)	Code	Family Medication Class
Alcohol (T51)	T51.0	Ethanol
	T51.1	Methanol
	T51.2	2-Propanol
	T51.3	Fuse Oil
	T51.8	Other alcohols
	T51.9	Alcohol, unspecified
Organic Solvents (T52)	T52.0	Petroleum products
	T52.1	Benzene
	T52.2	Homologues of benzene
	T52.3	Glycols
	T52.4	Ketones
	T52.8	Other organic solvents
	T52.9	Organic solvent, unspecified
Halogen Derivatives (T53)	T53.0	Carbon tetrachloride
	T53.1	Chloroform
	T53.2	Trichloroethylene
	T53.3	Tetrachloroethylene
	T53.4	Dichloromethane
	T53.5	Chlorofluorocarbons
	T53.6	Other halogen derivatives of aliphatic
		hydrocarbons
	T53.7	Other halogen derivatives of aromatic
		hydrocarbons
	T53.9	Halogen derivatives of aliphatic and
		aromatic hydrocarbons, unspecified
Corrosive Substances (T54)	T54.0	Phenol
	T54.1	Other corrosive organic compounds
	T54.2	Corrosive acids
	T54.3	Corrosive alkalis
	T54.9	Corrosive substances, unspecified
Toxic Effect of soaps and	T55	
detergents (T55)		
Toxic effect of Metals (156)	156.0	Lead
	156.1	Mercury
	156.2	Chromium
	156.3	Cadmium
	156.4	Copper
	156.5	
	T56.6	Tin
	156./	Beryllium
	156.8	Other metals
	T56.9	Metals, unspecified
Inorganic Substances (T57)	157.0	Arsenic
	T57.1	Phosphorous
	T57.2	Manganese
	T57.3	Hydrogen cyanide

	T57.8	Other specified inorganic substances
	T57.9	Inorganic substances, unspecified
Carbon Monoxide (T58)		
Other gases, fumes and vapours	T59.0	Nitrogen oxides
(T59)	T59.1	Sulfur dioxide
	T59.2	Formaldehyde
	T59.3	Lacrimogenic gas
	T59.4	Chlorine gas
	T59.5	Fluorine gas and hydrogen fluoride
	T59.6	Hydrogen sulfide
	T59.7	Carbon dioxide
	T59.8	Other specified gases, fumes and vapours
	T59.9	Gases, fumes and vapours, unspecified
Pesticides (T60)	T60.0	Organophosphates
	T60.1	Halogenated insecticides
	T60.2	Other and unspecified insecticides
	T60.3	Herbicides and fungicides
	T60.4	Rodenticides
	T60.8	Other pesticides
	T60.9	Pesticide, unspecified
Noxious substances eaten as	T61.0	Ciguatera poisoning
seafood (T61)	T61.1	Scombroid poisoning
	T61.2	Other fish and shellfish poisoning
	T61.8	Toxic effects of other seafoods
	T61.9	Toxic effects of unspecified seafoods
Noxious substances eaten as	T62.0	Mushrooms
food (T62)	T62.1	Berries
	T62.2	Other ingested parts of plants
	T62.8	Other specified noxious substances eaten as
		food
	T62.9	Noxious substances eaten as food,
		unspecified
Contact with Venomous animals	T63.0	Snake venom
(T63)	T63.1	Venom of other reptiles
	T63.2	Venom of scorpions
	T63.3	Venom of spiders
	T63.4	Venom of other arthropods
	T63.5	Toxic effect of contact with fish
	T63.6	Toxic effect of contact with other marine
		animals
	T63.8	Toxic effect of contact with other venomous
		animals
	т63.9	Toxic effect of contact with unspecified
		venomous animals
Aflatoxin and other mycotoxin		
food contaminants (T64)		
Other and unspecified	T65.0	Cyanides
substances (T65)	T65.1	Strychnine
	T65.2	Tobacco and nicotine

T65.3	Nitroderatives of benzene
T65.4	Carbon disulfide
T65.5	Nitroglycerin and other nitric acids
T65.6	Paints and dyes
T65.8	Toxic effect of other specified substances
T65.9	Toxic effect of unspecified substance
Table 4: ICD-10 codes for adverse effects caused by drugs

Drug Class (code)	Code	Family Medication Class
Systemic Antibiotics (Y40)		Penicillins
	Y40.0	
	Y40.1	Cephalosporins and other beta-
		lactam antibiotics
	Y40.2	Chloramphenicol group
	Y40.3	Macrolides
	Y40.4	Tetracyclines
	Y40.5	Aminoglycosides
	Y40.6	Rifamycins
	Y40.7	Antifungal antibiotics,
		systemically used
	Y40.8	Other systemic antibiotics
	Y40.9	Systemic antibiotic, unspecified
Other systemic anti-infectives	Y41.0	Sulfonamides
and anti-parasitics (Y41)	Y41.1	Antimycobacterial drugs
	Y41.2	Antimalarials and drugs acting
		on other blood protozoa
	Y41.3	Other antiprotozoal drugs
	Y41.4	Anthelminthics
	Y41.5	Antiviral drugs
	Y41.8	Other specified systemic anti-
		infectives and antiparasitics
	Y41.9	Systemic anti-infectives and
		antiparasitics, unspecified
Hormones and their synthetic	Y42.0	Glucocorticoids and synthetic
substitutes and antagonists, not		analogues
elsewhere classified (Y42)	Y42.1	Thyroid hormones and
		substitutes
	Y42.2	Thyroid hormones and
		substitutes
	Y42.3	Thyroid hormones and
		substitutes
	Y42.4	Oral contraceptives
	Y42.5	Other estrogens and
		progestogens
	Y42.6	Antigonadotropins,
		antiestrogens and
		antiandrogens not elsewhere
		classified
	Y42.7	Androgens and anabolic
		congeners
	Y42.8	Other and unspecified
		hormones and their synthetic
		substitutes
	Y42.9	Other and unspecified hormone
		antagonists

Primary Systemic Agents (Y43)	Y43.0	Antiallergic and antiemetic
		drugs
	Y43.1	Antineoplastic antimetabolites
	Y43.2	Antineoplastic natural drug
	Y43.3	Other antineoplastic drugs
	Y43.4	Immunosuppressive agents
	Y43.5	Acidifying and alkalizing agents
	Y43.6	Enzymes, not elsewhere
		classified
	Y43.8	Other primarily systemic agents
		not elsewhere classified
	Y43.9	Primarily systemic agent,
		unspecified
Agents primarily affecting blood	Y44.0	Iron preparations and other
constituents (Y44)		anti-hypochromic-anemic
		preparations
	Y44.1	Vitamin B12, folic acid and
		other anti-megaloblastic anemia
		preparations
	Y44.2	Anticoagulants
	Y44.3	Anticoagulant antagonists,
		vitamin K and other coagulants
	Y44.4	Antithrombotic drugs
	Y44.5	Thrombolytic drugs
	Y44.6	Natural blood and blood
		products
	Y44.7	Plasma substitutes
	Y44.9	Other and unspecified agents
		affecting blood constituents
Analgesics, antipyretics and	Y45.0	Opioids and related analgesics
anti-inflammatory drugs (145)	Y45.1	Salicylates
	Y45.2	Propionic acid derivatives
	Y45.3	Other NSAIDs
	Y45.4	Antirheumatics
	Y45.5	4-aminophenol derivatives
	Y45.8	Other analgesics and
		antipyretics
	Y45.9	Analgesic, antipyretics and anti-
		Inflammatory drugs, unspecified
Antiepileptics and	Y46.0	Succinimides
antiparkinsonism drugs	Y46.1	Oxazolidinediones
	Y46.2	Hydantoin derivatives
	Y46.3	Deoxybarbiturates
	Y46.4	Iminostilbenes
	Y46.5	Valproic Acid
	Y46.6	Other and unspecified
		antiepileptics
	Y46.7	Antiparkinsonism drugs
	Y46.8	Antispasticity drugs

