

Patterns and correlates of pregabalin use among a sample of people who inject drugs in Australia

R. Sutherland, P. Dietze, N. Gisev, R. Bruno, Campbell, G., Memedovic, S. & A. Peacock



 Gamma-aminobutyric acid (GABA) analogue that has analgesic and anticonvulsant effects (i.e. non-opioid)



- Gamma-aminobutyric acid (GABA) analogue that has analgesic and anticonvulsant effects
- Registered in Australia in 2005 as a Schedule 4 (prescription only) medicine for treatment of neuropathic pain and epilepsy



- Gamma-aminobutyric acid (GABA) analogue that has analgesic and anticonvulsant effects
- Registered in Australia in 2005 as a Schedule 4 (prescription only) medicine for treatment of neuropathic pain and epilepsy
- Listed for subsidy on the Pharmaceutical Benefits Scheme in 2013



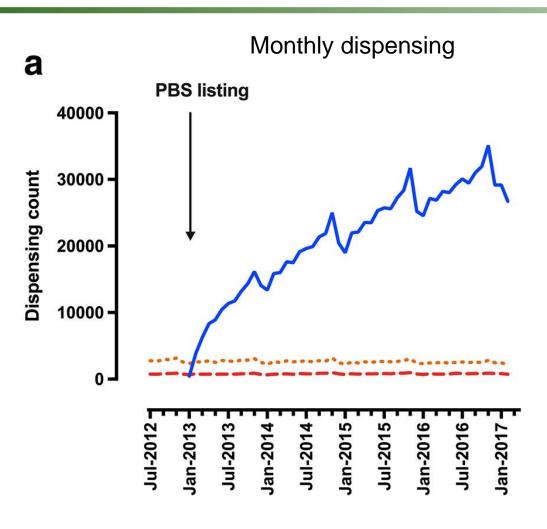
- Gamma-aminobutyric acid (GABA) analogue that has analgesic and anticonvulsant effects
- Registered in Australia in 2005 as a Schedule 4 (prescription only) medicine for treatment of neuropathic pain and epilepsy
- Listed for subsidy on the Pharmaceutical Benefits Scheme in 2013
- Sixth most prescribed subsidised drug in Australia, 2016-2017

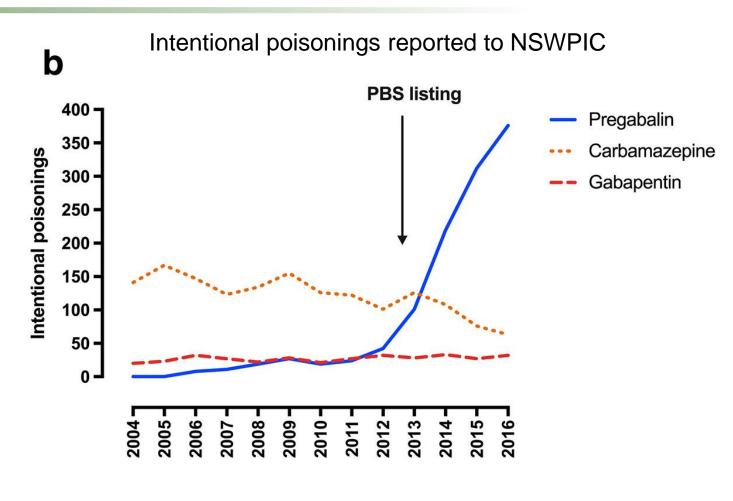


- Gamma-aminobutyric acid (GABA) analogue that has analgesic and anticonvulsant effects
- Registered in Australia in 2005 as a Schedule 4 (prescription only) medicine for treatment of neuropathic pain and epilepsy
- Listed for subsidy on the Pharmaceutical Benefits Scheme in 2013
- Sixth most prescribed subsidised drug in Australia, 2016-2017
- Increase in pregabalin-related harms



Background: Increasing evidence of harms





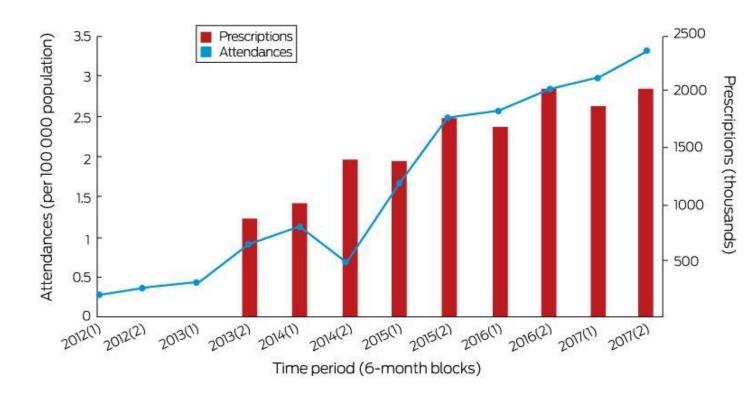


Cairns R, Schaffer AL, Ryan N, et al. Rising pregabalin use and misuse in Australia: trends in utilization and intentional poisonings. Addiction 2018; https://doi.org/10.1111/add.14412.

