To cite: Di Donato MF.

Mathieson S. Ferreira GE. et al.

Trends in opioid dispensing

to injured workers following

codeine scheduling changes

in Australia: a retrospective

2025;15:e092651. doi:10.1136/

cohort study. BMJ Open

bmjopen-2024-092651

Prepublication history

and additional supplemental

available online. To view these

online (https://doi.org/10.1136/

files, please visit the journal

bmjopen-2024-092651).

Received 19 August 2024

Accepted 25 February 2025

material for this paper are

BMJ Open Trends in opioid dispensing to injured workers following codeine scheduling changes in Australia: a retrospective cohort study

Michael F Di Donato ⁽¹⁾, ¹ Stephanie Mathieson, ² Giovanni E Ferreira ⁽¹⁾, ^{2,3} Ting Xia ⁽¹⁾, ⁴ Yonas Getaye Tefera, ¹ Christina Abdel Shaheed ⁽¹⁾, ^{2,3} Christopher Maher ⁽¹⁾, ^{2,3} Alex Collie ⁽¹⁾

ABSTRACT

Objectives To describe the prevalence and patterns of opioid analgesic and pain medicine dispenses, and the impact of up-scheduling of low-dose (\leq 15 mg) codeine-containing products to Australians with accepted workers' compensation time loss claims for musculoskeletal conditions between 2010 and 2019.

Design Interrupted time series.

Setting Workers' compensation scheme in Victoria, Australia.

Population Australians with accepted workers' compensation time loss claims for musculoskeletal conditions between 2010 and 2019.

Main outcome measures Number and proportion of workers dispensed pain medicines in the first year of claim and the monthly number, percentage of pain medicine dispenses and mean morphine equivalent dispense dose. Results Nearly one-third (28.4%, n=22807) of our sample of 80 324 workers were dispensed any opioid in the first year since the workers' compensation insurer received their claim. There were no significant step or trend changes in the number or percentage of pain medicines dispensed of up-scheduled low-dose codeine. Only 2.9% of workers were ever dispensed up-scheduled low-dose codeine, specifically 2.5% after up-scheduling (1 February 2018). After up-scheduling of low-dose codeine, workers were more likely to be dispensed opioids (excluding codeine) (prevalence ratio (PR) 1.21, 99% Cl 1.13, 1.31) or other pain medicines (eg, pregabalin, paracetamol) (PR 1.11, 99% CI 1.03, 1.19) compared with the year prior. There was a significant 28.5% (99% Cl 16.3, 41.9) step increase (ie, increase immediately after up-scheduling) in high-dose (>15 mg) codeine with a significant trend decrease (-1.3%, 99% CI -2.5, -0.2). Conclusion Up-scheduling low-dose codeine to prescription-only medicines did not significantly change the dispensing of low-dose codeine-containing products to workers with accepted workers' compensation time loss claims for musculoskeletal conditions.

INTRODUCTION

Recommendations for the use of opioid analgesics have changed in recent years with greater recognition that the risks may

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ Our study used a large sample of detailed claims and medicine data enabling us to gain new insights about a critical policy change.
- ⇒ This is the first study to report on the impact of codeine scheduling changes in the compensable patient population.
- ⇒ Our study is limited by background trends in medicine use, a relatively short 12-month follow-up period, and only including medicines funded by the workers' compensation scheme.

outweigh the potential benefits.¹² These side effects and the risk of opioid overuse and effects and the risk of opioid overuse and **a** overdose have become particularly pertinent **a** issues for policymakers globally. In Australia, **E** this has meant the introduction of real-time prescription drug monitoring programmes³ and restrictions on the access to some training, opioids. Low-dose codeine-containing products (ie, $\leq 15 \text{ mg}$ of codeine per dose unit) have been progressively restricted to reduce harmful use over several years.⁴ Originally available at pharmacies and licensed retailers in Australia, supply was first restricted to overthe-counter purchases (ie, available without a prescription) only at pharmacies in 2010.⁵ A major change was implemented on 1 February 2018, when low-dose codeine products were 'up-scheduled' from Schedule 3 to Schedule 4, restricting access to prescription-only.

