

Studying trends in fatal and non-fatal overdose: Findings from Drug Trends 2019

National Drug and Alcohol Research Centre, UNSW, Sydney

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Drug Trends Team:

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Conflicts of interest:

- Amy Peacock: untied educational grant from Mundipharma and Segirus for study of opioid medications
- Raimondo Bruno: untied educational grant from Mundipharma and Indivior for study of opioid medications
- Louisa Degenhardt: untied educational grant from Mundipharma, Seqirus and Indivior for study of opioid medications
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- Paul Dietze: untied educational grant from Gilead Sciences for HCV research and untied educational grant from Indivior



Drug Trends

To establish, maintain, and continuously improve monitoring of trends in illicit drug use, harms, and markets across Australia

National Monitoring: Secondary Data

Jurisdictional Monitoring: Secondary Data

Sentinel Sample Monitoring

Online **Monitoring**

Mortality Data

Drug-induced deaths from registry and coronial data

Hospitalisation Data

Drug-induced hospitalisations

Other Sources

Household survey, treatment data etc

Various sources
assessing drug use
and harms at the
population-level (e.g.,
emergency
department
presentations) and
subpopulation level
(e.g., needle-syringe
program visits)

Illicit Drug
Reporting
System (IDRS)
Ecstasy and
Related Drug
Reporting
System (EDRS)

Annual interviews
with people who
inject drugs (IDRS)
and who use
stimulants (EDRS)

Cryptomarket Data

Scraping listings on darknet drug markets

Input from researchers, national stakeholders, and jurisdiction stakeholders to inform priority research questions

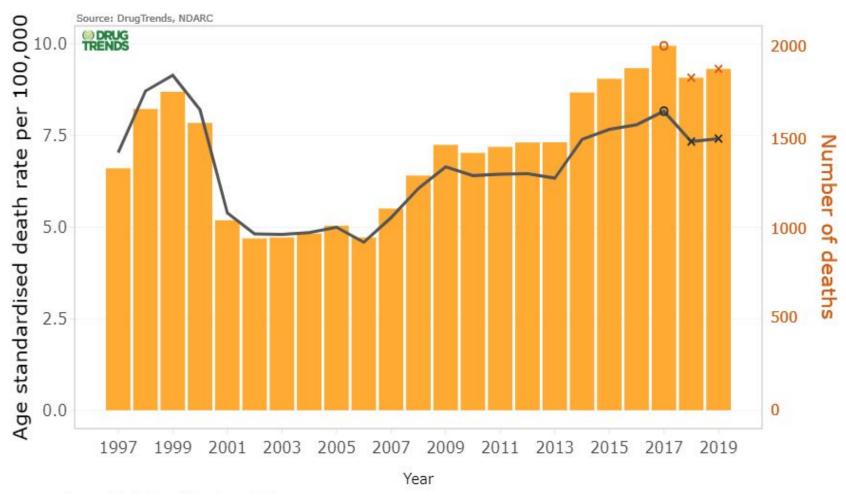
Analytical reports

Drug Trends

Caveats

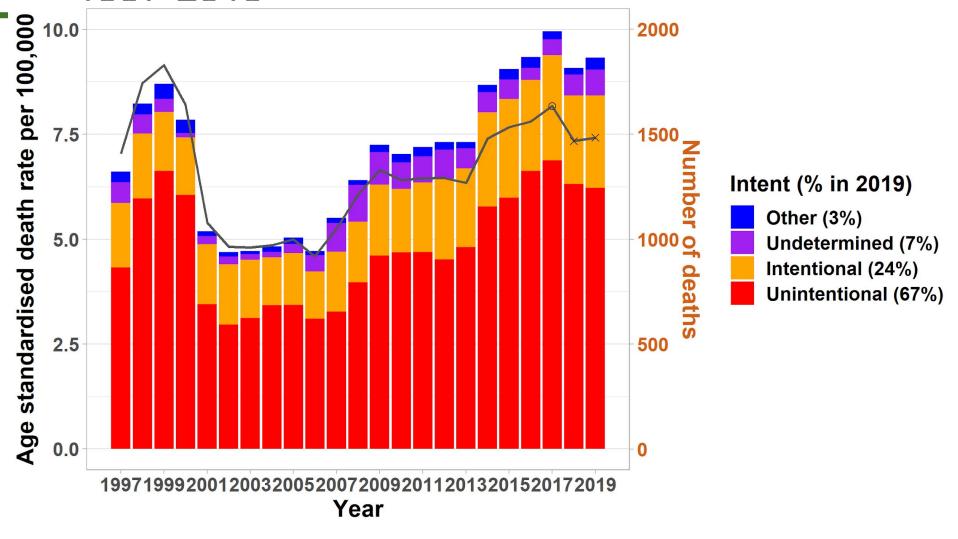
- ABS undertakes a revision process over a 3-year period; estimates for 2017, 2018, and 2019 are not final
- ICD codes have limited specificity for drugs and numbers may differ between organisations reporting on deaths due to codes used
 - Small numbers ≤5 are not shown to protect confidentiality
 - Does not include deaths caused by alcohol

