

Prescription Review Program

2017 Annual Report

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Prescription Review Program Overview

The Prescription Review Program (PRP) is an education-based program of the College of Physicians and Surgeons of Saskatchewan (CPSS) that monitors medications with known misuse, abuse and diversion potential for possible inappropriate prescribing by physicians, and possible in appropriate use by patients. The list of medications monitored by the PRP are listed in the CPSS Regulatory Bylaw 18.1 (Appendix A) as well as in Appendix B (*Prescription Review Program Monitored Medications*). In addition to this, Bylaw 18.1 outlines the requirements for prescribing these medications.

The PRP alerts physicians of possible inappropriate prescribing or of inappropriate use of PRP medications by their patients most commonly by letter. Depending on the situation, the PRP staff may contact the physician directly by phone. The PRP provides supportive and/or educational information, as well as recommendations to physicians in order to encourage appropriate prescribing practices. In some cases, physicians are required to provide explanations for their prescribing of medications to which the PRP applies. After reviewing a physician's reply, the PRP may make additional recommendations regarding best practices, to improve patient outcomes or reduce the possibility of inappropriate use of medications.

Staffing and Workflow

The staffing at the PRP for 2017 included:

- An administrative assistant (who also supports the Opioid Agonist Therapy Program Manager);
- A PRP Analyst (a registered pharmacy technician hired in August 2016);
- A part-time contract pharmacist providing up to two days of service a week (started September 1, 2017);
- The Program Manager (a pharmacist).

The main role of the administrative assistant is to process and prepare all correspondence to physicians, to identify instances of double doctoring each month, to assist in the monitoring of patients profiles, and to support the PRP team with administrative tasks. This role is divided between the PRP and the Opioid Agonist Therapy Program, and as such, the administrative assistant also is responsible to coordinate the approval of methadone exemption through Health Canada.

The main role of the Analyst is to monitor and review patient medication profiles, generate reports related to specific medication use or specific prescribers, and identify possible areas of concern. The Analyst is also the lead on law enforcement engagement, as well as receiving and responding to public information related to the possible diversion of PRP medications.

The main role of the contract pharmacist is to assist with responding to physician correspondence, and other PRP duties as assigned.

The main role of the Program Manager is to guide the work of the PRP and support physicians through education and recommendations related to the prescribing of PRP medications. The Program Manager also works with various stakeholder groups (e.g. FNIHB, NIHB, SCPP, SRNA, Ministry of Health, and many others) to help optimize prescribing and address prescription drug abuse in Saskatchewan.

There are numerous other roles, responsibilities and activities performed by the PRP team and they vary significantly from day-to-day, to month-to-month depending on Program demands, such as requests from Legal Counsel or the Quality of Care Department, the coordination of educational events, or stakeholder requests.

Generally, the day-to-day activities of the PRP for the period of this report can be summarized as follows:



Throughout 2017, the workflow and work processes continued to be refined, altered and optimized. The PRP continues to focus on striving for a paperless workflow process and further optimized their use of the electronic document management software.

Prescription Review Program Letters

There are four categories of letters most commonly sent to physicians by the PRP: Double Doctor, Explain, Alert, and Response. These are defined in the blue box on the right-hand side of the page, and the letter counts are in the table below.

Letter Counts for 2017

Letter Type	# sent out in 2017
Double Doctor	6002
Explain/Alert (1 st contact)	528
2 nd Request	40
Response/Recommendations	288
Law Enforcement Requests	79

Monthly computer generated **double doctor letters** are mailed out to alert physicians that a patient to whom they have prescribed has received a prescription for a PRP medications from at least two other prescribers. The reporting program is able to identify prescribers working at the same clinic address, based on the information provided to the College by individual physicians. Of note, the program cannot identify physicians providing services to federal corrections facilities, nor patients receiving medications from an in-hospital pharmacy.

Alert letters are also sent to physicians as a result of calls received by the PRP staff from individuals (sometimes anonymously) providing information that someone (who has been prescribed PRP medications) may possibly be misusing and/or diverting their medication. The PRP does not suggest in those letters that the physician cease prescribing to the patient. Rather, the PRP recommends that the prescriber put safeguards in place, such as treatment agreements, current random urine drug testing or surprise tablet counts in order to prevent prescription drug misuse or diversion. Alert letters will also be sent to physicians when the PRP receives information from law enforcement about the misuse or diversion of medications.

TYPES OF LETTERS

Alert – sent when the patient is identified as potentially misusing his/her meds (e.g. early refills, law enforcement investigation, information from public/HCP of misuse or diversion).

Double Doctor – sent when a patient received PRP meds from 3 or more physicians, at 3 different practice site addresses in a calendar month. These are system generated letters.

Explain – letters sent to physicians to get their rationale for prescribing (e.g. provide the medical indication and dosing) .

Law Enforcement Request – when a patient medication profile is provided to law enforcement for the purpose of an active investigation.

Prescription – letters to physicians regarding Bylaws 17.1 and 18.1 related to legibility and PRP requirements for a valid prescription.

Response/Recommendations – PRP Manager's response to a physician's explain letter response. These often contain recommendations and recommended resources.

Explain letters can be sent for a variety of reasons, but are always in reference to possible inappropriate or suboptimal prescribing. Common triggers that result in an explain letter include, but are not limited to:

- Double doctoring for an extended period of time (i.e. multiple months)
- A pattern of early refills
- Chronic use of benzodiazepines
- The combined use of a benzodiazepine and opioid
- The concurrent use of two benzodiazepines
- The concurrent use of two opioids
- Prescribing of large quantities of immediate-release opioids repeatedly with/without the use of a sustained-release preparation
- Prescribing of opioids and benzodiazepines for patients concurrently receiving opioid agonist therapy
- Patients with a history of unexpected urine drug screen results
- Large quantities of tablets being dispensed regularly
- Use of brand name preparations when a generic is available
- Specific medications very infrequently used (e.g. Demerol, Talwin, phenobarbital)

Once the physician provides a response to an Explain letter, the PRP can make an assessment of the appropriateness of the prescribing and provide recommendations for possible medication changes or general medication management, such as random urine drug screens, random pill counts, treatment agreements, or other approaches in the Response/Recommendations letter sent back to the physician.

Highlights of PRP Activities for 2017

For the activities below, the PRP team member(s) who attended are denoted by: PM – PRP Pharmacist Manager; PA – PRP Analyst; AA – PRP Administrative Assistant

Below are the PRP activities that occurred in 2017 in relation to education to various stakeholder groups, partnerships and collaborations with various stakeholder groups, as well as educational events attended.