Musculoskeletal conditions and injuries are the leading cause of workers' compensation claims in Australia, accounting for half (50.3%, n=64300) of time loss claims in 2021–2022.⁶ Workers' compensation schemes fund reasonable and necessary healthcare, including medicines.⁷ Costs for over-thecounter medicines may be reimbursed to the worker directly. However, prescription

Check for updates

© Author(s) (or their employer(s)) 2025. Re-use permitted under CC BY. Published by BMJ Group.

¹School of Public Health and Preventive Medicine, Monash University, Melbourne, Victoria, Australia

²Sydney Musculoskeletal Health, The University of Sydney, Sydney, New South Wales, Australia

 ³Institute for Musculoskeletal Health, Sydney Local Health District, Camperdown, New South Wales, Australia
 ⁴Monash Addiction Research Centre, Monash University, Melbourne, Victoria, Australia

Correspondence to

Michael F Di Donato; michael.didonato@monash.edu medicines require a consultation with a medical practitioner to first obtain a prescription. While workers' compensation funds this consultation, it is an additional step that may have impacted opioid-seeking behaviour in injured workers.

Several studies have highlighted the impact of up-scheduling low-dose codeine on codeine supply, other opioid supply,^{8 9} overdoses,¹⁰ opioid use disorders¹¹ and emergency department presentations.¹² However, these studies were in the general population. Work-related injuries present a unique set of factors that affect recovery and a funding mechanism for healthcare and medicines that differs from the mainstream universal public health model in Australia.¹³¹⁴ Therefore, we sought to (1) describe the prevalence and patterns of opioid and pain medicine dispensing over time and (2) examine the association between the up-scheduling of low-dose codeine and the number and proportion of opioids and pain medicines, and the dose of opioids, dispensed to injured workers with musculoskeletal conditions.

METHODS Setting

Workers' compensation schemes in each Australian state and territory fund income replacement and healthcare for workers where injury can be attributed to employment.⁷ Approximately 90% of workers in the state of Victoria, Australia, are covered by workers' compensation insurance. Victoria is Australia's second most populous state, with a labour force of 3.2 million people when lowdose codeine was up-scheduled in 2018.¹⁵ The Victorian workers' compensation scheme requires that employers fund the first 10 business days of income replacement and \$700 (AUD) of medical expenses.¹⁶

Data source

We used a sample of the workers' compensation claims data provided by the Victorian workers' compensation scheme regulator, WorkSafe Victoria.¹⁷ Data contained information about the worker (ie, sex, age, occupation, injury details, employer size and type, key dates) and detailed medicine dispense data (ie, medicine type and ingredients, dispense date, dispense dose, cost). A medicine dispense was considered the total number of units (eg, tablets) dispensed on a given date, typically a single packet of medicine, for example, 20 tablets of codeine and paracetamol. Preparing claim data for analysis involved quality assurance of several variables (eg, age, sex, occupation) and joining population socioeconomic (Index of Relative Socio-economic Advantage and Disadvantage (IRSAD)) and remoteness (Accessibility/Remoteness Index of Australia (ARIA)) measures by worker residential postcode.¹⁸¹⁹ Cleaning medicine data involved applying the Anatomical Therapeutic Chemical (ATC) coding scheme²⁰ and checking variables used in the calculation of opioid dose, morphine milligram equivalent dose and the calculation of opioid dose itself (ie, dispense quantity

* item strength * morphine multiplier). Access to data was approved by the Monash University Human Research Ethics Committee (Project ID 30718).

Sample

We included workers with accepted workers' compensation claims for musculoskeletal conditions received by insurers between 1 February 2010 and 31 January 2019. Musculoskeletal conditions were defined by the Type of Occurrence Classification System (TOOCS; online u supplemental table 1).²¹ Only workers with at least 1 day of income replaced by the workers' compensation scheme gewere included. Eligible workers were aged between 15 and 80 years.