Drug-induced deaths in Australia, all ages, 1997-2019



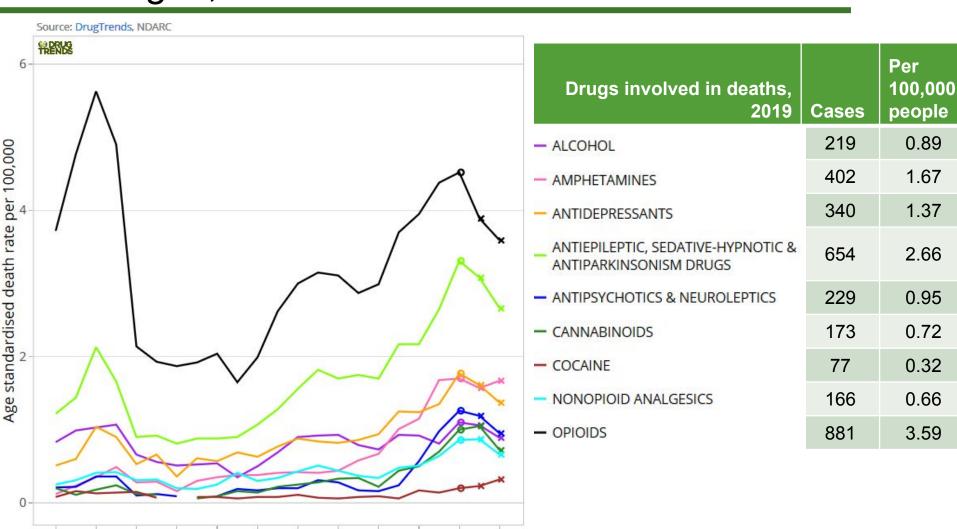


Drug-induced deaths in Australia, all ages, 1997-2019

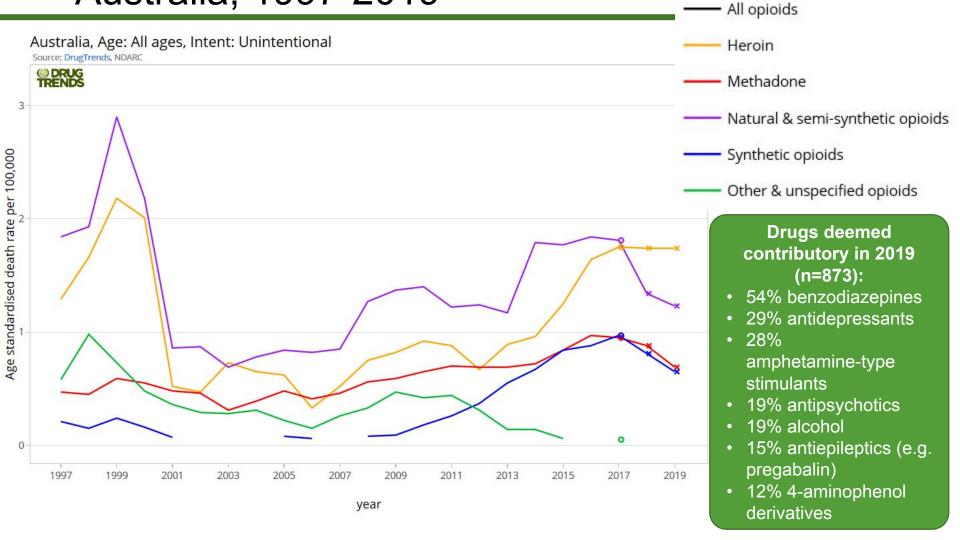




Unintentional drug-induced deaths in Australia, all-ages, 1997-2019



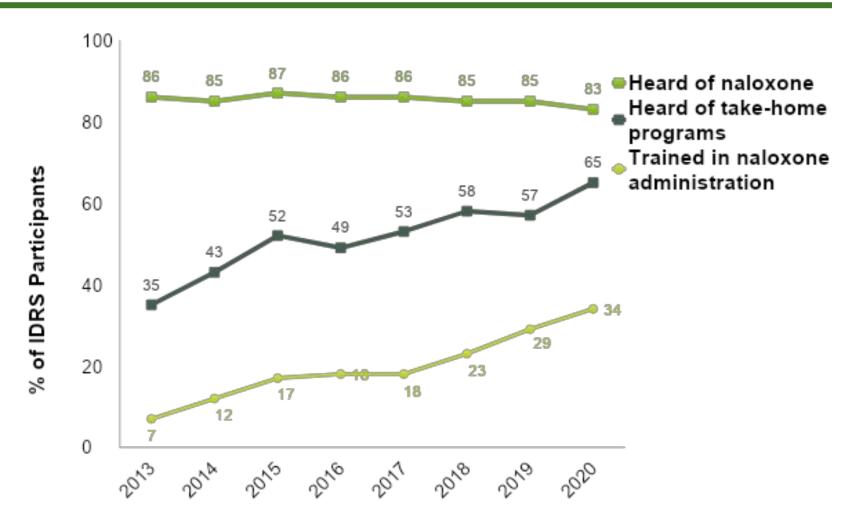
Unintentional opioid-induced deaths in Australia, 1997-2019





Chrzanowska et al. (2021)

Naloxone knowledge and training in the sample who inject drugs (Illicit Drug Reporting System)





Naloxone knowledge and training in the sample who inject drugs (Illicit Drug Reporting System)

"Have you been trained in naloxone administration?"



13% reported past 12-month opioid overdose

5%
had been resuscitated
by someone who had
participated in a
take-home naloxone
program

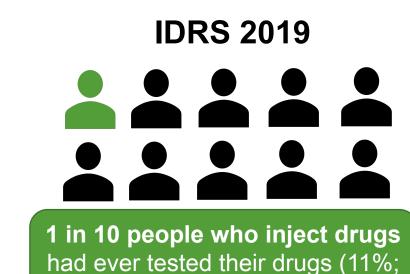




Preventing overdose: food for thought

Non-fatal overdose can lead to significant harms (e.g., brain injury). What are the other effective strategies for preventing <u>any</u> overdose?

- Naloxone
- Opioid agonist treatment (access, coverage, and retention)
- Supervised consumption facilities
- Overdose education (including peer-to-peer)
- Drug checking services (+ information-sharing)



71%
Personal testing kit last time

6% in the past year)



Preventing overdose: food for <u>thought</u>

Non-fatal overdose can lead to significant harms (e.g., brain injury). What are the other effective strategies for preventing any overdose?

- Naloxone
- Opioid agonist treatment (access, coverage, and retention)
- Supervised consumption facilities
- Overdose education (including peer-to-peer)
- Drug checking services (+ information-sharing)
- Wearable overdose detection technology
- Others? And things we know don't work?