Sedatives, hypnotics and	Y47.0	Barbiturates, not elsewhere
antianxiety drugs (Y47)		classified
	Y47.1	Benzodiazepines
	Y47.2	Cloral derivatives
	Y47.3	Paraldehyde
	Y47.4	Bromine compounds
	Y47.5	Mixed sedatives and hypnotics,
		not elsewhere classified
	Y47.8	Other sedatives, hypnotics and
		antianxiety drugs
	Y47.9	Sedative, hypnotic and
		antianxiety drug, unspecified
Anaesthetics and therapeutic	Y48.0	Inhaled anaesthetics
gases (Y48)	Y48.1	Parenteral anesthetics
	Y48.2	Other and unspecified general
		anaesthetics
	Y48.3	Local anaesthetic
	Y48.4	Anaesthetic, unspecified
	Y48.5	Therapeutic gases
Psychotropic drugs not	Y49.0	ТСА
elsewhere classified (Y49)	Y49.1	MAOi antidepressants
	Y49.2	Other and unspecified
		antidepressants
	Y49.3	Phenothiazine antipsychotics
		and neuroleptics
	Y49.4	Butyrophenone and
		thioxanthene neuroleptics
	Y49.5	Other antipsychotics and
		neuroleptics
	Y49.6	Psychodysleptics
	Y49.7	Psychostimulants with abuse
	×40.0	potential
	Y49.8	Other psychotropics drugs
	Y49.9	Psychotropic drug, unspecified
Central nervous stimulants	Y50.0	Analeptics
(150)	Y50.1	Opioid receptor antagonists
	Y50.2	Methylxantnines
	Y50.8	Other CNS stimulants
A 1	Y50.9	CNS, unspecified
Autonomic nervous system	Y51.0	Anticholinesterase agents
urugs (YSI)	Y51.1	Other parasympathomimetics
	Y51.2	
	Y51.3	Other parasympatholytics and spasmolytics
	Y51.4	Predominately alpha-
		adrenoreceptor agonists
	Y51.5	Predominately beta-
		adrenoreceptor agonists

	Y51.6	Alpha-adrenoreceptor
		antagonists
	Y51.7	Beta-adrenoreceptor
		antagonists
	Y51.8	Centrally acting and adrenergic-
		neuron-blocking agents
	Y51.9	Other and unspecified drugs
		affecting the ANS
Cardiovascular agents (Y52)	Y52.0	Cardiac stimulant glycosides
		and drugs of similar action
	Y52.1	Calcium channel blockers
	Y52.2	Other antidysrhythmic drugs
		not elsewhere classified
	Y52.3	Coronary vasodilators, not
		elsewhere classified
		-dipyridamole
	Y52.4	Angiotensin converting enzyme
		inhibitors
	Y52.5	Other antihypertensive drugs,
		not classified elsewhere
		-clonidine
	Y52.6	Antihyperlipidaemic and
		antiarteriosclerotic drugs
	Y52.7	Peripheral vasodilators
	Y52.8	Antivaricose drugs, including
		sclerosing agents
	Y52.9	Other and unspecified agents
		primarily affecting the
		cardiovascular system
Gastrointestinal System (Y53)	Y53.0	Histamine H ₂ Receptor
		Antagonists
	Y53.1	Other antacids and anti-gastric
		secretion drugs
	Y53.2	Stimulant laxatives
	Y53.3	Saline and osmotic laxatives
	Y53.4	Other laxatives
	Y53.5	Digestants
	Y53.6	Antidiarrheal drugs
	Y53.7	Emetics
	Y53.8	Other agents primarily affecting
		the gastrointestinal system
	Y53.9	Agent primarily affecting the
		gastrointestinal systemic,
		unspecified
Water-balance and mineral and	Y54.0	Mineralocorticoids
uric acid metabolism	Y54.1	Mineralocorticoids antagonists
	Y54.2	Carbonic anhydrase inhibitors
	Y54.3	Benzothiadiazide derivatives
	Y54.4	Loop diuretics

	Y54.5	Other diuretics
	Y54.6	Electrolytic, caloric and water
		balance agents
	Y54.7	Agents affecting calcification
	Y54.8	Agents affecting uric acid
		metabolism
	Y54.9	Mineral salts
Smooth and Skeletal muscles	Y55.0	Oxytocic drugs
and the respiratory system	Y55.1	Skeletal muscle relaxants
		[neuromuscular blocking
		agents]
	Y55.2	Other and unspecified agents
		primarily acting on muscles
	Y55.3	Antitussives
	Y55.4	Expectorants
	Y55.5	Anti-common cold drugs
	Y55.6	Antiasthmatics, not elsewhere
		classified
		-Salbutamol
	Y55.7	Other and unspecified agents
		primarily acting on the
		respiratory system
Topical agents primarily	Y56.0	Local antifungal, anti-infective
affecting skin and mucous		and anti-inflammatory drugs,
membranes and by		not elsewhere classified
Ophthalmological,	Y56.1	Antipruritics
otorhinolaryngological and	Y56.2	Local astringents and local
dental drugs (Y56)		detergents
	Y56.3	Emollients, demulcents and
		protectants
	Y56.4	Keratolytics, keratoplasties and
		other hair treatment drugs and
		preparations
	Y56.5	Ophthalmological drugs and
		preparations
		-eye anti-infectives
	Y56.6	Otorhinolaryngological drugs
		and preparations
		-ear, nose and throat anti-
		infectives
	Y56.7	Dental drugs, topically applied
	Y56.8	Other topical agents
		-spermicides
	Y56.9	Topical agent, unspecified
Unspecified drugs (Y57)	Y57.0	Appetite depressants
	Y57.1	Lipotropic drugs
	Y57.2	Antidotes and chelating agents
	Y57.3	Alcohol deterrents
	Y57.4	Pharmaceutical excipients

	Y57.5	X-ray contrast media
	Y57.6	Other diagnostic agents
	Y57.7	Vitamins
	Y57.8	Other drugs
	Y57.9	Drugs, unspecified
Bacterial Vaccines (Y58)	Y58.0	BCG Vaccine
	Y58.1	Typhoid and paratyphoid
		vaccine
	Y58.2	Cholera vaccine
	Y58.3	Plague vaccine
	Y58.4	Tetanus Vaccine
	Y58.5	Diphtheria vaccine
	Y58.6	Pertussis vaccine
	Y58.8	Mixed bacterial vaccines
	Y58.9	Other and unspecified vaccines
Other and unspecified vaccines	Y59.0	Viral vaccines
and biological substances (Y59)	Y59.1	Rickettsia vaccines
	Y59.2	Protozoal vaccines
	Y59.3	Immunoglobulins
	Y59.8	Other specified vaccines and
		biological substances
	Y59.9	Vaccines or biological
		substances, unspecified

 Table 5: ICD-10 Codes for mental and behavioral disorders due to psychoactive substance use