Background: Increasing evidence of harms

- Tenfold increase since 2012
- 18.4% increase for each additional 100,000 prescriptions
- Frequently used with other sedatives (e.g. benzodiazepines)

Pregabalin misuse related ambulance attendances (VIC)





Crossin, R., Scott, D., Arunogiri, S., Smith, K., Dietze, P. & Lubman, D. (2019) Pregabalin misuse-related ambulance attendances in Victoria, 2012–2017: characteristics of patients and attendances, Med J Aust; 210 (2); 75-79

Background: Deaths involving* pregabalin

Among Australians of all ages:

- <20 annually prior to 2015
- 72 deaths 2016 (total 1,858; 4%)
- 100 deaths 2017 (total 1,795; 6%)

*Most of these deaths were attributable to opioids

Concern that concomitant use of pregabalin and opioids can increase mortality risk



Aims

• Examine patterns of pregabalin use (prescribed and not prescribed) among a sample of people who frequently inject drugs (PWID) in Australia.



Aims

- Examine patterns of pregabalin use (prescribed and not prescribed) among a sample of people who frequently inject drugs (PWID) in Australia.
- Identify correlates of prescribed and non-prescribed pregabalin use among PWID.



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 populationlevel (e.g., emergency
department
presentations) and
subpopulation level (e.g.,
needle-syringe program
visits)

Illicit Drug Reporting System (IDRS)

Annual interviews with people who inject drugs (IDRS) ~900 per/year

Cryptomarket Data

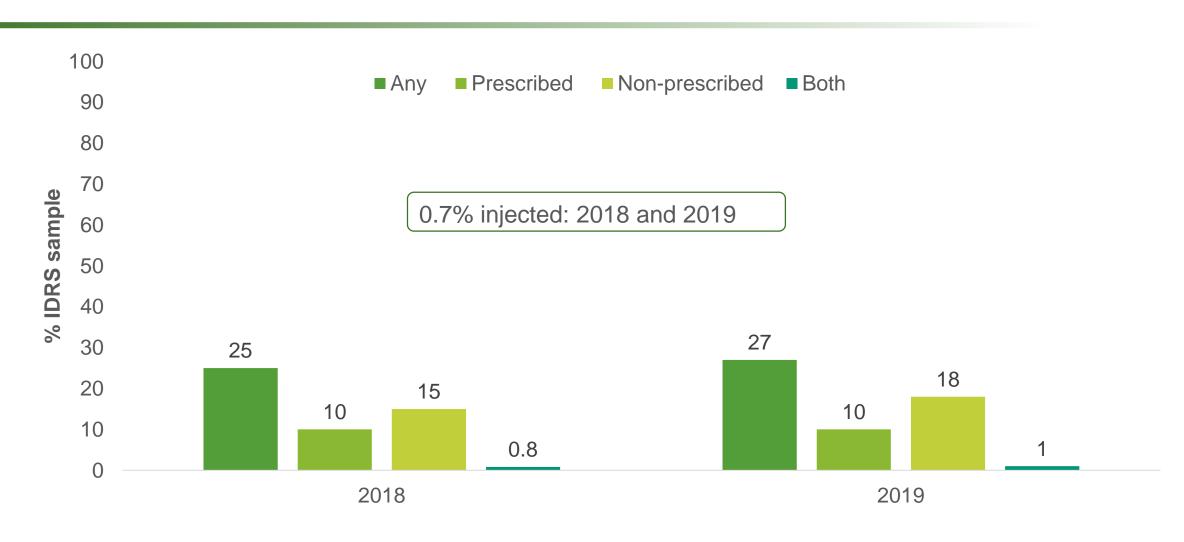
Scraping listings on darknet drug markets