SCIENCE TRANSLATIONAL MEDICINE | RESEARCH ARTICLE

OPIOID OVERDOSE

Opioid overdose detection using smartphones

Rajalakshmi Nandakumar¹, Shyamnath Gollakota¹*, Jacob E. Sunshine²*

Early detection and rapid intervention can prevent death from opioid overdose. At high doses, opioids (particu larly fentanyl) can cause rapid cessation of breathing (apnea), hypoxemic/hypercarbic respiratory failure, and death, the physiologic sequence by which people commonly succumb from unintentional opioid overdose. We present algorithms that run on smartphones and unobtrusively detect opioid overdose events and their precursors. Our proof-of- concept contactless system converts the phone into a short-range active sonar using frequency shifts to identify respiratory depression, apnea, and gross motor movements associated with acute opioid toxicity. We develop algorithms and perform testing in two environments: (i) an approved supervised injection facility (SIF), where people self-inject illicit opioids, and (ii) the operating room (OR), where we simulate rapid, opioidinduced overdose events using routine induction of general anesthesia. In the SIF (n = 209), our system identified postinjection, opioid-induced central apnea with 96% sensitivity and 98% specificity and identified respiratory depression with 87% sensitivity and 89% specificity. These two key events commonly precede fatal opioid overdose. In the OR, our algorithm identified 19 of 20 simulated overdose events. Given the reliable reversibility of acute opioid toxicity, smartphone-enabled overdose detection coupled with the ability to alert naloxoneequipped friends and family or emergency medical services (EMS) could hold potential as a low-barrier, harm reduction intervention.