Educational Outreach

- Saskatchewan International Physician Practice Assessment (SIPPA) PM
 - o Presentation Title: What is the PRP?
 - o February 10, 2017; June 8, 2017; September 22, 2017
 - Saskatoon, Saskatchewan
- FNIHB Elder Meeting PM
 - o Presentation Title: What is the PRP?
 - o Ministry of Justice also in attendance
 - o March 23, 2017
 - Saskatoon, Saskatchewan
- Presentation to Saskatoon Tribal Council PM
 - o Presentation Title: What is the PRP?
 - o April 28, 2017
 - Saskatoon, Saskatchewan
- Law Enforcement Educational Event PM, PA, AA
 - o Presentation Title: PRP Meets Saskatchewan's Law Enforcement
 - In attendance: RCMP, City Police, Ministry of Health, Ministry of Justice, RCMP Academy
 - Presented via WebEx to a live audience
 - o June 7, 2017
 - CPSS, Saskatoon, Saskatchewan
- Community Navigator Workshop PM, PA
 - Presentation Title: Prescription Drug Abuse
 - o July 25, 2017
 - Regina, Saskatchewan
- No Future in a Flatline Saskatchewan Fentanyl & Opioid Seminar PM
 - o Presentation Title: Prescription Review Program
 - Objective: To provide a high level overview of various law enforcement and justice community-based initiatives within the province of Saskatchewan that target abuse or illegal use of opioids. Presenters included representatives from policing, corrections, the Crown Attorney's Office along with government ministries and professional governance bodies.
 - o Presented via WebEx to a live audience

- Target audience: Front line policing, Community Based Organizations, Health partners, Ministry of Justice, Canada Border Services Agency
- September 19, 2017
 - RCMP Heritage Centre in Regina, Saskatchewan
- The College of Physicians and Surgeons of Saskatchewan presents the Opioid Substitution Therapy Conference Passion. Hope. Recovery. PM, PA, AA
 - Target Audience: Physicians, pharmacists, nurses, pharmacy technicians, counselors and other individuals working with patients undergoing opioid substitution therapy
 - 176 registrants (90 of which were physicians); 17 presentation topics; 15 speakers
 - o April 29 & 30, 2017
 - Saskatoon Inn, Saskatoon, Saskatchewan
- Current Options for Managing Pain and Addiction Conference (COMPAC) PM, PA
 - The target audience of healthcare providers included: family physicians, physician specialists, medical residents, nurse practitioners, nurses, pharmacists, physical and occupational therapists, psychologists, dentists, social workers and pharmacy technicians.
 - Combined innovative education sessions with interprofessional networking opportunities to address complexities of care for people with acute, chronic, and recurrent pain, and/or substance use disorders.
 - o 252 registrants; 38 speakers
 - o October 27 & 28, 2017
 - o Sheraton, Saskatoon, Saskatchewan
- National Civil Forfeiture Executive Committee Meeting PM & Field Officer from SCPP
 - Presentation explaining the role of both Colleges, what the PRP is, and possible opportunities for collaboration
 - o November 11, 2017
 - Regina, Saskatchewan
- Presentation to Saskatoon HIV/AIDS, Health Canada, and FNIHB PM
 - Presentation Objective: Increase audience's ability to provide clients accurate information about Suboxone alternative treatment and advocate on their behalf when they suspect overprescribing and/or diversion.
 - o November 14, 2017
 - Saskatoon, Saskatchewan

Partnerships and Collaborative Efforts

- Meeting with Minster of Justice Gord Wyant and Deputy Minister Dale McFee PM
 - Meeting Objective: To discuss the effects of fentanyl in Saskatchewan and the collaboration of PRP with the Ministry of Justice.
 - o February 13, 2017
 - Saskatoon, Saskatchewan
- Meeting with Missinipi Broadcasting Corporation PM
 - o Nap Gardiner (MBC) & Katy Windl (NIHB) were present on the conference call
 - Meeting Objective: To discuss how MBC is proposing to help the College of Physicians and Surgeons of Saskatchewan educate the aboriginal public on impact of misuse and diversion of prescription drugs of prescription drugs.
 - o April 27, 2017
- Canadian Community Epidemiology Network on Drug Use (CCENDU) Meeting of the National Network – PM
 - o Meeting Objectives:
 - Identify emerging alcohol and drug use trends in order to identify possible evidence-informed responses
 - Identify and prioritize opportunities to increase linkages and capacity across and beyond the network
 - Build and reflect on activities of the past year to inform the 2017-2018
 CCENDU work plan
 - o May 18 & 19, 2017
 - Montreal, QC (Delta Hotels by Marriott)
 **Funded by CCENDU
- Saskatchewan Regional Prescription Drug Abuse Coordinating Meeting FNIHB-SK PDA Meeting - PM
 - o May 26, 2017
 - Saskatoon, Saskatchewan (Correctional Learning & Development Centre)
- Collaboration with Dr. Luke Terret & Jeff Elder of the Saskatoon Health Region PM,
 PA
 - Saskatoon Emergency Department Opioid Use Study
 - The PRP Analyst provided de-identified data that included physicianspecific opioid prescribing data (fentanyl, hydromorphone, morphine, Percocet, Tylenol #3 and all generics) prescribed pre- and postguideline implementation in the Saskatoon ERs
 - Results have been submitted to the Canadian Journal of Emergency Medicine for 2018
 - o July to August 2017
- medSask Advisory Board Meeting PM
 - o August 24, 2017
 - Saskatoon, Saskatchewan (University of Saskatchewan)

Saskatchewan Regional Prescription Drug Abuse Coordinating Meeting - FNIHB-SK PDA Meeting - PM

- o October 4 & 5, 2017
 - Regina, Saskatchewan (First Nations University of Canada)

Pharmacy Technician Society of Saskatchewan (PTSS) conference – PA

- Presentation Title: What is the PRP and what does the Analyst do?
 - Objective: Discuss how pharmacies can collaborate with PRP to make patients and communities safer.
- Presented to 50+ technician/assistants from hospital/community pharmacies
- o October 21, 2017
 - Travelodge, Saskatoon, Saskatchewan

Northern Alcohol Strategy Event – Changing the Story

- Multidisciplinary event provided over six sessions (each session was repeated once to facilitate attendance).
- The object of the education session was to provide the community with evidence-based cross sector tools on mentoring, education and management of alcohol and opioid use disorders, on the premise that when communities address the issue, every sector is positively impacted.
- 3 speakers: Leslie Molnar (Clinical Social Work/Therapist), Dr. Ashok
 Krishnamurthy (MD), and Irene Njoroge (advanced practical nurse) all from the
 Women's College Hospital (Ontario)
- Topics included: mentoring, education, and clinical tools for addiction, primary care management of alcohol and opioid use disorders, management of alcohol and opioid use disorders in the community and in the workplace
- 192 registrants (including physicians, pharmacists, nurses, business management representatives, law enforcement, and members of the public)
- o October 23 to 26, 2017
 - Eagle Point Resort, La Ronge, Saskatchewan