copy We included pain medicines defined by ATC level 2 codes relating to the musculoskeletal and nervous systems: M01, M02, M03, N01, N02, N03, N05 and N06 (online supplemental table 2). Included medicines were classified as either up-scheduled low-dose code ($\leq 15 \text{ mg}$), highdose codeine (>15 mg), opioids (excluding codeine) or other pain medicines using ATC codes and item strength (see online supplemental table 3). We included any opioid or other pain medicine dispensed to eligible workers 31 uses rela days before to 365 days after the date the insurer received the workers' compensation claim (ie, their first year of claim). Although rare, healthcare funded by workers' compensation may be retrospectively reimbursed, hence the 31 days prior. Our follow-up data allowed for a 1-year \overline{a} (ie, 365 day) follow-up period. Opioid dispenses with ŧ missing dose information or illogical dose quantities (eg, >240 tablets for codeine and paracetamol) were excluded from analyses. This accounted for approximately 0.59% from analyses. This accounted for approximately 0.59% and of all opioid dispenses and 0.21% of opioid dispenses are accounted for approximately 0.59%included in final interrupted time-series models (online supplemental table 4).

Outcome variables

I training, Changes in the prevalence of any pain medicines dispensed were measured by the number and proportion of workers dispensed each type of pain medicine in the year since an insurer received their claim. Changes in monthly pain medicine dispenses were measured by the similar number of each type of pain medicine, the percentage of pain medicines dispensed and mean morphine equiva-

lent (MME) dispense dose (for opioids).
Analysis
We first grouped workers into 1 year intervals that g aligned with the up-scheduling of low-dose codeine (1 February 2018) by the date the insurer received their claim. That is, each year commenced on 1 February and ended 31 January the following year. For example, if a worker's claim was received by the insurer on 2 January 2013, they would be assigned as a claim commencing in 2012. We used descriptive statistics to report the number and proportion of workers per year who were dispensed each category of pain medicine (online supplemental table 2) at any time in the first year of their claim. We

then used Poisson models to compare the likelihood of being dispensed each of those pain medicines relative to the year prior to up-scheduling (1 February 2017 to 31 January 2018). This allowed us to compare the prevalence of pain medicines in the year after up-scheduling to the year prior, but also to previous years in the sample. Poisson models adjusted for worker sex, age group (ie, 15-24, 25-34, 35-44, 45-54, 55-64 and 65 or more years), employment type (ie, full-time, part-time, casual or other), employer size (ie, government, large, medium or small), nature of injury (TOOCS),²¹ bodily location of injury (TOOCS), occupation (Australian Standard Classification of Occupations major groups),²² socioeconomic status (IRSAD quintiles)¹⁸ and remoteness (ARIA).¹⁹ We used the log of the total number of workers as offsets and robust standard errors in each model. Results were reported as prevalence ratios (PR) with 99% CI (PR 99% CI) and considered statistically significant where p<0.01.

We then visualised the monthly percentages of (1) pain medicines by pain medicine category and (2) the five most frequently dispensed opioids, both over the entire time series (ie, 2010 to 2019).

We used interrupted time-series analyses to measure the impact of the up-scheduling of low-dose codeine on pain medicine dispensing. We selected a time series of medicines dispensed 2 years before to 1 year after lowdose codeine was up-scheduled (ie, dispensed between 1 February 2016 and 31 January 2019). In total, 36 time points were included in the analyses. We used descriptive statistics to report the monthly mean and SD of the number, percentage of pain medicines dispensed and mean dispensed dose of each category of pain medicine. Negative binomial models were used to measure changes in the number of dispenses. The output of negative binomial models was converted to a percentage change (ie, 100 * incidence rate ratio - 100). Generalised least squares (GLS) models were used to measure changes in the rate and mean dose of dispenses. We log-transformed the response variable in GLS models (ie, rate and dispense dose) to make the output a percentage change. We tested for seasonality in all models by adding six sine and six cosine terms to the initial models. Seasonality terms that were significant (p<0.05) were retained in the final models. Autoregressive moving average methods were used to assess and adjust models for autocorrelation and partial autocorrelation.²³ Akaike and Bayesian Information Criteria were used to compare and select the final models. We considered results as statistically significant where p<0.01. We reported results in percentage change in step (ie, immediate change in outcome following the month of up-scheduling) and trend (ie, a change in trend slope following the month of up-scheduling) with 99% CI.^{23 24} Results were visualised with time-series figures reporting the original data, trends before and after the up-scheduling of low-dose codeine and the counterfactual trend after up-scheduling (ie, trend if up-scheduling had not occurred). We performed analyses in RStudio

related to

text

a

tra

using R 4.2.2 and several R packages (online supplemental table 5).