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

Analytical reports

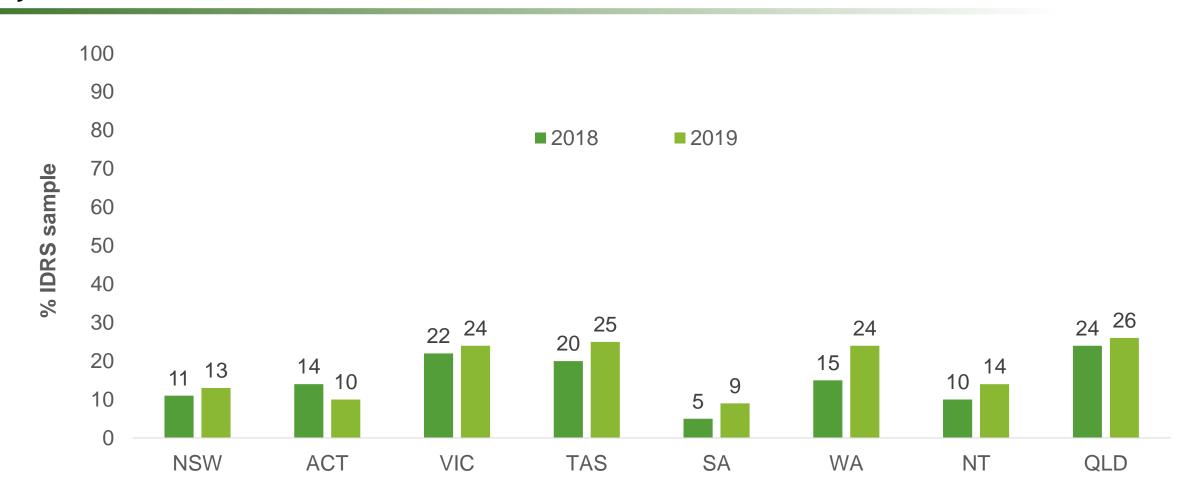
Aim 1: Patterns of pregabalin use, 2018-2019

Past six month pregabalin use, 2018-2019



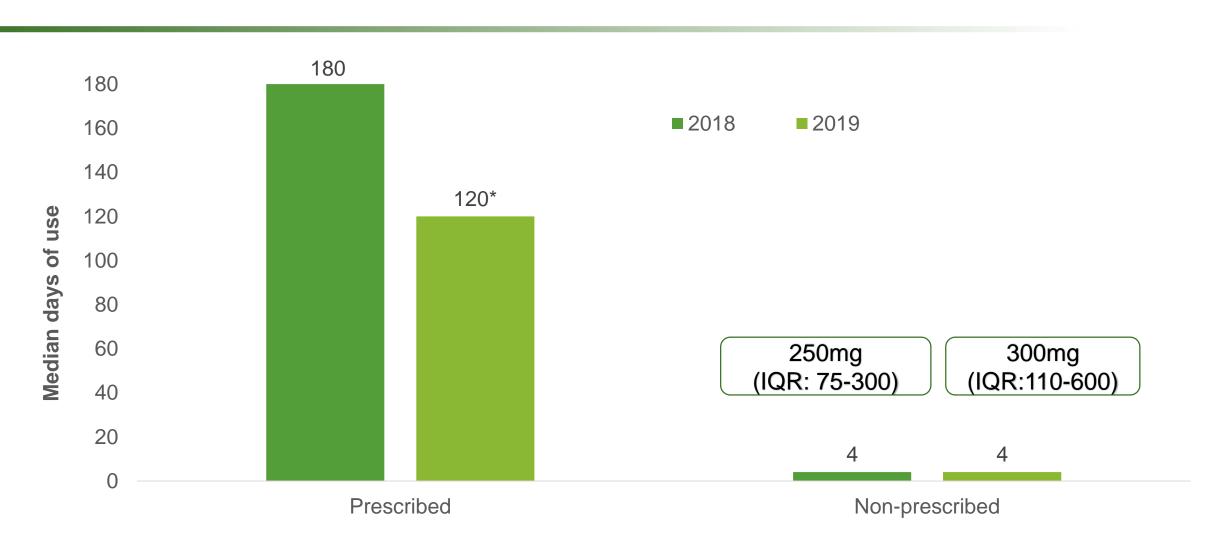


Past six month non-prescribed pregabalin use, by jurisdiction





Frequency of use, 2018-2019



*p<0.05





- Age
- Gender
- Aboriginal or Torres Strait Islander
- Employment status
- Accommodation
- Substance use (heroin, OST, pharmaceutical opioids, benzodiazepines, stimulants)
- Dependence (opioids, stimulants)
- Hazardous alcohol consumption
- Bingeing
- Overdose
- Psychological distress (K10)
- Self-reported mental health problems
- Mobility problems
- Pain/discomfort