Government of Saskatchewan Coordinated Health Stakeholders - Problematic Use of Opioids Meeting – PM

- Meeting purpose: The Ministry of Health organized a stakeholders meeting on the subject of problematic use and misuse of opioids. Most provinces and territories including Saskatchewan have agreed to contribute to reporting on a range of activities that are underway in their respective jurisdiction.
- o February 8, 2017; September 13, 2017
 - Regina, Saskatchewan

Meeting with Moose Jaw Police Service – PM, PA

- Meeting purpose: Networking opportunity with Chief Bourassa and the senior team. The discussion included how PRP and MJPS can collaborate on addressing diversion and trafficking investigations in the community. Also discussed future educational sessions for their internal staff on prescription drug abuse in Saskatchewan.
- July 25, 2017
 - Moose Jaw, Saskatchewan

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- Meeting: Addressing barriers to opioid agonist treatments in Saskatchewan PM & OATP Manager
 - Meeting purpose: addressing barriers to treatment for opioid use disorder in order to increase access to buprenorphine/naloxone combination.
 - Attendees: CPSS, U of S, Northern Medical Services, FNIHB office of population & public health, Controlled substances directorate – opioid response team – Health Canada
 - o November 17, 2017
 - Saskatoon, Saskatchewan

Attendance at Education Events

- Pharmacist Information Session: Substitution Therapy for Opioid Use Disorder The Role of Suboxone (buprenorphine/naloxone)- PM
 - Coordinated by Indivior in collaboration with CPDPP (request made after prescriber session was organized)
 - Speaker: Brett Baumback
 - o February 1, 2017
 - Saskatoon, Saskatchewan
- Presentation by Dr. Murray Opdahl on Medical Cannabis PM
 - o March 2, 2017
 - Saskatoon, Saskatchewan
- Webinar: Fentanyl and Its Chemical Cousins: Abuse Patterns, Surveillance, and Treatment – PM
 - o March 7, 2017
- Pharmacy Association of Saskatchewan Annual Conference PM
 - o April 5 to 7, 2017
 - Saskatoon, Saskatchewan
- Call to Action: The time is right for a Provincial Pain Strategy (webinar) PM
 - Event Description: An educational webinar to learn more about Pain Management, the Opioid Crisis and strategies to support improved pain services. The SRNA Pain Management Professional Practice Group has collaborated with key stakeholders to develop a provincial pain strategy framework. Join our moderated panel session to hear about national and provincial pain initiatives, the impact of the opioid crisis and the experiences of individuals living with pain. The time is right to get involved and provide input into this important topic!
 - o May 31, 2017
- Canadian Pharmacy Association Annual (CPhA) Conference PM
 - o June 1 to 5, 2017

- Quebec City, Quebec
- Federation of Medical Regulatory Authorities of Canada (FMRAC) Annual Meeting and Conference - PM
 - o June 10 to 12, 2017
 - Winnipeg, Manitoba
- Webinar: Physician Opioid Prescribing Understanding, Identifying and Re-mediating Problematic Behaviors - PM
 - o June 13, 2017
- Webinar: Opioids in the Acute Care Setting: Safety is Within Our Reach PM
 - o July 11, 2017
- Dialogue to Action on Discussing Substance Use with Women PM & Dr. Karen Shaw
 - o Hosted by: Centre of Excellence for Women's Health
 - o September 21, 2017
 - Saskatoon, Saskatchewan
- Webinar: Street Fentanyl and Its Analogues What Pharmacists Need to Know PM
 - September 28, 2017
- Webinar: Street Fentanyl and Its Analogues What Pharmacists Need to Know PM
 - September 28, 2017
- CADTH Lecture Canada's Opioid Crisis: The Changing Reality Between Exam Rooms and Ivory Towers; Presented by Dr. Hakique Virani PM
 - o October 12, 2017
- Webinar: Braiding together Indigenous wellness, trauma- and gender-informed approaches in the substance use field- PM
 - o October 19, 2017
- Perspective on Substance Use and Recovery PA
 - Hosted by: Break the Barrier
 - o 200+ attendees
 - Panelist included physicians, law enforcement, pharmacist, addictions counsellor, and social worker
 - o November 13, 2017
 - Asher Auditorium, City Hospital, Saskatoon
- Webinar: Pharmacists facing the opioid crisis with evidence and experience PM
 - o November 15, 2017
- RxFiles Presentation to CPSS Staff
 - o Topic: Opioids in Chronic Non-cancer Pain
 - o December 22, 2017

In addition to the above listed activities, the PRP continues to collaborate with the College of Pharmacy Professionals (mainly through Lori Postnikoff, field officer) to identify apparent inappropriate dispensing of PRP drugs. The PRP also continued its work with the National Advisory Council on Prescription Drug Misuse in partnership with the Canadian Centre on Substance Abuse, a comprehensive 10 year pan-Canadian strategy, First Do No Harm: Responding to Canada's Prescription Drug Crisis which was released in March 2013. The strategy highlights the actions required to address the harm associated with the misuse of prescription drugs in Canada in the areas of prevention, education, treatment, monitoring and surveillance and enforcement. The PRP also continues to work with NIHB, the FNIHB Prescription Drug Abuse Saskatchewan group, both the Ministry of Health and Ministry of Justice (including Corrections, Policing and the Chief Coroner's Office), the Provincial Lab, College of Dental Surgeons, Saskatchewan Registered Nurses Association, CCENDU (PRP manages the CCENDU Saskatchewan Facebook page), and CRISM.