Patient and public involvement

Patients and/or the public were not involved in the design, conduct, reporting or dissemination plans of this research, due to the nature of the de-identified data. We sought expert input on the final manuscript from the workers' compensation regulator, WorkSafe Victoria, prior to submission.

RESULTS

Protected by copyri Our sample included 80324 workers. Nearly one-third (28.4%, n=22807) were dispensed any pharmaceutical .ight opioid analgesic and 25.9% (n=20790) other medicines to manage pain conditions in the first year since the insurer received their claim (see table 1). Specifically, 2.9% (n=2367) were dispensed up-scheduled lowdose codeine, 12.9% (n=10358) high-dose codeine and 22.6% (n=18154) opioids (excluding codeine) (see ð table 2). The proportion of workers who were dispensed each medicine by all available covariates is available in the supplementary materials (online supplemental table 6).

Changes in the prevalence of pain medicines

Opioids (including codeine) and other pain medicines were significantly more prevalent in workers whose claims began in the year after the up-scheduling of low-dose codeine relative to the year prior (see table 1). The lowest proportion of workers dispensed opioids was observed in the year prior to the up-scheduling of codeine. Opioids were significantly more prevalent in workers whose claims began in 2010 and 2011. There were no significant differ- ▶ ences in the prevalence of other pain medicines in any year prior to 2017.

There were no significant differences in the prevalence of up-scheduled low-dose codeine or high-dose codeine ھ in the year after low-dose codeine was up-scheduled. The greatest number and proportion of workers dispensed up-scheduled low-dose codeine and high-dose codeine were in workers whose claims were received by the insurer in 2010 at 3.9% and 16.7%, respectively (see table 2). High-dose codeine was dispensed to a significantly greater percentage of workers with claims commencing between & 2010 and 2015 compared with the reference year (2017). There were no significant differences in the prevalence of opioids (excluding codeine) over the study period, except for those workers whose claims commenced after the up-scheduling of low-dose codeine (PR 1.21, 99% CI 1.13, 1.31). The prevalence of workers whose claims were received by the insurer in this year increased to 26.7% from 22.3% in the previous year. Full models are available in supplementary materials (online supplemental table 7).

texi

ar

data mini

, AI training, and similar technologies

Protected by copyright, including for uses related to

Table 1	Statistical comparison of the prevalence of all opioids and other pain medicines by the year insurers received
workers'	' claims

Year the insurer	Workers All opioids (incl. codeine)		odeine)	Other pain medicines	
received the claim*	N	N (%)	PR (99% CI)†‡	N (%)	PR (99% Cl)†‡
2010	9141	2751 (30.1)	1.13 (1.06, 1.20)§	2426 (26.5)	1.05 (0.98, 1.13)
2011	9011	2616 (29.0)	1.09 (1.02, 1.16)§	2390 (26.5)	1.06 (0.99, 1.13)
2012	9400	2644 (28.1)	1.04 (0.98, 1.11)	2356 (25.1)	0.99 (0.93, 1.07)
2013	8835	2433 (27.5)	1.03 (0.97, 1.10)	2274 (25.7)	1.02 (0.96, 1.10)
2014	8873	2475 (27.9)	1.04 (0.98, 1.11)	2310 (26.0)	1.05 (0.98, 1.12)
2015	8644	2457 (28.4)	1.07 (1.00, 1.14)	2309 (26.7)	1.08 (1.01, 1.16)
2016	8680	2347 (27.0)	1.01 (0.94, 1.08)	2202 (25.4)	1.02 (0.95, 1.10)
2017	8749	2310 (26.4)	1.00 (ref)	2113 (24.2)	1.00 (ref)
2018	8991	2774 (30.9)	1.18 (1.11, 1.26)§	2410 (26.8)	1.11 (1.03, 1.19)§
All years	80324	22807 (28.4)	-	20790 (25.9)	-

*Year that the insurer received the claim, where each year commenced on 1 February and ended on 31 January.

†Prevalence ratio and 99% Cl.

*Poisson model adjusted for worker sex, age group, employment type, employer size, nature of injury, bodily location of injury, occupation, socioeconomic status and remoteness. Full models available in supplementary materials (online supplemental table 7).