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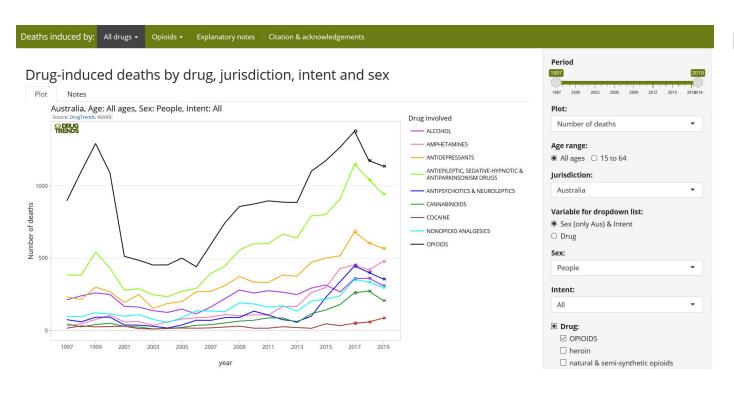
Acceptability of technological solutions for overdose monitoring: Perspectives of people who use drugs

V. W. L. Tsang^a 📵, K. Papamihali, MPH^b, A. Crabtree, MD, PhD^b 📵, and J. A. Buxton, MBBS, MHSc, FRCPC^{a,b} 📵 ^aFaculty of Medicine, University of British Columbia, Vancouver, Canada; ^bBC Centre for Disease Control, Vancouver, Canada

Canada, Despite the availability of observed consumption sites in British Columbia (BC), data suggest people who use drugs (PWUD) alone in private residences are most at risk of overdose death. These individuals may not use consumption sites due to fear of stigmatization, lack of privacy, or personal preference. It is this population that would benefit from overdose monitoring alternatives. Methods: This 2018 study employed two sources of data. (1) A provincial harm reduction distribu-tion site client survey administered at 27 sites across BC asked about cell phone possession and functionality. (2) Structured interviews with PWUD in Vancouver gathered perspectives regarding monitoring devices and alerting systems. *Results*: The client survey was administered to 486 individuals. Among 443 respondents, 48% (n=212) owned a cellphone and 68% (n=115) of individuals with a cellphone with access to internet (n=168) would use an application to mitigate opioidrelated overdose deaths. Thirty qualitative interviews were performed; thematic analysis of the responses identified three major themes – safety, public versus private drug use, and technological monitoring and function. The relevance of technological devices was affected by the inconsistent availability of cellphones, concerns about anonymity, as well as personal comfort while using drugs. Conclusions: Technological applications may not be suitable for clients with transient life

KEYWORDS people who use drugs; technology; fentanyl

For further information: visualisation



Disaggregate by:

- Drug
- Intent
- Age
- Sex
- Jurisdiction
- Remoteness



For more visualisations, go to:

https://ndarc.med.unsw.edu.au/drug-trends-visualisations





















For further information

IDRS 2021 results published late Nov/early Dec 2021

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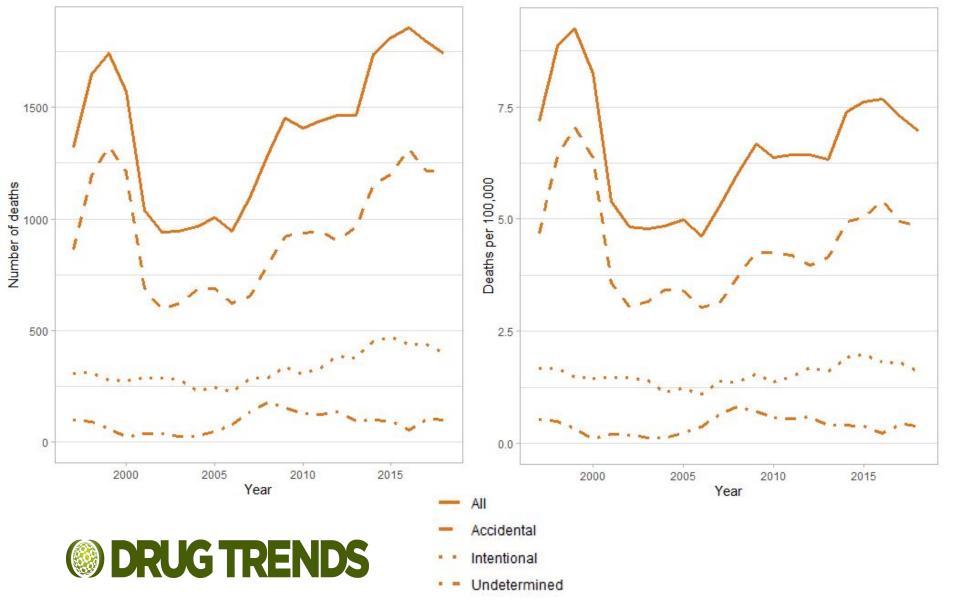