Other PRP activities of note that occurred in 2017 include:

- Collaboration with the Saskatchewan Medical Association (SMA) to establish criteria for the Medical Services Branch (MSB) Chronic Pain Billing Code
- Participation with the Current Options for Managing Pain and Addiction Conference (COMPAC) Planning Committee & Curriculum Sub-Committee
- Collaboration with the Ministry of Justice Fentanyl Opioid Overdose Task Force (Cory Lerat, Kait Quinn)
- Collaboration with eHealth to redesign MicroStrategies to meet the needs of PRP
- Collaboration with the Safer Communities and Neighbourhoods (SCAN) Unit
- Collaboration with Dr. Nathaniel Osgood Professor, Department of Computer Science;
 Associate Faculty, Department of Community Health & Epidemiology; Associate Faculty,
 Bioengineering Division on health informatics and analysis of the PRP data
- Initiated Project ECHO (Extension for Community Healthcare Outcomes) work with Dr Susan Tupper and Dr Cathy Jeffery
 - Project ECHO links expert specialist teams at an academic hub with primary care clinicians in local communities.
- Collaboration with College of Physicians and Surgeons of British Columbia (CPSBC)
 Prescription Monitoring Program by spending time at the Program in Vancouver comparing processes and approaches to monitoring
- Participated in the national consultation on the Section 56 methadone exemption with Health Canada
- Continued with the development and growth of the saskpainaddiction.com website

SPEP Students

For the first time, the PRP accepted two 4th year pharmacy SPEP students on their specialty rotations in 2017. Below is a quick overview of their work and experience at PRP. An overview of their experience was provide in DocTalk Volume 4, Issue 1, 2017 available: http://www.cps.sk.ca/iMIS/Documents/Newsletters/DOCTALVol4_Issue1.pdf

1. Lin Wang

- o Offsite experiences that included shadowing different health professionals
 - o Kirsty (pharmacist), Mayfair Pharmacy
 - Dr. R McAllister & Dr. M Markentin, Methadone Clinic on 20th
 - Methadone for Addiction
 - Naloxone training, Mayfair Pharmacy
 - o Dr. M Opdahl
 - Methadone for Pain
 - o Dr. L Pawluck
 - Methadone for Addiction
 - o Christi (RN), Brief Detox Centre
- Presentation
 - o Experience at CPSS
- o Projects
 - Opioid prescribing checklist for chronic pain

2. Arielle Sherman

- o Offsite experiences that included shadowing different health professionals
 - o Dr. R McAllister & Dr. M Markentin, Methadone Clinic on 20th
 - Methadone for addiction
 - Kim Roblin (Addictions Counsellor)
 - o Kirsty (pharmacist), Mayfair Pharmacy
 - o Dr. P Butt, Mayfair Pharmacy
 - o Christi (RN), Brief Detox Centre
- o Presentation
 - Experience at CPSS
- Projects
 - o Opioid myths and facts, Fentanyl patch return tool box
 - DocTalk article: 2C-B and Other Illicit Drugs Surfacing Today

PRP Medication Use in Saskatchewan for 2017 Trends and Insight

An overview of the PRP medications prescribed and dispensed in Saskatchewan are available in Appendices C through F. Dispensing quantities from 2013 to 2017 were provided to allow for a comparison and to identify possible trends.

Stimulants (Methylphenidate)

• Concerta use has gradually increased from 2013 to 2017, while all other forms of methylphenidate have seen a general toward trend. It would appear that prescribers are opting for the Concerta formulation more frequently. While it is assumed that the Concerta formulation is less likely to be abused, law enforcement has indicated that this is not the case. Due to the increasing use of methylphenidate, the PRP will focus on ADD/ADHD and the use of stimulants in 2018.

Opioids

- The fentanyl injection, available as a 50mcg/mL vial showed a peak, or increased use, in 2016, but has dropped significantly in 2017. Fentanyl patches (all strengths) continued to show a decline in use. The concerns around illicit fentanyl and subsequent overdoses, along with the significant attention the media is giving to this topic, may have affected the use of prescription fentanyl.
- The **hydromorphone** 2mg/mL injection continues to be frequently dispensed. The oral immediate-release hydromorphone all saw a slight decrease in use, with the exception of the lowest strength of 1 mg. For the sustained-release preparations (Hydromorph Contin), the higher strengths of 12, 18, 24 and 30mg all decreased very slightly. Conversely, the lower strengths of 3, 4.5, 6 and 9mg all saw slight increases. Overall, the hydromorphone dispensed, in morphine equivalents saw a decrease for the first time since 2013 (however the level still exceeds that of 2015 and previous years).
- The morphine 10mg/mL and 15 mg/mL injection saw a decline in use from 2016, but the use of the 50mg/mL preparation increased very slightly. However, the use of the oral syrup increased for both the 1mg/mL and the 5mg/mL strength. In terms of oral preparations, the use of 5mg immediate-release morphine has steadily increased since 2013, whereas the 10mg strength has decreased. The sustained-release preparations for morphine (both 12-hour and 24-hour preparations) generally saw a decrease in use with a few exceptions: M-Eslon 100mg and Kadian 10mg.
- The use of the Oxycodone immediate-release preparations across all three strengths continues to remain rather consistent. The use of the sustained-release product, OxyNEO, has continued to decrease. However, the prescribing of other preparations has resulted in an increase in the morphine equivalent of oxycodone in dispensed in Saskatchewan in 2017.

Acetaminophen/codeine combination products have seen very consistent use over the
five years, with the 30mg codeine containing product (e.g. Tylenol #3) being prescribed
more commonly than the 20mg and 60mg codeine-containing preparations (Tylenol #2
and Tylenol #4).

Gabapentin

• The 300mg capsule continues to be the most commonly used preparation, and the prescribing of gabapentin continues to increase in Saskatchewan. Gabapentin is a common drug of abuse, and due to its growing use, the PRP will also focus educational efforts toward this medication in 2018.

Benzodiazepines

• The most commonly dispensed benzodiazepines in 2017 was **clonazepam** (specifically the 0.5mg strength) followed closely by **lorazepam** (specifically the 1mg strength). With the exception of clonazepam and lorazepam, the quantity of all other benzodiazepines listed decreased slightly in 2017 (alprazolam, diazepam, oxazepam, and temazepam).

Opioid-Associated Deaths

• According to the Office of the Chief Coroner report entitled 'Drug Overdose Deaths' (see Appendix H), there were 40 opioid associated deaths in 2017. However, this statistic is subject to change in 2018 as new investigations are undertaken and/or ongoing investigations are concluded. In 2016, there were 100 deaths (accidental, suicide and undetermined), and 2015 had 121 deaths. While Saskatchewan saw a significant increase in the opioid associated deaths from 2014 to 2015, hopefully this trend does not continue. Regardless, it speaks to the need for further promotion of the use of naloxone.