\$p<0.01

PR, prevalence ratio.

Changes in pain medicine dispensing

Opioid analgesics were a common pain medicine dispensed throughout the sample period (figure 1). Up-scheduled low-dose codeine contributed to a consistently small proportion of pain medicines dispensed. High-dose codeine products appeared to decrease over the 10 year sample period.

The five most frequent opioids dispensed fluctuated over the study period. Single-ingredient oxycodone and codeine in combination with paracetamol were the most frequent opioids throughout the sample period. A complete list of the most frequent pain medicines and pain medicine strengths is available in the supplementary materials (online supplemental table 8). Figure 1 also highlights the up-scheduling of low-dose codeine in February 2018 (dotted line) and 3 years of data used in the interrupted time series (highlighted segment).

There were no significant step or trend changes in the number of up-scheduled low-dose codeine dispenses or the percentage of pain medicines that were up-scheduled ō low-dose codeine (see table 3 and figure 2). However, there was a significant 18.5% (99% CI -27.7, -12.7%) step decrease in mean dispense dose at the month of up-scheduling implementation, but with a significant

Year the	Workers	Up-schedu	Up-scheduled low-dose codeine		High-dose codeine		Opioids (excl. codeine)	
insurer received the claim*	N	N (%)	PR (99% Cl)†‡	N (%)	PR (99% CI)†‡	N (%)	PR (99% Cl)†‡	
2010	9141	359 (3.9)	1.52 (1.21, 1.92)§	1530 (16.7)	1.75 (1.57, 1.94)§	1999 (21.9)	0.97 (0.90, 1.05)	
2011	9011	340 (3.8)	1.48 (1.17, 1.87)§	1382 (15.3)	1.61 (1.44, 1.80)§	1996 (22.2)	0.98 (0.91, 1.06)	
2012	9400	313 (3.3)	1.27 (1.00, 1.61)	1370 (14.6)	1.52 (1.36, 1.69)§	1978 (21.0)	0.92 (0.86, 1.00)	
2013	8835	263 (3.0)	1.16 (0.91, 1.49)	1197 (13.5)	1.42 (1.27, 1.60)§	1887 (21.4)	0.94 (0.88, 1.02)	
2014	8873	210 (2.4)	0.93 (0.72, 1.21)	1194 (13.5)	1.41 (1.26, 1.58)§	1964 (22.1)	0.98 (0.91, 1.06)	
2015	8644	222 (2.6)	1.00 (0.78, 1.30)	1053 (12.2)	1.29 (1.14, 1.45)§	1996 (23.1)	1.03 (0.96, 1.11)	
2016	8680	228 (2.6)	1.05 (0.82, 1.36)	909 (10.5)	1.10 (0.97, 1.24)	1984 (22.9)	1.01 (0.94, 1.09)	
2017	8749	210 (2.4)	1.00 (ref)	815 (9.3)	1.00 (ref)	1947 (22.3)	1.00 (ref)	
2018	8991	222 (2.5)	1.05 (0.82, 1.36)	908 (10.1)	1.09 (0.96, 1.23)	2403 (26.7)	1.21 (1.13, 1.31)§	
All years	80324	2367 (2.9)	-	10358 (12.9)	-	18154 (22.6)	-	

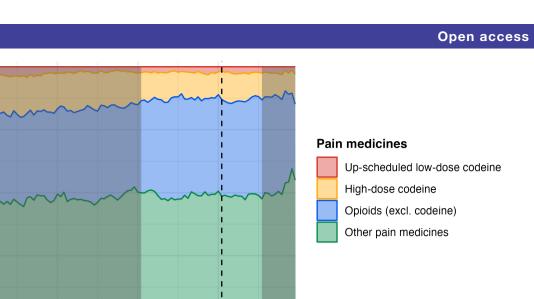
*Year that the insurer received the claim, where each year commenced on 1 February and ended on 31 January.