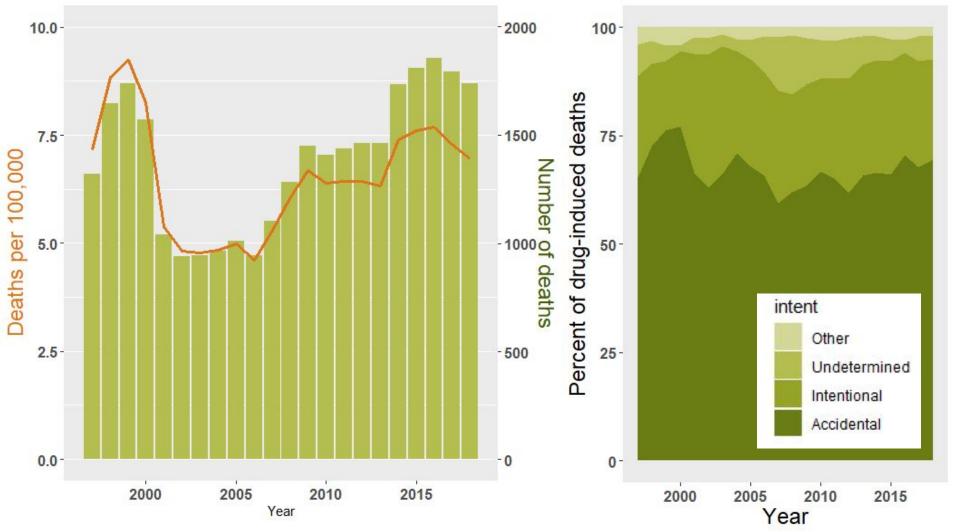




Drug-induced deaths in Australia, all ages, 1997-2018

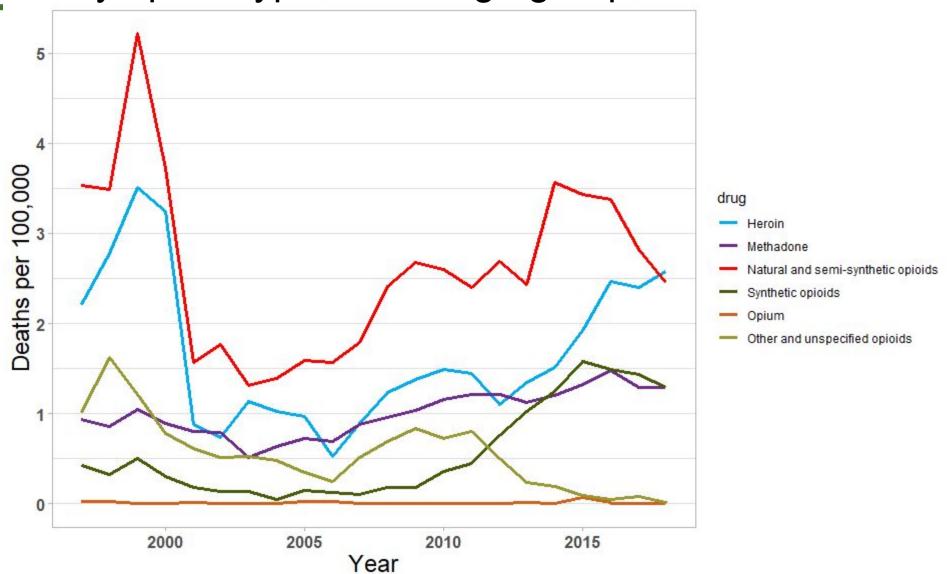


Drug-induced deaths in Australia, all ages, 1997-2018





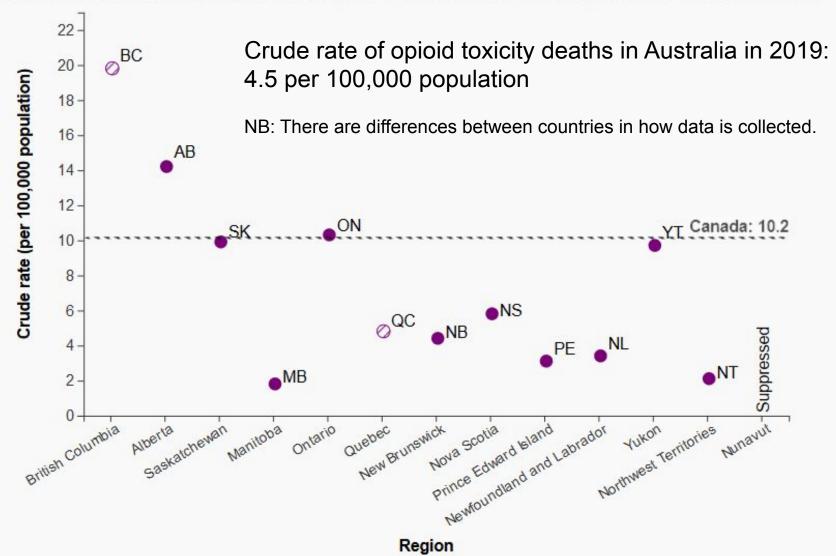
Accidental opioid-induced deaths in Australia by opioid type, 15-64 age group, 1997-2018



Total apparent opioid toxicity deaths in

Canada