Category	Code	Specifics	
Alcohol (F10)	F10.0	Acute Intoxication	
	F10.1	Harmful Use	
	F10.2	Dependence	
	F10.3	Withdrawal	
	F10.4	Withdrawal with delirium	
	F10.5	Psychotic Disorder	
	F10.6	Amnesia Syndrome	
	F10.7	Residual, late onset, psychotic	
		disorder	
	F10.8	Other mental health and	
		behavioral disorders	
	F10.9	Unspecified mental health and	
		behavior disorders	
Opioids (F11)	F11.0	Acute Intoxication	
,	F11.1	Harmful Use	
	F11.2	Dependence	
	F11.3	Withdrawal	
	F11.4	Withdrawal with delirium	
	F11.5	Psychotic Disorder	
	F11.6	Amnesia Syndrome	
	F11 7	Residual late onset insychotic	
		disorder	
	F11 8	Other mental health and	
	111.0	behavioral disorders	
	F11 9	Unspecified mental health and	
	1 11.5	behavior disorders	
Cannahinoids (E12)	F12.0		
	F12.0	Harmful Lise	
	F12.1	Dependence	
	F12.2	Withdrawal	
	F12.5	Withdrawal with dolirium	
	F12.4	Developtic Dicordor	
	F12.5	Ampasia Sundrama	
	F12.0	Amnesia Syndrome	
	F12.7	Residual, late onset, psychotic	
	512.0	alsorder	
	F12.8	Other mental health and	
		behavioral disorders	
	F12.9	Unspecified mental health and	
		behavior disorders	
Sedatives and Hypnotics (F13)	F13.0	Acute Intoxication	
	F13.1	Harmtul Use	
	F13.2	Dependence	
	F13.3	Withdrawal	
	F13.4	Withdrawal with delirium	
	F13.5	Psychotic Disorder	
	F13.6	Amnesia Syndrome	

	F13.7	Residual, late onset, psychotic	
		disorder	
	F13.8	Other mental health and	
		behavioral disorders	
	F13.9	Unspecified mental health and	
		behavior disorders	
Cocaine (F14)	F14.0	Acute Intoxication	
	F14.1	Harmful Use	
	F14.2	Dependence	
	F14.3	Withdrawal	
	F14.4	Withdrawal with delirium	
	F14.5	Psychotic Disorder	
	F14.6	Amnesia Syndrome	
	F14.7	Residual, late onset, psychotic disorder	
	F14 8	Other mental health and	
	1110	behavioral disorders	
	F14.9	Unspecified mental health and	
		behavior disorders	
Stimulants including cocaine	F15.0	Acute Intoxication	
(F15)	F15.1	Harmful Use	
	F15.2	Dependence	
	F15.3	Withdrawal	
	F15.4	Withdrawal with delirium	
	F15.5	Psychotic Disorder	
	F15.6	Amnesia Syndrome	
	F15.7	Residual, late onset, psychotic	
		disorder	
	F15.8	Other mental health and	
		behavioral disorders	
	F15.9	Unspecified mental health and	
		behavior disorders	
Hallucinogens (F16)	F16.0	Acute Intoxication	
	F16.1	Harmful Use	
	F16.2	Dependence	
	F16.3	Withdrawal	
	F16.4	Withdrawal with delirium	
	F16.5	Psychotic Disorder	
	F16.6	Amnesia Syndrome	
	F16.7	Residual, late onset, psychotic	
		disorder	
	F16.8	Other mental health and	
	516.0	benavioral disorders	
	F16.9	Unspecified mental health and	
		benavior disorders	
	F17.0		
	F17.2		
	+1/.3	withdrawai	

	F17.4	Withdrawal with delirium
	F17.5	Psychotic Disorder
	F17.6	Amnesia Syndrome
	F17.7	Residual, late onset, psychotic
		disorder
	F17.8	Other mental health and
		behavioral disorders
	F17.9	Unspecified mental health and
		behavior disorders
Volatile substances (F18)	F18.0	Acute Intoxication
	F18.1	Harmful Use
	F18.2	Dependence
	F18.3	Withdrawal
	F18.4	Withdrawal with delirium
	F18.5	Psychotic Disorder
	F18.6	Amnesia Syndrome
	F18.7	Residual, late onset, psychotic
		disorder
	F18.8	Other mental health and
		behavioral disorders
	F18.9	Unspecified mental health and
		behavior disorders
Multiple drug use and other	F19.0	Acute Intoxication
psychoactive substances (F19)	F19.1	Harmful Use
	F19.2	Dependence
	F19.3	Withdrawal
	F19.4	Withdrawal with delirium
	F19.5	Psychotic Disorder
	F19.6	Amnesia Syndrome
	F19.7	Residual, late onset, psychotic
		disorder
	F19.8	Other mental health and
		behavioral disorders
	F19.9	Unspecified mental health and
		behavior disorders

Table 6: ICD-10 Codes for Misadventures to patients during surgical and medical care relate to medication poisonings

Category	Code	Specifics
Failure in dosage during surgical	Y63.0	Excessive amount of blood or
or medical care (Y63)		other body fluid given
	Y63.1	Incorrect dilution of fluid used
		during infusion
	Y63.4	Failure in dosage in
		electroshock or insulin-shock
		therapy
	Y63.6	Nonadministration of necessary
		drug, medicament or biological
		substance
	Y63.8	Failure in dosage during other
		surgical and medical care
	Y63.9	Failure in dosage during
		unspecific

Appendix D: Variables

List 1: Antidotes Stocked at SJHH as per OPC Guidance¹²

Antidote	Strength	Route	Indication
Acetylcysteine	200 mg/mL	IV	Acetaminophen and other
(Mucomyst)			hepatotoxins
Atropine sulfate	0.6 mg/mL	IV/IM	Carbamate and
			organophosphate
			insecticides
Botulism A-G Heptavalent	18 mL	IV	Botulism Toxin
Antitoxin			
Bromocriptine	2.5 mg	Oral	Neuroleptic malignant
			syndrome
Calcium chloride	1g/10 mL Syringe	IV	Calcium channel blockers,
			hydrofluoric acid burns
Calcium gluconate	1g/10mL (10%)	IV	Calcium channel blockers,
	0.5%		hydrofluoric acid burns
Calcium gluconate	2.5% gel	Topical	
Cyproheptadine	4 mg	Oral	Serotonin Syndrome
(Periactin)	20	N/	
Dantrolene	20 mg injection	IV	Malignant nypertnermia,
			neuroleptic malignant
Deferencemine meaulate	2 a inication	1) / /1) /	syndrome
(Desferal)	z g injection		Iron
(Desierai)	25 a/50 ml	1)/	Insulin sulfonduroos with
Dextrose	25 g/ 50 IIIL		insulin for BB_CCB
Digoxin immune fah	40 mg vial	IV	Digoxin and other cardiac
(Digifab/Digibind)			glycosides
Dimercaprol (BAL)	100 mg/mL (3 mL)	IM	Acute arsenic, inorganic
			mercury. lead
Esmolol	10 mg/mL	IV	
Ethyl alcohol	100 % (dehydrated)	IV	Methanol, ethylene glycol,
,			or diethylene glycol
Flumazenil	0.1 mg/mL injection	IV	Benzodiazepines
	(5mL)		
Folic acid	5 mg/mL injection	IV	Formaldehyde, methanol,
			methotrexate,
			trimethoprim
Fomepizole	1 mg/mL injection (1.5	IV	Methanol, ethylene glycol
	mL)		
Glucagon	1 mg vial	IM/IV	Beta-blockers
Hydroxocobalamin	5 g/vial	IV	Cyanide
(Cyanokit)			
Idarucizumab (Praxbind)	2.5 g/50 mL injection	IV	Dabigatran
Insulin (regular)	100 units/mL	IV	Beta-blockers, calcium
-			channel blockers
Labetalol	5 mg/mL injection	IV	