Appendix A: CPSS Regulatory Bylaw 18.1

18.1 The Prescription Review Program

(a) Panel of Monitored Drugs – The Prescription Review Program shall apply to all dosage forms of the following drugs, except where indicated otherwise:

ACETAMINOPHEN WITH CODEINE - in all dosage forms except those containing 8 mg or less of codeine

ACETYLSALICYLIC ACID (ASA) WITH CODEINE - in all dosage forms except those containing 8 mg or less of codeine

AMPHETAMINES - in all dosage forms

ANABOLIC STEROIDS ANILERIDINE - in all dosage forms

BARBITUATES

BENZODIAZEPINES – in all dosages and forms

BUPRENORPHINE - in all dosages and forms

BUTALBITAL - in all dosage forms

BUTALBITAL WITH CODEINE - in all dosage forms

BUTORPHANOL

CHLORAL HYDRATE

COCAINE - in all dosage forms

CODEINE - as the single active ingredient, or in combination with other active ingredients, in all dosage forms except those containing 20 mg per 30 ml or less of codeine in liquid for oral administration

DIETHYLPROPION - in all dosage forms

FENTANYL - in all dosage forms

GABAPENTIN

HYDROCODONE - DIHYDROCODEINONE - in all dosage forms

HYDROMORPHONE - DIPHRYDROMORPHONE - in all dosage forms

LEVORPHANOL - in all dosage forms

MEPERIDINE - PETHIDINE - in all dosage forms

METHADONE - in all dosage forms

METHYLPHENIDATE - in all dosage forms

MORPHINE - in all dosage forms

NORMETHANDONE-P-HYDROXYEPHEDRINE - in all dosage forms

OXYCODONE - as the single active ingredient or in combination with other active ingredients in all dosage forms

OXYMORPHONE

PANTOPON - in all dosage forms

PENTAZOCINE - in all dosage forms

PHENTERMINE - in all dosage forms

PROPOXYPHENE - in all dosage forms (

- b) Prescriptions for drugs covered by the Prescription Review Program shall be issued by physicians according to the policies and procedures agreed to and amended from time to time by the College of Dental Surgeons of Saskatchewan, the College of Physicians and Surgeons of Saskatchewan, the Saskatchewan Registered Nurses Association and the Saskatchewan College of Pharmacists.
- (c) In order to prescribe a drug to which the Prescription Review Program applies, physicians shall complete a written prescription which meets federal and provincial legal requirements and includes the following:
 - (i) The patient's date of birth;
 - (ii) The patient's address;
 - (iii) The total quantity of medication prescribed, both numerically and in written form;
 - (iv) The patient's health services number; and,
 - (v) The prescriber's name and address.
- (d) For the purpose of this bylaw, "written prescription" includes an electronic prescription that meets the requirements for electronic prescribing under the Pharmaceutical Information Program.
- (e) A physician who prescribes a drug to which the Prescription Review Program applies, and who provides the prescription directly to a pharmacy by electronic prescribing, by email or by FAX, or who transmits a prescription in accordance with the policies and protocols of the Pharmaceutical Information Program, need not include both the quantity numerically and in written form.
- (f) If a physician is registered on the Educational Register, the physician shall, in addition to the information in paragraph (c) above, include the following in a prescription for a drug to which the Prescription Review Program applies:
 - (i) The training level of the physician writing the prescription;
 - (ii) The legibly printed name of the Most Responsible Physician (the physician to whom queries regarding the prescription should be addressed);
 - (iii) The legibly printed name of the physician writing the prescription.
- (g) Physicians shall only prescribe part-fills of medications to which the Prescription Review Program applies if the following information is specified in the prescription:
 - (i) The total quantity;
 - (ii) The amount to be dispensed each time; and
 - (iii) The time interval between fills.

- (h) The office of the Registrar may gather and analyze information pertaining to the prescribing of medications to which the Prescription Review Program applies in Saskatchewan for the purpose of limiting the inappropriate prescribing and inappropriate use of such drugs. In order to fulfill that role, the office of the Registrar may, among other activities:
 - (i) Generally, provide education to physicians in order to encourage appropriate prescribing practices by physicians registered by the College;
 - (ii) Alert physicians to possible inappropriate use of medications to which the Prescription Review Program applies by patients to whom they have prescribed such drugs;
 - (iii) Alert physicians to possible inappropriate prescribing of medications to which the Prescription Review Program applies;
 - (iv) Make recommendations to a physician with respect to the physician's prescribing of medications to which the Prescription Review Program applies;
 - (v) Require physicians to provide explanations for their prescribing of medications to which the Prescription Review Program applies. In making requests for explanations, the office of the Registrar may require the physician to provide information about the patient, the reasons for prescribing to the patient, and any knowledge which the physician may have about other narcotics or controlled drugs received by the patient;
 - (vi) Cause information, concerns or opinions of general application to the profession to be communicated to the physicians registered by the College without identifying the particular physician to whom such information relates;
 - (vii)Provide information gathered in connection with the Prescription Review Program to another health professional body including the College of Dental Surgeons of Saskatchewan, the Saskatchewan College of Pharmacists or the Saskatchewan Registered Nurses Association, provided the information gathered is required by that body to perform and carry out the duties of that health professional body pursuant to an Act with respect to regulating the profession. Where the personal health information relates to a member of the health professional body seeking disclosure, disclosure by the Registrar of that information may only be made in accordance with The Health Information Protection Act, and in particular section 27(5) or that Act.
- (i) Physicians shall respond to such requests for explanation, as described in paragraph (h)(v) above, from the office of the Registrar within 14 days of receipt of such a request for information.
- (j) The Registrar, Deputy Registrar, or Prescription Review Program Supervisor may extend the deadline for reply at their discretion, upon receipt of a written request for extension from the physician.
- (k) All physicians who receive such a request for information will comply, to the best of their ability, fully and accurately with such requests for information.
- (I) Failure to comply with paragraphs (h)(v), (i) and (k) above is unbecoming, improper, unprofessional or discreditable conduct.

- (m) Members shall keep a record of all drugs to which the Prescription Review Program applies that are purchased or obtained for the member's practice and a record of all such drugs administered or furnished to a patient in or out of the physician's office, showing:
 - (i) the name, strength and quantity of the drug purchased or obtained;
 - (ii) the name, strength, dose and quantity of the drug administered or furnished;
 - (iii) the name and address of the person to whom it was administered or furnished, and, if applicable, the name and address of the person who took delivery of the drug; and
 - (iv) the date on which the drug was obtained and the date(s) on which the drug was administered, furnished or otherwise disposed of.
- (n) The record referred to in paragraph (m) shall be kept separate from the patient's medical record.

Appendix B: Prescription Review Program Monitored Medications

The following section lists the chemical name of the monitored medication, the type of dosage form that is monitored, and all the tradename (or brand name) products that are currently available.