Crude rate (per 100,000 population) of total apparent opioid toxicity deaths by province or territory in 2019



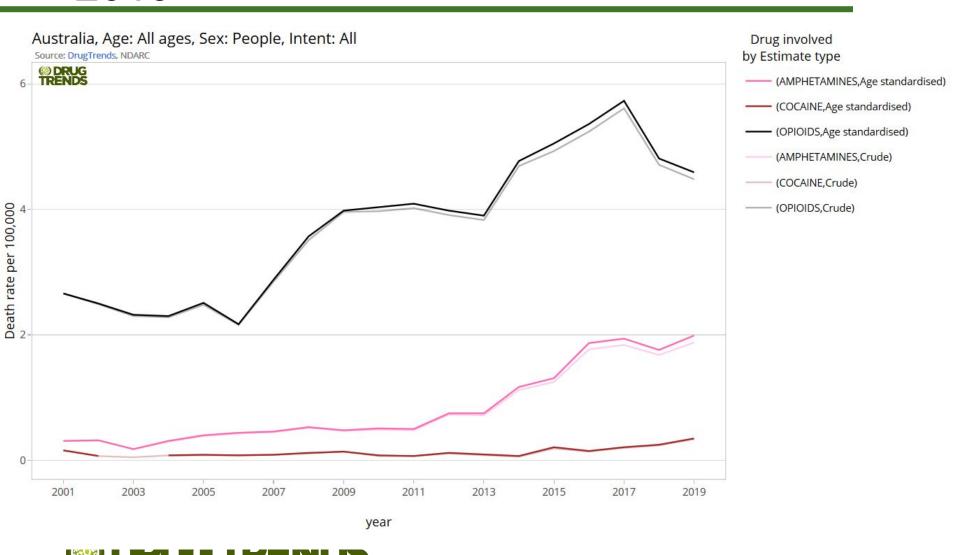
https://health-infobase.canada.ca/substance-related-harms/opioids-stimulants/graphs?index=422

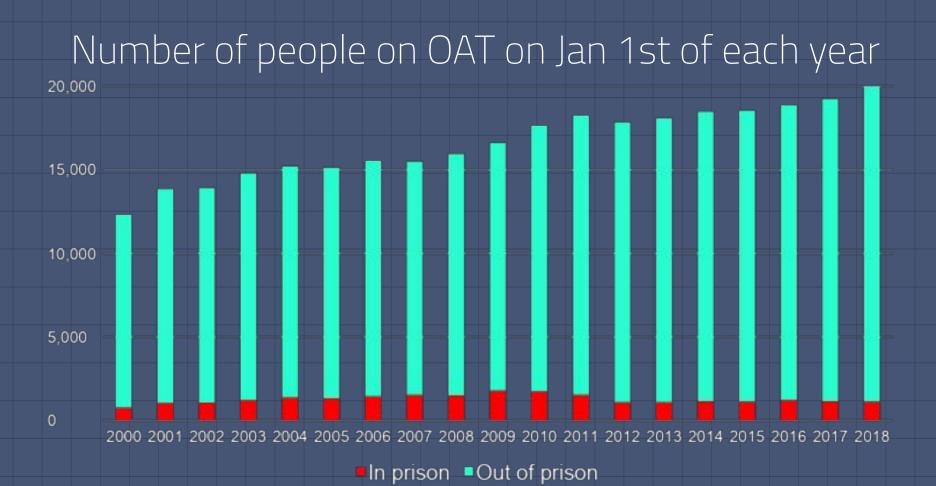
Opioid toxicity deaths, 2019

	Crude rate per 100,000				
	Canada	Australia			
2019	10.2	4.5			
2018	11.8				
2017	10.7	5.6			
2016	7.8	5.2			