L-Carnitine	200 mg/mL injection	IV	Hyperammonemia or valproic acid
Leucovorin	50 mg/mL injection	IV	Formaldehyde, methanol, methotrexate, trimethoprim
Lipid 20% (Intralipid)	1L	IV	Lipid soluble toxin
Methylene Blue	10 mg/mL	IV	Methemoglobinemia
Midazolam	1 mg/mL injection 5 mg/mL injection	IM/IV	
Naloxone (Narcan)	0.4 mg/mL injection	IM/IV	Opioids
Octreotide (Sandostatin)	500 mcg/mL injection	IV	Sulfonylureas, repaglinide and related drugs
Phentolamine	5 mg/mL	IV	Dopamine, epinephrine, norepinephrine and phenylephrine
Physostigmine salicylate	1 mg/mL injection	IM/IV	Anticholinergic syndrome
Phytonadione (Vitamin K)	10 mg/mL injection	IV	Warfarin
Pralidoxime (2PAM)	1 g injection	IV/IM	Organophosphate insecticides
Prothrombin Complex Concentrate (Octaplex)		IV	Warfarin
Pyridoxine (Vit B6)	100 mg/mL injection	IM/IV	Isoniazid (INH), ethylene glycol (cofactor)
Sodium Bicarbonate	1 mEq/mL injection	IV	Tricyclic antidepressants (bolus), cocaine (bolus), salicylates (infusion)
Sodium thiosulfate	25% injection	IV	Cyanide
Thiamine (Vitamin B1)	100 mg/mL injection	IM/IV	Ethanol, ethylene glycol
Tranexamic Acid	100 mg/mL injection	IV	
Additional Agent	Used in Poisonings not o	n the Recommended	OPC Antidote List
			A .111

Activated Charcoal50g /225 mLPOAdsorbent*Protamine is not listed on antidote list but carried in pharmacy

**Dimoval to replace Dimercaprol on formulary due to drug shortages, but for the study period, Dimercaprol was available.

List 2: Selected consults that can be ordered/documented in EPIC[®] relating to acute drug poisonings and mental health

- Concurrent capacity building team inpatient consult
- Psychiatric emergency services (PES) consult
- Psychologist consult
- Consultation liaison psychiatry services inpatient consult
- Behavioral therapy consult
- Consult to mood disorders tertiary mental health
- Consult to Schizophrenia tertiary mental health
- Social Work inpatient consult
- Inpatient consult to clinical pharmacology and toxicology
- Integrated Comprehensive care (ICC) inpatient consult

List 3: Selected laboratory tests that can be ordered/documented in EPIC® relating to drug poisonings

- Drug Screen Panel (emergency)
 - Includes; methamphetamines, cocaine, cannabis, benzodiazepines, tricyclic antidepressants, barbiturates, oxycodone, MDMA (ecstasy), amphetamines, opiates, and methadone
 - $\circ \quad \text{Urine} \quad$
- Drug Screen Faeces
- Drug Screen, hair
- Drug Screen, Urine
 - Includes EDDP-methadone metabolite, amphetamines/MDMA, oxycodone, opiates, benzodiazepines, cocaine, THC metabolite (Cannabis)
- POCT rapid urine drug panel
- Barbiturate Screen
- Phenothiazine Screen, urine
 - Drug Levels available:
 - Acetaminophen level
 - Alcohols and ethylene glycol
 - o Amiodarone level
 - o Amitriptyline level
 - Benzodiazepine, urine, quantitative
 - Carbamazepine level, total
 - Clonazepam level
 - Cocaine, urine, qualitative
 - o Cyclosporine level
 - Dabigatran level
 - o Digoxin level
 - o Doxepin level
 - o Ethanol
 - Ethanol, urine
 - Fentanyl, urine
 - Gabapentin level
 - Haloperidol level
 - Imipramine level
 - Protime-INR
 - XA Heparin standard

- Factor 10 activity
 - Lamotrigine level
 - o Lidocaine level
 - o Lithium level
 - Methotrexate level
 - Nortriptyline level
 - Opiate, urine, qualitative
 - o Phenobarbital level
 - Phencyclidine (PCP), urine
 - Phenytoin level
 - o Primidone level
 - Salicylate level
 - o Sirolimus level
 - Tacrolimus level
 - Theophylline level
 - Topiramate level
 - Valproic acid level

List 4: Selected comorbidities in patients relating to drug poisonings³⁶⁻³⁸

- Mental Health Illness
 - Major Depressive Disorder
 - o Generalized Anxiety Disorder
 - o Agoraphobia
 - Obsessive-compulsive disorder
 - Post-traumatic Stress Disorder
 - o Schizophrenia
 - o Schizoaffective Disorder
 - Bipolar Disorder Type 1
 - o Bipolar Disorder Type 2
 - o Borderline Personality Disorder
 - o Antisocial personality disorder
 - Attention Deficit Hyperactivity Disorder
 - Eating Disorder
- Addiction Disorders
 - Opioid Use Disorder
 - o Cannabis Use Disorder
 - Alcohol Use Disorder
 - Substance Use Disorder
- Miscellaneous disorders previously linked to higher rates of poisonings
 - Chronic Pain
 - Human Immunodeficiency Virus
 - \circ Cancer
 - Cardiovascular Disease (HF, MI, Stroke, PVD)
 - Diabetes (T1DM and T2DM)
 - o COPD
 - o Dementia

Appendix E: Data Collection

Table 1: Variables Collected

	Variable	CIHI Coded Data	Description	Location	
Poisoning	Drug(s) Involved	Yes	ICD-10-CA Coding (Appendix C, Table 2)	Discharge Abstracts	
	Intention	Yes	ICD-10-CA Coding (Appendix C, Table 1)	(DAD/NACR)	
In-hospital Management	Antidotes Administered	No	Ontario Poison Control recommended antidotes and activated charcoal (Appendix D, List 1)	Electronic health record system	
	Consults	No	Teams referred to assist in care (Appendix D, List 2)		
	Providers Specialities Involved in Care	Yes	Medical specialities/disciplines of Providers	Discharge Abstracts (DAD/NACR)	
In-hospital Outcomes	Length of Stay	Yes	Time from admission to discharge (days)	Discharge Abstracts	
	Admission to Special Care Units	Yes	Admission to intensive care unit, medical step-down unit, general internal medicine, or acute mental health, etc.	(DAD/NACR)	
	Discharge Location	Yes	Home, another hospital, long term care, community shelter, or no discharge location (death)		
Laboratory	Urine Drug Levels	No	Toxicology levels measured	Electronic	
Data	Blood Drug Levels	No	during admission (Appendix D, List 3)	health record system	
Demographics	Age	Yes	Years	Discharge	
	Gender	Yes	Female, Male, Non-binary	Abstracts	
	Forward Sortation Area	Yes	First 3 characters of a postal code	(DAD/NACR)	
	Comorbidities	Yes	Medical Conditions of Interest (Appendix D, List 4)		

Appendix F: Data Organization

Table 1: (Grouping	of Intention	Codes
------------	----------	--------------	-------

ICD-10		Study
Code	Details	Grouping
X40	Non-Intentional: Nonopioid analgesics, antipyretics and antirheumatics (NSAIDs, acetaminophen, salicylate, etc.)	Non- intentional
X41	Non-Intentional: Antiepileptic, sedative, hypnotic, antiparkinsonism and psychotropic drugs, not elsewhere classified (antidepressants, neuroepileptics, tranquilizers, stimulants, etc.)	Poisoning
X42	Non-Intentional: Narcotics and psychodysleptics (Cannabis, opioids', cocaine, heroin, LSD, methadone, etc.) [hallucinogens], not elsewhere classified	
X43	Non-Intentional: Drugs acting on the autonomic nervous system (Parasympathomimetics, Sympatholytics, etc.)	
X44	Non-Intentional: Other and unspecified drugs, medicaments and biological substance (Cardio, hormones, anaesthetics, GI, etc.)	
X60	Intentional: Nonopioid analgesics, antipyretics and antirheumatics (NSAIDs, acetaminophen, salicylate, etc.)	Intentional Poisoning
X61	Intentional: Antiepileptic, sedative hypnotic antiparkinsonism and psychotropic drugs, not elsewhere classified (antidepressants, neuroepileptics, tranquilizers, stimulants, etc.)	
X62	Intentional: Narcotics and psychodysleptics (Cannabis, opioids, cocaine, heroin, LSD, methadone, etc.) [hallucinogens], not elsewhere classified	
X63	Intentional: Drugs acting on the autonomic nervous system (Parasympathomimetics, Sympatholytics, etc.)	
X64	Intentional: Other and unspecified drugs, medicaments and biological substance (Cardio, hormones, anaesthetics, GI, etc.)	
Y10	Unknown: Nonopioid analgesics, antipyretics and antirheumatics (NSAIDs, acetaminophen, salicylate, etc.)	Unknown Intent of Poisoning
Y11	Unknown: Antiepileptic, sedative hypnotic antiparkinsonism and psychotropic drugs, not elsewhere classified (antidepressants, neuroepileptics, tranquilizers, stimulants, etc.)	
Y12	Unknown: Narcotics and psychodysleptics (Cannabis, opioids, cocaine, heroin, LSD, methadone, etc.) [hallucinogens], not elsewhere classified	
Y13	Unknown: Drugs acting on the autonomic nervous system (Parasympathomimetics, Sympatholytics, etc.)	
Y14	Unknown: Other and unspecified drugs, medicaments and biological substance (Cardio, hormones, anaesthetics, GI, etc.)	