ACETAMINOPHEN WITH CODEINE - in all dosage forms except those containing 8mg or less of codeine

- EXCLUDES: Tylenol #1, Mersyndol
- Tylenol #2
- Tylenol #3
- Tylenol #4

ACETYLSALICYLIC ACID (ASA) WITH CODEINE - in all dosage forms except those containing 8 mg or less of codeine

- EXCLUDES: 222
- 282
- 292

AMPHETAMINES - in all dosage forms

- Adderall XR
- Dexedrine
- Vyvanse

ANABOLIC STEROIDS (testosterone)

- Andriol
- Androgel
- Testim
- Androderm
- Delatestryl

ANILERIDINE - in all dosage forms

BARBITUATES

Phenobarbital

BENZODIAZEPINES – in all dosages and forms

- Alprazolam (Xanax)
- Bromazepam (Lectopam)
- Chlordiazepoxide
- Clonazepam (Rivotril)
- Clorazepate
- Diazepam (Valium)
- Flurazepam
- Lorazepam (Ativan)
- Nitrazepam (Mogadon)
- Oxazepam
- Temazepam (Restoril)
- Triazolam

BUPRENORPHINE – in all dosages and forms

- Butran Patch
- Suboxone (naloxone combo product)

BUTALBITAL - in all dosage forms & BUTALBITAL WITH CODEINE - in all dosage forms

• Fiorinal, Fiorinal C1/2, Fiorinal C1/4 (ASA, caffeine, codeine [15mg or 30mg], butalbital)

BUTORPHANOL

CHLORAL HYDRATE

COCAINE - in all dosage forms

CODEINE - as the single active ingredient, or in combination with other active ingredients, in all dosage forms except those containing 20 mg per 30 ml or less of codeine in liquid for oral administration

Controlled-release:

Codeine Contin

DIETHYLPROPION - in all dosage forms

FENTANYL - in all dosage forms

- Duragesic patch
- Onsolis buccal film (cancelled post market)
- Abstral sublingual
- Fentora sublingual

GABAPENTIN

Neurontin

HYDROCODONE - DIHYDROCODEINONE - in all dosage forms

- Dalmacol
- Hycodan
- Tussionex

HYDROMORPHONE - DIPHRYDROMORPHONE - in all dosage forms

Immediate-release:

Dilaudid

Controlled-release:

• Hydromorph-Contin

LEVORPHANOL - in all dosage forms

MEPERIDINE - PETHIDINE - in all dosage forms

Demerol

METHADONE - in all dosage forms

Metadol

METHYLPHENIDATE - in all dosage forms

- Ritalin & Ritalin SR
- Biphentin
- Concerta

MORPHINE - in all dosage forms

$\underline{Immediate\text{-}release\text{:}}$

- M.O.S.
- MS-IR
- Statex

Controlled-release:

- MS Contin
- MOS-SR
- M-Eslon
- Kadian

NORMETHANDONE-P-HYDROXYEPHEDRINE - in all dosage forms

OXYCODONE - as the single active ingredient or in combination with other active ingredients in all dosage forms

Immediate-release:

- OXY-IR
- Supeudol

Controlled-release:

OxyNEO

OXYMORPHONE

PANTOPON - in all dosage forms

PENTAZOCINE - in all dosage forms

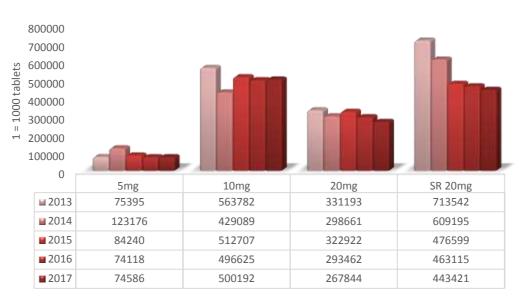
• Talwin

PHENTERMINE - in all dosage forms

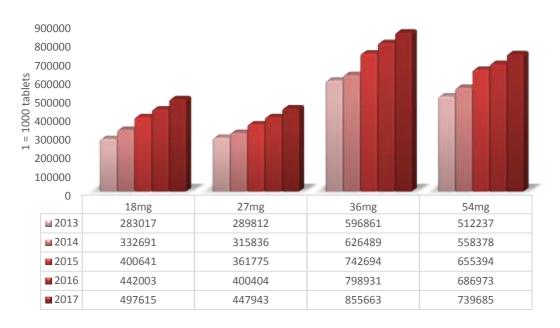
PROPOXYPHENE - in all dosage forms

Appendix C: Stimulants



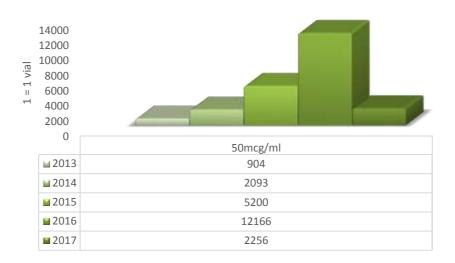


Concerta

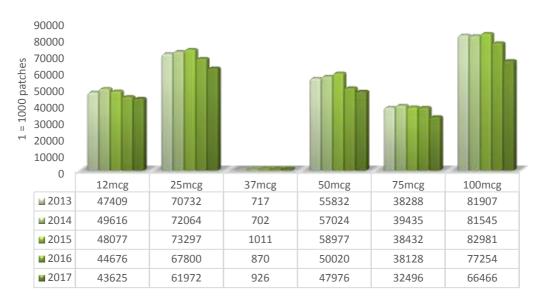


Appendix D: Opioids

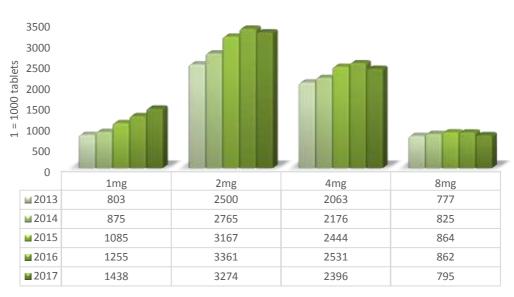
Fentanyl Citrate INJECTION



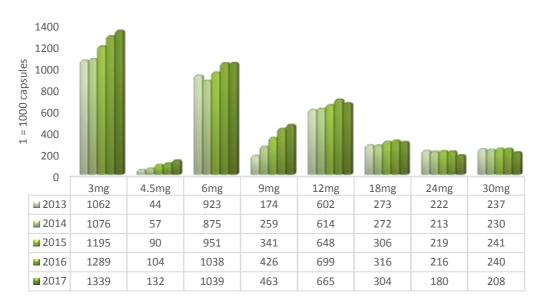
Fentanyl PATCH







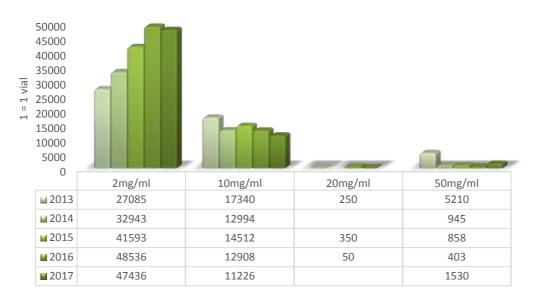
Hydromorph Contin



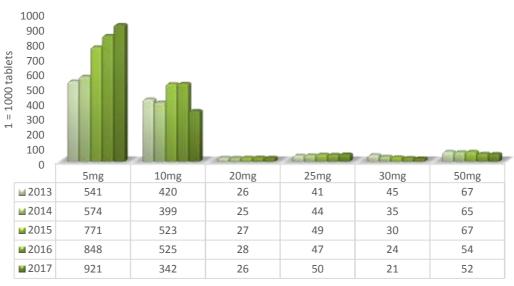
Morphine equivalence – hydromorphone specific