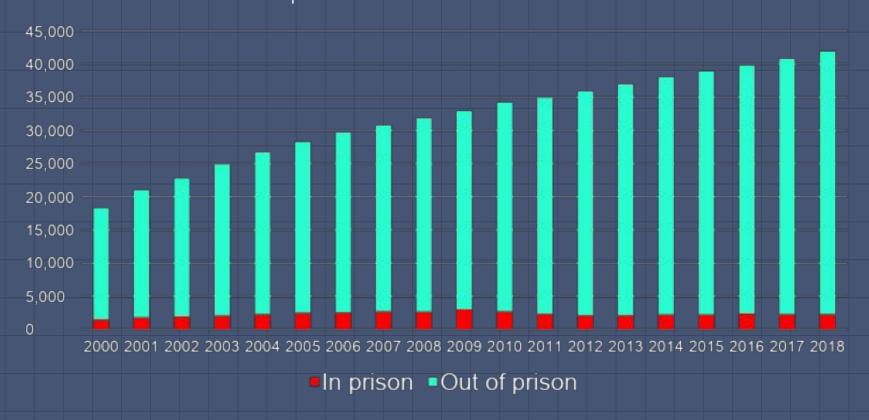
NB: There are differences in between countries in how data is collected.

Opioid and stimulant toxicity deaths, 2019

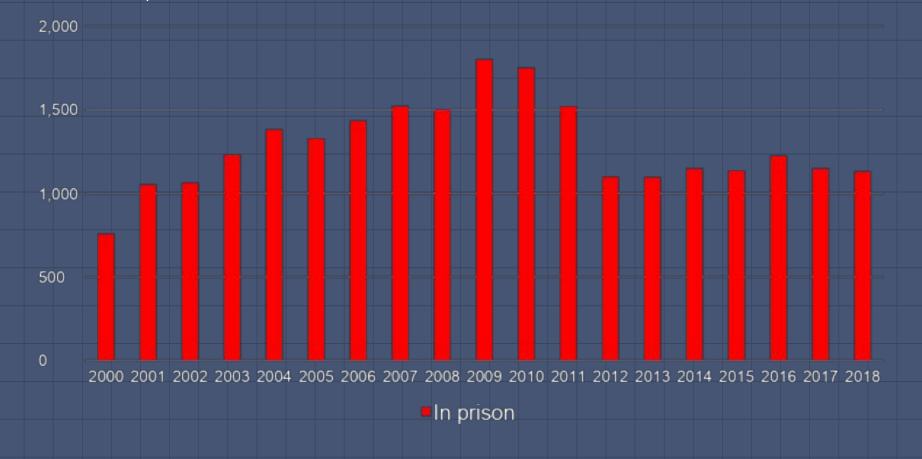




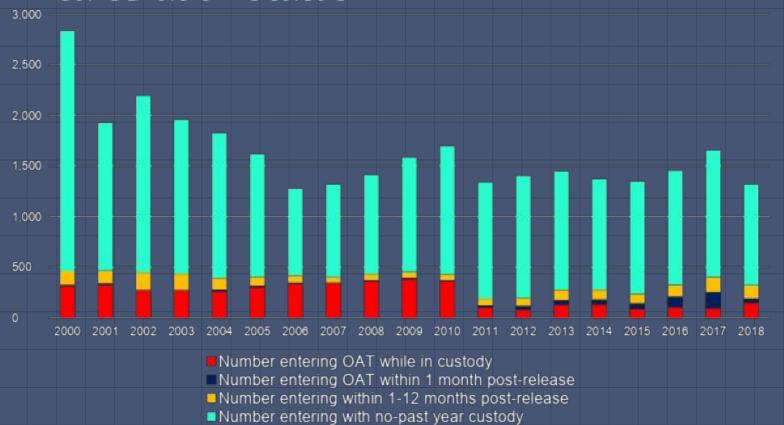
Number of people in the OAT cohort (with at least one OAT episode since 2001)