Table 2: Grouping of Poison Codes

ICD-10 Code	Details	Grouping
36,0	Penicillins	Antimicrobials
36.1	Cephalosporins	
36.3	Macrolides	
36.4	Tetracyclines	
36.9	Systemic antibiotic, unspecified	
37.2	Antimalarials and drugs acting on other blood protozoa	
37.5	Antiviral drugs	
37.8	Other specified systemic anti-infectives and antiparasitics	
38,0	Glucocorticoids and synthetic analogues	Hormones
38.1	Thyroid hormones and substitutes	
38.3	Insulin and oral hypoglycemic [antidiabetic] drugs	
38.4	Oral contraceptives	
38.5	Other estrogens and progestogens	
38.8	Other and unspecified hormones and their synthetic substitutes	
39,0	Salicylates	Salicylates
39.1	4-Aminophenol derivatives	Acetaminophen
39.2	Pyrazoline derivatives	NSAIDs
39.3	Other Non-steroidal anti-inflammatory drugs (NSAIDs)	
39.4	Antirheumatics, not elsewhere classified	Other analgesics
39.8	Other nonopioid analgesics and antipyretics not elsewhere classified	
39.9	Nonopioid analgesics, antipyretics and antirheumatics, unspecified	
40,0	Opium	Other Opioids
40.2	Other opioids not specified elsewhere	
40.28	Other opioid	
40.41	Tramadol	
40.48	Other synthetic opioid	
40.1	Heroin	Heroin
40.,20	Codeine	Prescription
40.21	Morphine	Opioids Excluding
40.22	Hydromorphone	Methadone
40.23	Oxycodone	
40.3	Methadone	Methadone
40,40	fentanyl	Fentanyl
40.5	Cocaine	Cocaine
40.7	Cannabis (derivatives)	Cannabis

40.6	Other and unspecified narcotics	Psychedelics
40.8	Lysergide (LSD)	
40.9	Other and unspecified psychedelics (hallucinogens) (Mescaline, psilocin)	
41.2	Other and unspecified general anesthetics	Anesthetics
41.3	Local anesthetics	
42,0	Hydantoin derivatives	Antiepileptics
42.1	Iminostilbenes (carbamazepine)	
42.2	Succinimides and oxazolidinediones	
42.3	Barbiturates	
42.5	Mixed antiepileptics not classified elsewhere	
42.6	Other antiepileptic and sedative-hypnotic drugs (valproic acid)	
42.7	Antiepileptic and sedative hypnotic drugs, unspecified	
42.4	Benzodiazepines	Benzodiazepines
42.8	Antiparkinsonism drugs and other central muscle tone depressants (amantadine)	Antiparkinsonisms
43,0	Tricyclic and tetracyclic antidepressants	Tricyclic Antidepressants
43.1	Monoamine-oxidase inhibitor antidepressants	Other
43.2	Other and unspecified antidepressants	Antidepressants
43.3	Phenothiazine antipsychotics and neuroleptics	
43.4	Butyrophenone and thioxanthene neuroleptics	Antipsychotics
43.5	Other and unspecified antipsychotics and neuroleptics	
43.6	Psychostimulants with abuse potential	Other Psychostimulants
43.8	Other psychotropic drugs, not elsewhere classified	Other
43.9	Psychotropic drug, unspecified	Psychotropics
44.0	Anticholinesterase agents	Drugs Acting on
44.1	Other parasympathomimetics (cholinergics)	Autonomic
44.2	Ganglionic blocking drugs, not elsewhere classified	
44.3	Other parasympatholytics and spasmolytics not elsewhere classified (papaverine)	-
44.4	Predominately alpha-adrenoreceptor agonists, not elsewhere classified	
44.5	Predominately beta-adrenoreceptor agonists, not elsewhere classified	
44.6	Alpha-adrenoreceptor antagonists, not elsewhere classified (ergots)	1
44.7	Beta-adrenoreceptor antagonists, not elsewhere classified	
44.8	Centrally acting and adrenergic-neuron-blocking agents, not elsewhere classified	
44.9	Other and unspecified drugs primarily affecting the autonomic nervous system (stimulate both alpha and beta)	

T45.0	Antiallergic and antiemetic drugs	Antiemetics and Antiallergics
T45.1	Antineoplastic and immunosuppressive drugs (antineoplastic, cvtarabine)	Hematological Agents
T45.2	Vitamins not elsewhere classified	
T45.3	Enzymes, not elsewhere classified	-
T45.4	Iron and its compounds	-
T45.5	Anticoagulants	1
T45.6	Fibrinolysis-affecting drugs	7
T45.7	Anticoagulant antagonists, vitamin k and other coagulants	
T45.8	Other primarily systemic and haematological agents (natural blood products, plasma substitute, liver preparations)	-
T45.9	Primarily systemic and haematological agents, unspecified	
46,0	Cardiac stimulant glycosides and drugs of similar action	Cardiovascular
46.1	Calcium channel blockers	Drugs
46.2	Other antidysrhythmic drugs not elsewhere classified	
46.3	Coronary vasodilators, not elsewhere classified (dipyridamole)	7
46.4	Angiotensin converting enzyme inhibitors	
46.5	Other antihypertensive drugs, not classified elsewhere (clonidine)	
46.6	Antihyperlipidemics and antiarteriosclerotic drugs	
46.7	Peripheral vasodilators	
46.8	Antivaricose drugs, including sclerosing agents	
46.9	Other and unspecified agents primarily affecting the cardiovascular system	
47,0	Histamine H2 Receptor Antagonists	Gastrointestinal
47.1	Other antacids and anti-gastric secretion drugs	Drugs
47.2	Stimulant laxatives	
47.4	Other laxatives	1
47.5	Digestants	7
47.6	Antidiarrheal drugs	1
47.7	Emetics	1
47.8	Other agents primarily affecting the gastrointestinal system	
48,0	Oxytocic drugs	Drugs affecting
48.1	Skeletal muscle relaxants [neuromuscular blocking agents]	Smooth Muscle
48.2	Other and unspecified agents primarily acting on muscles	
48.3	Antitussives	1
48.4	Expectorants	1
48.5	Anti-common cold drugs	
48.6	Antiasthmatics, not elsewhere classified (Salbutamol)	

48.7	Other and unspecified agents primarily acting on the respiratory	
49,0	Local antifungal, anti-infective and anti-inflammatory drugs, not elsewhere classified	Topical Drugs
49.2	Local astringents and local detergents	
49.4	Keratolytics, keratoplasties and other hair treatment drugs and preparations	
49.6	Otorhinolaryngological drugs and preparations -ear, nose and throat anti-infectives	
49.7	Dental drugs, topically applied	
T50.0	Mineralocorticoids and their antagonists	Diuretics
T50.1	Loop diuretics	1
T50.2	Carbonic-anhydrase inhibitors, benzothiadiazides and other diuretics	
T50.3	Electrolytic, caloric and water balance agents	Other Agents
T50.4	Drugs affecting uric acid metabolism	
T50.5	Appetite depressants	
T50.6	Antidotes and chelating agents, not classified elsewhere	
T50.7	Analeptics and opioid receptor antagonists	
T50.8	Diagnostic agents	
T50.9	Other unspecified drugs, medicaments and biological substances	1
	(acidifying agents, immunoglobulins, immunologicals, lipotropic, PTH)	