2013 2014		2015	2016	2017		
277,240,828	286,011,403	313,800,478	329,656,298	315,145,148		

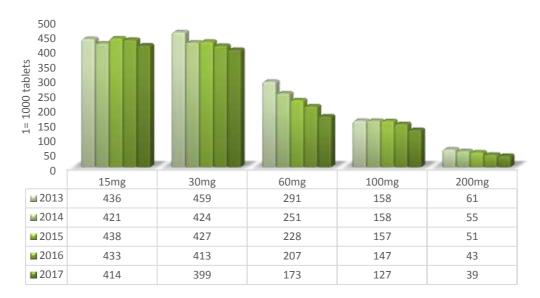
Hydromorphone INJECTION





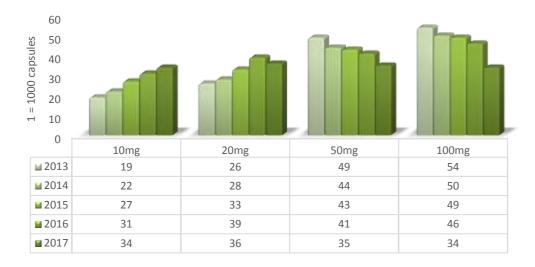


Morphine SR - 12 hour

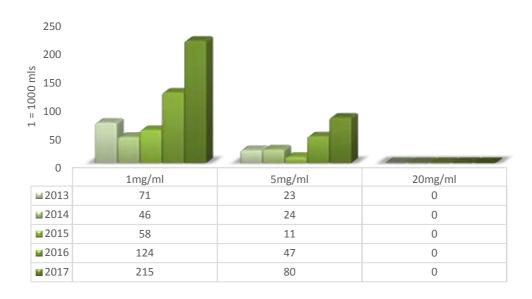




Kadian Morphine SR - 24 hour



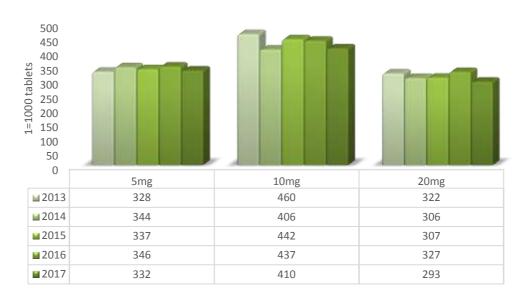




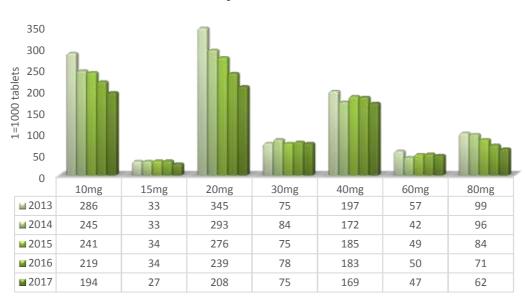
Morphine INJECTION



Oxycodone IR







Morphine equivalence - oxycodone specific*

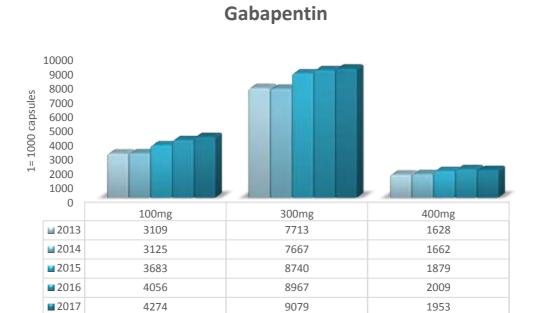
2013 2014		2015	2016	2017	
221,774,355	226,385,865	232,513,924	257,091,458	276,095,108	

^{*} In addition to the oxycodone immediate-release and OxyNeo charts above, the morphine equivalence chart also includes all past and present oxycodone products dispensed in Saskatchewan (e.g. OxyContin, Targin, Percocet, Percocet Demi, Endocet and all marketed oxycodone generics).