	No use (n=678)	Prescribed use only (n=86)	Non-prescribed use only (n=133)
Non-prescribed pharmaceutical opioid use %	26	46	49
Prescribed benzodiazepine use %	25	54	34
Non-prescribed benzodiazepine use %	24	31	55
Stimulant use %	76	80	92
Overdose (past year) %	18	31	36
Pain/discomfort (day of interview) %	50	81	53

Note: those who had used both prescribed and non-prescribed pregabalin (n=7) excluded from analysis



	No use (n=678)	Prescribed use only (n=86)	Non-prescribed use only (n=133)
Non-prescribed pharmaceutical opioid use %	26	46	49
Prescribed benzodiazepine use %	25	54	
Non-prescribed benzodiazepine use %	24	31	55
Stimulant use %	76	80	
Overdose (past year) %	18	31	36
Pain/discomfort (day of interview) %	50	81	

No use as the referent category; those who had used both prescribed and non-prescribed pregabalin (n=7) excluded from analysis



	No use (n=678)	Non-prescribed use only (n=133)
Non-prescribed pharmaceutical opioid use %	26	49 🕇
Prescribed benzodiazepine use %	25	34
Non-prescribed benzodiazepine use %	24	55
Stimulant use %	76	92
Overdose (past year) %	18	36
Pain/discomfort (day of interview) %	50	53

No use as the referent category; those who had used both prescribed and non-prescribed pregabalin (n=7) excluded from analysis



	Prescribed use only (n=86)	Non-prescribed use only (n=133)
Non-prescribed pharmaceutical opioid use %	46	49
Prescribed benzodiazepine use %	54	34
Non-prescribed benzodiazepine use %	31	55
Stimulant use %	80	92
Overdose (past year) %	31	36
Pain/discomfort (day of interview) %	81	53 ↓

Prescribed use is the referent category; those who had used both prescribed and non-prescribed pregabalin (n=7) excluded from analysis



Little overlap, indicating sufficient prescribing



- Little overlap, indicating sufficient prescribing
- Harm reduction messages re: concomitant use of opioids and pregabalin (consumers and prescribers), incorporated into existing overdose materials



- Little overlap, indicating sufficient prescribing
- Harm reduction messages re: concomitant use of opioids and pregabalin (consumers and prescribers)
- Non-prescribed pregabalin consumers appear to be a riskier group of consumers (e.g. non-fatal overdose, stimulant use)



- Little overlap, indicating sufficient prescribing
- Harm reduction messages re: concomitant use of opioids and pregabalin (consumers and prescribers)
- Non-prescribed pregabalin consumers appear to be a riskier group of consumers (e.g. non-fatal overdose, stimulant use)
- Motivations for non-prescribed use unclear



- Little overlap, indicating sufficient prescribing
- Harm reduction messages re: concomitant use of opioids and pregabalin (consumers and prescribers)
- Non-prescribed pregabalin consumers appear to be a riskier group of consumers (e.g. non-fatal overdose, stimulant use)
- Motivations for non-prescribed use unclear

	Hazard ratio (95% CI)	Hazard ratio (95% CI)	
Suicidal behaviour and deaths from suicide			
All gabapentinoids	-	1.26 (1.20 to 1.32)	Molero et al BMJ 2019; 365: l2147
Pregabalin only		1.26 (1.19 to 1.32)	
Gabapentin only		1.04 (0.89 to 1.21)	



Acknowledgements

Funding

Australian Government Department of Health for funding provided under the Drug and Alcohol Program

Current Team

- National Drug and Alcohol Research Centre: Amy Peacock, Daisy Gibbs, Toni Karlsson, Anant Mathur, Julia Uporova, Rosie Swanton, Agata Chrzanowska, Louisa Degenhardt and Michael Farrell
- Burnet Institute: Amy Kirwan, Cristal Hall, Campbell Aitken and Paul Dietze
- School of Medicine, University of Tasmania: Ellie Bucher, Callula Sharman and Raimondo Bruno
- National Drug Research Institute (WA): Jodie Grigg, Seraina Agramunt and Simon Lenton
- School of Public Health, The University of Queensland (QLD): Caroline Salom
- Northern Territory Department of Health (NT): Chris Moon

And the many people here at NDARC and elsewhere who have contributed to the program over the last two+ decades!

Other Acknowledgements

- IDRS participants: for the time they give to complete the interview
- Data custodians: for the timely provision of data and input on analysis and interpretation
- Stakeholders and Advisory Committee: for engagement with and input on Drug Trends

Further questions?

rachels@unsw.edu.au



