NACR			
ICD-10 Code	Description	Grouping	
10	Death after arrival	Died	
72	Died in facility - Excludes MAID and in facility suicide		
3	Left AMA after triage & registration	Left Against Medical Advice	
4	Left AMA after triage & registration & assessment		
5	Left AMA after triage & registration & assessment & treatment		
62	Leave post initial treatment - patient left following registration, further assessment by a service provider and initiation of treatment		
63	Left after triage- patient left the ED at his/her own risk following registration and triage		
64	Left after initial assessment- patient left after registration, triage and further assessment (initial treatment did not occur)		
1	Discharged home, no support services(private dwelling)	Home (with or without	
15	Discharged to place of residence	support, correctional facility, transitional home, etc.)	
16	Home WITH Support/Referral- Discharged to private home with supports from the community at home or referred services		
17	Home WITHOUT Support/Referral- Discharged to private home		
	without supports from the community at home or referred services		
40	Group/Supportive Living- Transfer to assisted living/supportive housing or transitional housing, including shelters, do not have 24- hour nursing care		
90	Correctional Facility- Transfer to jail or halfway house		
6	Admit to reporting facility as inpatient to SCU or Ambulatory care visit functional centre	Transfer to another acute care locations within	
7	Admit to reporting facility as INP to another unit of the reporting facility from the ambulatory care visit functional centre	hospital or between hospitals (can include	
8	Transfer to another acute care facility directly from ambulatory care visit functional centre (includes transfer to another acute care facility with entry through the ED)	Mental Health)	
12	Transfer to surgery		
9	Transfer to another non-acute care facility directly from ambulatory	Transfer to non-acute care	
	care functional centre (stand-alone rehab, mental health)	facilities; LTC, Mental	
30	Residential Care- Transfer to long term care home (24-hour nursing, mental health, or addiction treatment centre)	health or Addiction Facilities	

Table 4: Grouping of Discharge Location from Inpatient Admissions

DAD		
ICD-10	Description	Study Grouping
Code		
7	Died	Died
72	Died in facility	
6	Left AMA	Left Against Medical
61	Absent without leave (AWOL)	Advice
62	Left against medical advice (LAMA)	
65	Did not return from leave	
4	Discharge home with support	Home (with or without
5	Discharge home without support	support, correctional
40	Transfer to group/supportive living	facility, transitional home,
90	Transfer to correctional	c.c.,
3	Transfer to OTHER care (ambulatory, palliative, etc.).	Transfer to another acute
10	Transfer to reporting or another facility for inpatient	care locations within
		hospital or between
		hospitals (can include
		Mental Health)
2	Transfer to Continuing Care	Transfer to non-acute care
30	Transfer to residential care	facilities; LTC, Mental
		health or Addiction
		Facilities

Disease State	Grouping
Major Depressive Disorder, single episode	Mental Health Disorder
Major Depressive Disorder, recurrent episode	
Anxiety Disorders (GAD, mixed, panic)	
Agoraphobia	
Obsessive-compulsive disorder	
Post-traumatic stress disorder (+ reaction to severe stress)	
Schizophrenia	
Schizoaffective	
Persistent mood [affective] disorders	
Unspecified mood [affective] disorder	
Bipolar Disorder Type 1 and 2	
Manic Disorder	
Personality disorders	
Attention Deficit Hyperactivity Disorder	
Eating Disorder	
Homelessness	Homelessness
Alcohol Related Disorders	Addiction Disorder
Opioid Related Disorders	
Cannabis Related Disorders	
Sedative, hypnotic or anxiolytic related disorder	
Cocaine related disorder	
Other stimulant related disorder	
Hallucinagen related disorder	
Other psychoactive substance related disorder	
Chronic Pain	Chronic Health Disorder
Human Immunodeficiency Virus (any related)	
Active Cancer	
Cardiovascular Disease (HF, MI, Stroke, PVD)	
Diabetes	
COPD	
Dementia	
Kidney Disease (Acute kidney + chronic kidney disease)	
Liver cirrhosis	

Epic [®] Procedure Order Display Name	Grouping
NA	No Consult
Psychiatric Emergency Services (PES) consult	Mental Health Consult
Psychologist consult	
Neuropsychology consult	
Consultation Liaison Psychiatry Services inpatient consult	
Behavioral Therapy Consult	
Behavioral Supports Ontario inpatient consult	
Cognitive Behavioral Therapy inpatient consult	
Consult To Mood Disorders tertiary mental health	
Consult To Schizophrenia tertiary mental health	
Social Work inpatient consult	Social Work Consult
PES Social Work Consult	
Integrated Comprehensive Care (ICC) inpatient consult	
	Concurrent Capacity and
Concurrent Capacity Building Team inpatient consult	Addiction Consult
Inpatient Consult to Clinical Pharmacology and Toxicology	CPICs Consult
Pharmacy general consult	Pharmacy Consult

Table 7: Grouping of Admissions to Special Care Units

ICD-10 Code	Description	Grouping
99	No Special Care Unit	No admission to a special
		care unit
45	Coronary Intensive Care Nursing Unit (medical)	Admission to the
		Intensive Care Unit
95	Step-down Surgical Unit	Admission to the Medical
		or Surgical Step-Down
		Unit
30	Combined Medical, Surgical Care Nursing Unit	Admission to both the
		Intensive Care and
		Medical/Surgical Step-
		Down units

Table 8: Grouping of Admitting Physician Services

Service	Groupings
Psychiatry	Psychiatry
Internal Medicine, Cardiology, Nephrology, Gastroenterology, Respirology, Neurology and Urology	Internal Medicine
Critical Care	Critical Care
General, orthopaedic, thoracic and otolaryngology surgery	Surgery
Emergency	No admitting Service

Epic [®] Lab Value Order Name	Grouping
Drug Screen Panel (emergency)	Drug Screen
Drug Screen Faeces	
Drug Screen, hair	
Drug Screen, Urine	
POCT rapid urine drug panel	
Barbiturate Screen	
Phenothiazine Screen, urine	
Acetaminophen level	Acetaminophen
Salicylate level	Salicylate
Alcohols and ethylene glycol	Ethanol
Ethanol	
Ethanol, urine	
Opiate, urine, qualitative	Elicit Substances
Phencyclidine (PCP), urine	
Fentanyl, urine	
Cocaine, urine, qualitative	
Amiodarone level	Therapeutic Drug
Amitriptyline level	
Carbamazepine level, total	
Clonazepam level	
Cyclosporine level	
Dabigatran level	
Digoxin level	
Doxepin level	
Gabapentin level	
Haloperidol level	
Imipramine level	
Lamotrigine level	
Lidocaine level	
Lithium level	
Methotrexate level	
Nortriptyline level	
Phenobarbital level	
Phenytoin level	
Primidone level	
Sirolimus level	
Tacrolimus level	
Theophylline level	

Topiramate level	
Valproic acid level	
Protime-INR	Coagulation
XA Heparin standard	
Factor 10 activity	