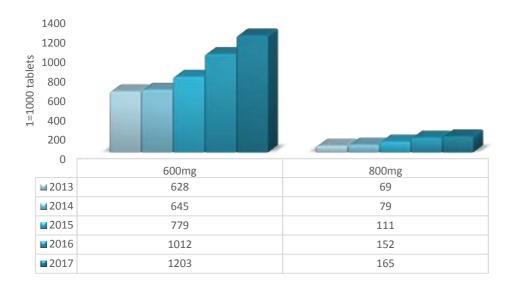
Tylenol with Codeine



Appendix E: Gabapentin



Gabapentin

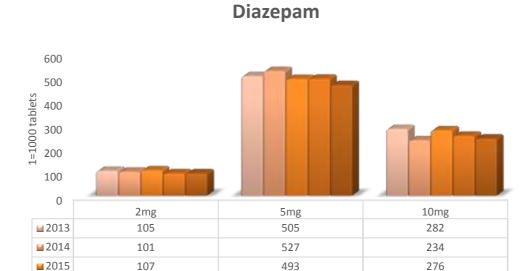


Appendix F: Benzodiazepines



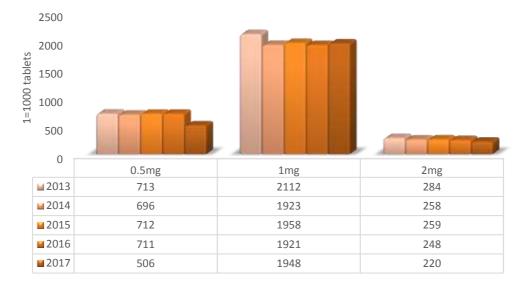
Clonazepam



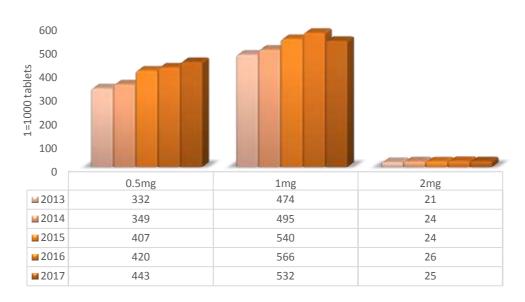


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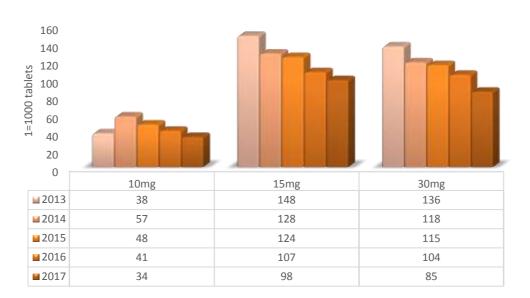
Lorazepam



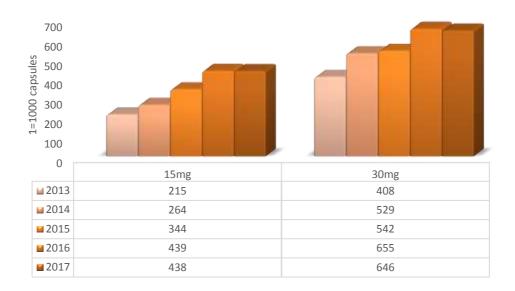
Lorazepam Sublingual



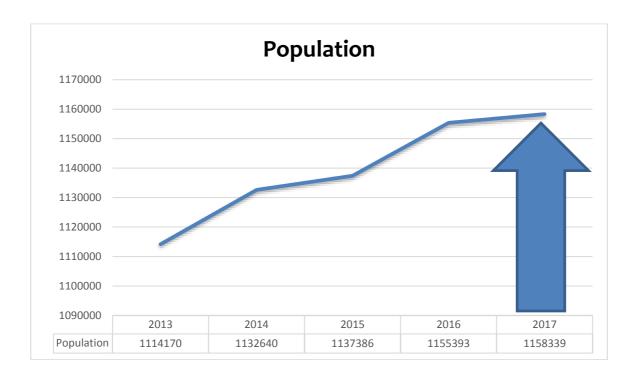
Oxazepam



Temazepam



Appendix G: Saskatchewan Population Growth



Appendix H: Coroner Report – Opioid Related Deaths



Office of the Chief Coroner

DRUG TOXICITY DEATHS Saskatchewan, 2010 to 2017 (Updated – January 11, 2018)

The data in the following tables include all death investigations concluded by the Office of the Chief Coroner (OCC) between January 1, 2010 and December 31, 2017 where the cause of death was due to a Drug Toxicity (Single or Combined Drug Toxicity). The statistics shown are subject to change as new investigations are undertaken and/or on-going investigations are concluded.

For the following tables please note:

- 'Undetermined' indicates that after completing an investigation, there is equal evidence, or a significant contest between one or more classifications.
- 2016 and 2017 data consists of concluded death investigations from January 1, 2016 to December 31, 2017; the data does not include deaths that are still under investigation.

	2010	2011	2012	2013	2014	2015	2016	2017
Accident	52	56	60	62	67	91	84	31
Suicide	21	24	17	21	13	23	13	5
Homicide	255.3		100	270	***	0.000	27	
Undetermined	5	6	9	5	5	7	3	4
Total	78	86	86	88	85	121	100	40

		Codeine	Fentanyl	Herain	Hydrocodone	Hydromorphone	Methadone	Morphine	Oxycodone	Opicid (Unknown)	W-18*
2010	Accident	4	2	***	-	12	11	12	10		
	Suicide	2	1	+	-	2	4	3	-	17	
	Hamicide	1665		-	-	-	(4)	**	-	-	
	Undetermined	100	99	97	-	-	3	1	1	111	
2011	Accident	7	2	4	7.00	19	20	12	5	100	
	Suicide	2	77	**	2.66	1	3	3	4	177	
	Homicide	100	***	-	-	-	-	***	-	-	
	Undetermined	155.0	1			3	1	0.00		144	
2012	Accident	12	6	1	-	16	14	19	3	+	
	Suicide	4	240	***	-	1	1	**	2	\leftrightarrow	
	Homicide	**	***	-	-	-	+	++	-	***	
	Undetermined	2	1	-	-	-	2	2	2		
2013	Accident	3	9	-	-	17	21	10	7	1	
	Suicide	2	1	**	100	4	2	4	1	100	
	Homicide	(+1)	**	***	100	-	(+)	***	-	177.0	
	Undetermined	100	25	-	-	2	2		1	-	
2014	Accident	5	9	· *:		22	20	15	4	-	
	Suicide		2			1	4	2	1	144	
	Homicide	One	-	-	-	2	-		-		
	Undetermined	100	2	***	-	1	-	1	2	140	
2015	Accident	10	21	0+	-	30	27	21	5	+0	1"
	Suicide	1	1	94	Check	4	2	3	2	277	
	Homicide	799	**	94	100		001	**	**	***	
	Undetermined	1	**	***	100	3	(40)	3	77.	77.	
2016	Accident	8	16	1	-	23	31	18	4		
	Suicide	1	-	-		1	2	01	100	-	
	Homicide	10	44	-	100	100	-	90	-	40	
	Lindetermined	1	1	-	75.5	1	1	140			
017	Accident	6	7		3	6	10	10	.)		10
	Suidde	1	***	-	-	1	-		-	544.1	100
	Homicide	-07	-		-		(ac-	190	-	40	
	Undetermined	- 40			-	1	1	44	. 1	2.00	-

^{*} Shot Drugs Containing W-18 — As part of a 2015 investigation into the combined drug toxicity death of a male, age 25, there were tablets found at the scene which were analyzed and found to contain fentanty and W-18. Given the limitations of toxicologies yearing, it is not possible to quantify W-18 beneath a currain level within a person's blood. The Office of the Chief Coroner was unable to determine whether W-18 contributed to this individual's death. Also, based on the circumstances of the death, it could not be confirmed whether the deceased ingested any of the totalets that contained the festivary and W-18. The individual's cause of death was combined drug toxicity involving a number of drugs including festianyl and morphine which are reflected in the statistics contained in the tablets of this report.

Appendix I: Budget and Actuals

Appendix J: Audited Financial Statements 2017