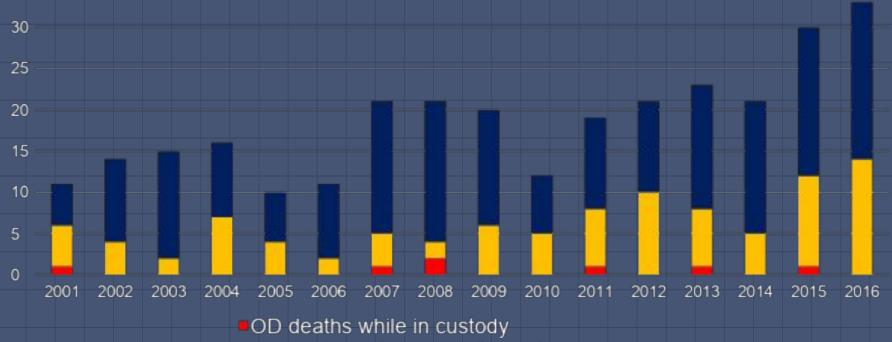
Number of people on OAT in prison on Jan 1st of each year



New (first time) entries into the OAT program by incarceration status

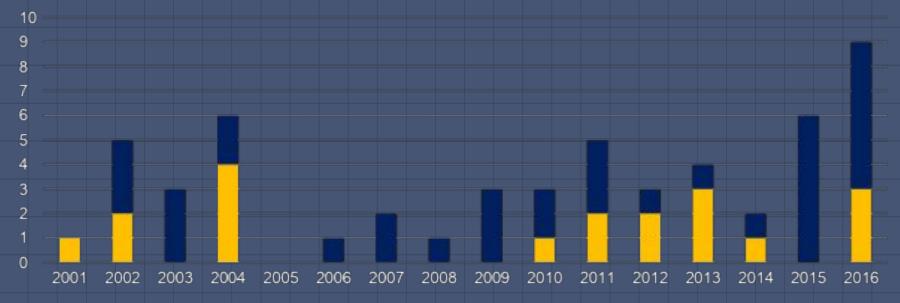


Opioid overdose death by incarceration status -among those incarcerated in the past year-



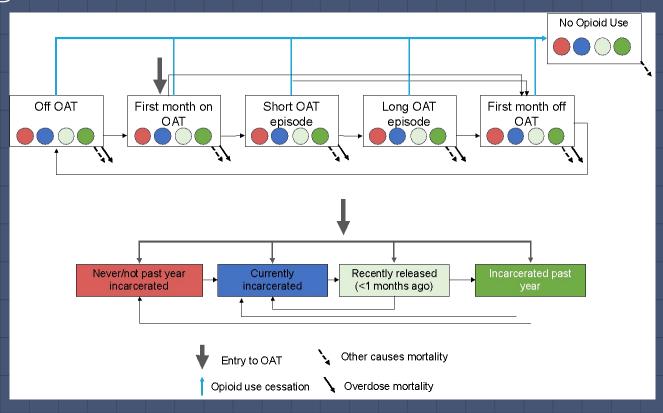
- OD deaths within 1 month post-release
- ■OD deaths within 1-12 months post-release

Overdose deaths on OAT by incarceration status -among those incarcerated in the past year-

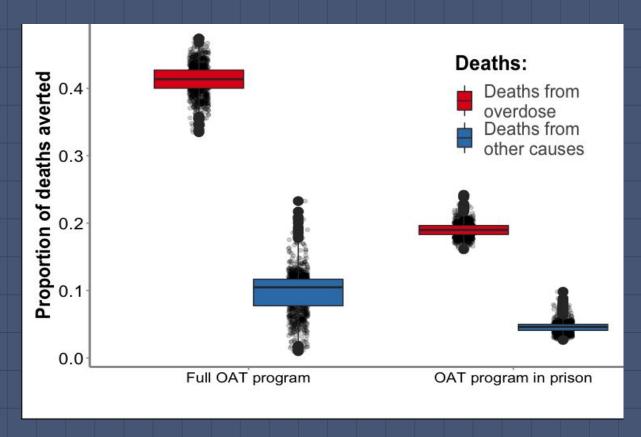


- OD deaths on OAT while in custody
- OD deaths on OAT within 1 month post-release
- OD deaths on OAT within 1-12 months post-release

Mathematical model to estimate impact of OAT program in NSW



Proportion of deaths averted through OAT program from 2001-2010



Number of deaths averted through the OAT program between 2001–2010

	Overdose deaths	Change in overdose	Deaths from other causes	Change in other cause	Total Deaths	Change in total deaths
	Mean [95% I]	deaths vs. baseline Mean	Mean [95% I]	deaths vs. baseline Mean	Mean [95% I]	vs. baseline Mean
Baseline	723	[95% I] -	1541	[95% I] -	2264	[95% I] -
No Opiate Agonist	[540-895] 1233		[1156-1873] 1711		[1692-2755] 2944	
Therapy	[900-1552]		[1257-2133]		[2166-3636]	
No prison OAT program	893 [664-1112]		1616 [1198-1974]		2508 [1871-3057]	