Appendix G: Tabulation of All Results

	All Drug	Intentional Drug	Unintentional	Unknown Intent
	Poisonings	Poisonings	Drug Poisonings	of Drug Poisoning
Antimicrobials	27 (0.91)	15 (0.10)	12 (0.97)	0 (0)
Hormones	111 (3.72)	62 (4.12)	45 (3.64)	4 (1.65)
Salicylates	33 (1.11)	26 (1.73)	7 (0.57)	0 (0)
Acetaminophen	436 (14.62)	329 (21.86)	93 (7.52)	14 (5.79)
NSAIDs	149 (4.09)	122 (8.11)	26 (2.10)	1 (0.41)
Other non-opioid	5 (0.17)	3 (0.20)	2 (0.16)	0 (0)
analgesics				
Fentanyl	238 (7.98)	26 (1.73)	197 (15.94)	15 (6.20)
Heroin	182 (6.10)	10 (0.66)	142 (11.49)	30 (12.40)
Methadone	21 (0.70)	5 (0.33)	15 (1.21)	1 (0.41)
Prescription	174 (5.83)	97 (6.45)	70 (5.66)	7 (2.89)
Narcotics				
excluding fentanyl				
and methadone				
Other opioids	259 (8.68)	39 (2.59)	185 (14.97)	35 (14.46)
Cocaine	105 (3.52)	30 (1.99)	58 (4.69)	17 (7.02)
Cannabis	61 (2.04)	11 (0.73)	43 (3.48)	7 (2.89)
Psychedelics	87 (2.92)	9 (0.60)	57 (4.61)	21 (8.68)
Anaesthetics	6 (0.20)	2 (0.13)	3 (0.24)	1 (0.41)
Benzodiazepines	515 (17.26)	375 (24.92)	112 (9.06)	28 (11.57)
Antiparkinsonisms	52 (1.74)	28 (1.86)	19 (1.54)	5 (2.07)
Antiepileptics	337 (11.30)	253 (16.81)	75 (6.07)	9 (3.72)
Other	163 (5.46)	51 (3.39)	95 (7.69)	17 (7.02)
Psychostimulants				
Tricyclic	47 (1.58)	41 (2.72)	5 (0.40)	1 (0.41)
antidepressants				
Other	449 (15.05)	364 (24.19)	74 (5.99)	11 (4.55)
antidepressants				- (>
Antipsychotics	273 (9.15)	210 (13.95)	54 (4.37)	9 (3.72)
Other	25 (0.84)	9 (0.60)	14 (1.13)	2 (0.83)
Psychotropics	() () () ()			
Drugs primarily	102 (3.42)	63 (4.19)	33 (2.67)	6 (2.48)
affecting the				
autonomic				
nervous system	452 (5.40)	110 (7.04)	20 (2 25)	5 (2.07)
Antiallergic and	152 (5.10)	118 (7.84)	29 (2.35)	5 (2.07)
antiemetic	20 (4.27)	24/4 50)	42 (4.05)	1 (0.44)
Hematologic	38 (1.27)	24 (1.59)	13 (1.05)	1 (0.41)
Agents		E4 (2 E0)	20 (2 26)	2 (1 24)
Druge	85 (2.88)	54 (3.59)	28 (2.20)	3 (1.24)
Gastrointectinal	25 /1 17)	24 (1 50)	0 (0 72)	2 (0 02)
Drugs	33 (1.17)	24 (1.35)	5 (0.75)	2 (0.03)

 Table 1: Classes of Drugs Involved in Diagnosed Drug Poisonings (% per visit)

Drugs affecting	39 (1.30)	30 (1.99)	8 (0.65)	1 (0.41)
smooth muscles				
Topical Drugs	16 (0.54)	7 (0.47)	9 (0.73)	0 (0)
Diuretics	15 (0.50)	8 (0.53)	7 (0.57)	0 (0)
Other drugs	147 (4.93)	58 (3.86)	60 (4.85)	29 (11.98)

 Table 2: Selected Lab Tests Ordered in Diagnosed Drug Poisonings (% per visit)

	All Drug	Intentional Drug	Unintentional	Unknown Intent
	Poisonings	Poisonings	Drug Poisonings	of Drug Poisoning
Drug Screen	276 (9.25)	148 (9.83)	99 (8.01)	29 (11.98)
Acetaminophen	1055 (35.37)	629 (41.79)	330 (26.70)	96 (39.67)
Salicylate	1046 (35.07)	629 (41.79)	322 (26.05)	95 (39.26)
Ethanol	1050 (35.20)	628 (41.73)	327 (26.46)	95 (39.26)
Illicit Drug	9 (0.30)	0	5 (0.40)	4 (1.65)
Therapeutic Drug	111 (3.72)	66 (4.39)	32 (2.59)	13 (5.37)
Coagulation	214 (7.17)	113 (7.51)	80 (6.47)	21 (8.68)

 Table 3: Selected Antidotes Ordered in Diagnosed Drug Poisonings (% per visit)

	All Drug	Intentional Drug	Unintentional	Unknown Intent
	Poisonings	Poisonings	Drug Poisonings	of Drug Poisoning
Acetylcysteine (Mucomvst)	150 (5.03)	119 (7.91)	29 (2.35)	2 (0.83)
Activated Charcoal	67 (2.25)	62 (4.12)	3 (0.24)	2 (0.83)
Atropine sulfate	3 (0.10)	2 (0.13)	1 (0.08)	0
Botulism A-G	0	0	0	0
Heptavalent				
Antitoxin				
Bromocriptine	1 (0.03)	1 (0.07)	0	0
Calcium chloride	83 (2.78)	37 (2.46)	40 (3.24)	6 (2.48)
or gluconate				
Cyproheptadine	2 (0.07)	2 (0.13)	0	0
(Periactin)				
Dantrolene	0	0	0	0
Deferoxamine	0	0	0	0
mesylate				
(Desteral)	427 (4 50)	66 (4.20)	C2 (5.40)	0 (2 24)
Dextrose	137 (4.59)	66 (4.38)	63 (5.10)	8 (3.31)
Digoxin immune	2 (0.07)	1 (0.07)	1 (0.08)	0
Digitah (Digihind)				
Dimercanrol (BAL)	0	0	0	0
Esmolol	1 (0.03)	0	1 (0.8)	0
Ethyl alcohol	1 (0:03)	0	0	0
Flumazenil	2 (0 07)	0	2 (0 16)	0
Folic acid	12 (0.40)	4 (0.27)	6 (0.49)	2 (0.83)
Fomepizole	5 (0.17)	3 (0.20)	2 (0.16)	0
Glucagon	17 (0.57)	12 (0.80)	5 (0.40)	0
Hydroxocobalamin	0	0	0	0
(Cyanokit)				
Idarucizumab	0	0	0	0
(Praxbind)				
Insulin (regular)	30 (1.00)	12 (0.80)	15 (1.21)	3 (1.24)
Labetalol	14 (0.47)	3 (0.20)	9 (0.73)	2 (0.83)
L-Carnitine	1 (0.03)	1 (0.07)	0	0
Leucovorin	2 (0.07)	1 (0.07)	1 (0.08)	0
Lipid 20%	3 (0.10)	3 (0.20)	0	0
(Intralipid)				
Methylene Blue	0	0	0	0
Midazolam	0	0	0	0
Naloxone (Narcan)	342 (11.46)	92 (6.11)	211 (17.07)	39 (16.12)
Uctreotide	7 (0.23)	2 (0.13)	4 (0.32)	1 (0.41)
(Sandostatin)	0	0		0
Physostiamine	0	0	0	0
salicylate	U	U		U
salicylate				

Phytonadione	34 (1.14)	14 (0.93)	19 (1.54)	1 (0.41)
(Vitamin K)				
Pralidoxime	0	0	0	0
(2PAM)				
Prothrombin	0	0	0	0
Complex				
Concentrate				
(Octaplex)				
Pyridoxine (Vit B6)	2 (0.07)	1 (0.07)	1 (0.08)	0
Sodium	0	0	0	0
Bicarbonate				
Sodium	0	0	0	0
thiosulfate				
Thiamine (Vitamin	235 (7.88)	141 (0.37)	81 (6.55)	13 (5.37)
B1)				
Tranexamic Acid	7 (0.23)	5 (0.33)	2 (0.16)	0

 Table 4: Forward Sortation Area as described Per Drug Poisoning Visit

Forward Sortation	All Drug	Intentional Drug	Unintentional	Unknown Intent
Area	Poisonings	Poisonings	Drug Poisonings	of Drug Poisoning
B1K	1	1	0	0
H2E	1	1	0	0
LOC	1	1	0	0
LOG	1	0	1	0
LOP	1	0	1	0
LOR	88	52	33	3
LOS	1	1	0	0
L1N	1	1	0	0
L2E	2	2	0	0
L2H	1	0	1	0
L2M	1	1	0	0
L2N	1	1	0	0
L2R	1	0	1	0
L2S	2	1	0	1
L3B	3	2	1	0
L3C	2	2	0	0
L3M	17	12	4	1
L3W	1	1	0	0
L3Y	1	0	0	1
L4L	1	1	0	0
L4S	2	0	2	0
L5A	1	1	0	0
L5E	1	1	0	0
L5H	1	0	0	1
L5J	0	0	0	0
L5K	1	1	0	0
L5N	1	1	0	0
L6A	2	2	0	0
L6H	3	2	1	0
L6J	3	0	3	0
L7A	1	1	0	0
L7G	1	0	1	0
L7H	1	0	0	1
L7L	3	3	0	0
L7M	6	2	2	2
L7N	2	1	1	0
L7P	6	4	2	0
L7S	2	1	1	0

L7T	3	1	2	0
L8B	14	10	3	1
L8E	105	56	40	9
L8G	70	34	29	7
L8H	198	103	77	18
L8J	59	34	20	5
L8K	127	65	56	6
L8L	271	117	133	21
L8M	138	63	61	14
L8N	197	98	89	10
L8P	223	102	102	19
L8Q	0	0	0	0
L8R	153	73	60	20
L8S	100	59	33	8
L8T	65	37	21	7
L8U	0	0	0	0
L8V	99	56	38	5
L8W	72	38	27	7
L9A	107	74	24	9
L9B	60	37	20	3
L9C	199	120	64	15
L9G	58	42	14	2
L9H	79	44	32	3
L9K	37	25	12	0
L9N	2	1	1	0
L9P	1	1	0	0
L9T	4	2	2	0
L9W	3	1	2	0
L9Y	1	1	0	0
M1B	3	2	1	0
M1J	1	0	0	1
M1V	2	2	0	0
M2J	1	0	1	0
M2R	1	1	0	0
M4E	1	1	0	0
M4M	0	0	0	0
M5A	2	2	0	0
M5B	1	0	1	0
M5J	1	0	1	0
M5Z	0	0	0	0
M8W	2	2	0	0

65

Total	2983	1505	1236	242
BLANK	237	41	168	28
N5Z	1	0	1	0
V2N	1	0	1	0
T5G	1	0	0	1
T1K	1	0	1	0
S4R	1	0	0	1
P9N	1	0	1	0
N9G	1	1	0	0
N6A	1	1	0	0
N5Y	2	0	1	1
N5A	1	0	1	0
N4N	1	0	0	1
N4B	1	0	1	0
N3Y	1	0	0	1
N3W	18	8	10	0
N3V	1	1	0	0
N3T	4	3	1	0
N3S	7	6	0	1
N3R	2 8	2	5	1
N3P	2	1 0	2	0
N3I	1	1	0	0
N2S	1	1 0	1	0
N2R	1	1	1 0	0
	2	1	1	0
		1	0	0
		1	0	0
		1	0	0
NIG			0	0
N1A	6	3	2	1
NOK	1	1	0	0
NOJ	2	2	0	0
NOG	1	0	1	0
NOE	6	3	2	1
NOC	1	0	1	0
NOB	7	5	2	0
NOA	29	14	10	5
M9R	2	0	2	0
М9С	1	1	0	0

Table 5: Forward Sortation	n Area as descr	ibed Per Patient
----------------------------	-----------------	------------------

Forward Sortation	All Drug	Intentional Drug	Unintentional	Unknown Intent
Area	Poisonings	Poisonings	Drug Poisonings	of Drug Poisoning
B1K	1	0	1	0
H2E	1	0	1	0
LOC	1	0	1	0
LOG	1	1	0	0
LOP	1	1	0	0
LOR	79	31	45	3
LOS	1	0	1	0
L1N	1	0	1	0
L2E	2	0	2	0
L2H	1	1	0	0
L2M	1	0	1	0
L2N	1	0	1	0
L2R	1	1	0	0
L2S	2	0	1	1
L3B	3	1	2	0
L3C	2	0	2	0
L3M	13	4	9	0
L3W	1	0	1	0
L3Y	1	0	0	1
L4L	1	0	1	0
L4S	2	2	0	0
L5A	1	0	1	0
L5E	1	0	1	0
L5H	1	0	0	1
L5J	0	0	0	0
L5K	1	0	1	0
L5N	1	0	1	0
L6A	2	0	2	0
L6H	2	1	1	0
L6J	2	2	0	0
L7A	1	0	1	0
L7G	1	1	0	0
L7H	1	0	0	1
L7L	3	0	3	0
L7M	5	2	2	1
L7N	2	1	1	0
L7P	4	2	2	0
L7S	1	1	0	0

L7T	3	2	1	0
L8B	13	3	10	0
L8E	95	40	49	6
L8G	56	26	25	5
L8H	147	69	68	10
L8J	55	19	32	4
L8K	99	47	49	3
L8L	211	118	77	16
L8M	91	44	41	6
L8N	132	79	47	6
L8P	164	82	71	11
L8Q	0	0	0	0
L8R	79	45	24	10
L8S	71	28	37	6
L8T	51	16	29	6
L8U	0	0	0	0
L8V	73	35	34	4
L8W	61	25	32	4
L9A	77	21	50	6
L9B	50	18	30	2
L9C	138	56	74	8
L9G	42	13	27	2
L9H	63	31	30	2
L9K	30	10	20	0
L9N	2	1	1	0
L9P	1	0	1	0
L9T	4	2	2	0
L9W	3	2	1	0
L9Y	1	0	1	0
M1B	1	1	0	0
M1J	1	0	0	1
M1V	1	0	1	0
M2J	1	1	0	0
M2R	1	0	1	0
M4E	1	0	1	0
M4M	0	0	0	0
M5A	1	0	1	0
M5B	1	1	0	0
M5J	1	1	0	0
M5Z	0	0	0	0
M8W	1	0	1	0

Total	2211	1038	1021	152
BLANK	131	104	13	14
N5Z	1	1	0	0
V2N	1	1	0	0
T5G	1	0	0	1
Т1К	1	1	0	0
S4R	1	0	0	1
P9N	1	1	0	0
N9G	1	0	1	0
N6A	0	0	0	0
N5Y	2	1	0	1
N5A	1	1	0	0
N4N	1	0	0	1
N4B	1	1	0	0
N3Y	1	0	0	1
N3W	16	9	7	0
N3V	1	0	1	0
N3T	<u>л</u>	1	3	0
N3S	7	n	6	1
N3R	7	5	1	1
N3P	2	2	1 0	0
N3I	1	1 0	1	0
N25		1	1	0
	1	1	1	0
	2	1	1	0
		0	1	0
		0	1	0
NIH		0	1	0
NIG		0	1	0
N1A	6	2	3	1
NOK	1	0	1	0
NOJ	2	0	2	0
NOG	1	1	0	0
NOE	6	2	3	1
NOC	1	1	0	0
NOB	6	2	4	0
NOA	25	10	12	3
M9R	2	2	0	0
М9С	1	0	1	0


Figure 1: Geographical Heat Map for All Drug Poisonings Analyzed Per Patient