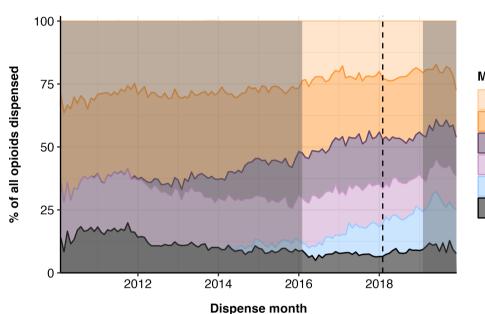
+Prevalence ratio and 99% CI.

Poisson model adjusted for worker sex, age group, employment type, employer size, nature of injury, bodily location of injury, occupation, socioeconomic status and remoteness. Full models available in supplementary materials

§p<0.01.

PR. prevalence ratio





2014

2016

2018

6

% of pain medicines dispensed

100

75

50

25

0

2012

Most frequent opioids (incl. codeine)

Codeine and paracetamol (N02AJ06) Oxycodone (N02AA05) Oxycodone and naloxone (N02AA55) Tramadol (N02AX02) Tapentadol (N02AX06) All other opioids

Figure 1 Trends in pain medicines and most frequent opioids throughout the sample period. Dotted line indicates upscheduling of codeine; highlighted period indicates time series (ie, 2016 to 2019) used in interrupted time-series analyses.

trend increase of 2.2% per month following the up-scheduling implementation (99% CI 0.8%, 3.6%). We observed a significant 32.3% (99% CI 17.4%, 49.0%) step increase in the number of high-dose codeine dispenses following up-scheduling of low-dose codeine. This was accompanied by a 28.5% (99% CI 16.3%, 41.9%) step increase in the percentage of high-dose codeine dispenses that were pain medicines but, with a significant trend decrease of -1.3% (99% CI -2.5%, -0.1%). There were significant trend increases in the number of opioids (excluding codeine) (1.4%, 99% CI 0.0%, 2.8%) and other pain medicines (2.0%, 99% CI 1.0%, 3.1%), but not in the percentage of pain medicines. Finally, there was a significant -12.3% (99% CI -19.0%, -5.1%) step decrease in the mean dispense dose of opioids (excluding codeine) following the up-scheduling of low-dose codeine.

DISCUSSION

The up-scheduling of low-dose codeine did not signifitechnol cantly change the prevalence, monthly number or monthly percentage of low-dose codeine-containing products dispensed to injured Australian workers with workers' compensation claims for musculoskeletal conditions. Less than 3% of workers were ever dispensed $\overline{\mathbf{g}}$ up-scheduled low-dose codeine compared with the nearly third ever dispensed other types of opioids. There was a significant step increase in the monthly number and percentage of high-dose codeine dispenses at the time of low-dose codeine up-scheduling. This was accompanied by a significant decreasing trend over the subsequent year, and high-dose codeine was significantly less prevalent in workers whose claims commenced every year after 2010. Opioids other than codeine were significantly more

Table 3 Results of interrupted time series

	Monthly mean (SD) values		Step change	Trend change
	2 years before*	1 year after†	% (99% CI)‡	% (99% CI)
N of dispenses				
Up-scheduled low-dose codeine	55.5 (8.7)	58.8 (12.5)	-7.9% (-29.4%, 19.5%)	2.3% (-0.9%, 5.6%)
High-dose codeine	259.8 (36.4)	265.5 (31.7)	32.3% (17.4%, 49.0%)§	0.0% (-1.4%, 1.5%)
Opioids (excl. codeine)	948.5 (89.6)	924.7 (60.7)	1.1% (–9.9%, 13.5%)	1.4% (0.0%, 2.8%)§
Other pain medicine	1213.4 (118.1)	1188.8 (89.9)	-3.6% (-12.1%, 5.9%)	2.0% (1.0%, 3.1%)§
% of pain medicines dispensed				
Up-scheduled low-dose codeine	2.2 (0.4)	2.4 (0.4)	-13.0% (-32.8%, 12.7%)	1.0% (-2.1%, 4.2%)
High-dose codeine	10.5 (0.8)	10.9 (1.0)	28.5% (16.3%, 41.9%)§	-1.3% (-2.5%, -0.2%)§
Opioids (excl. codeine)	38.3 (1.6)	38.0 (0.8)	-4.9% (-10.9%, 1.5%)	0.2% (-0.6%, 1.0%)
Other pain medicine	49.0 (1.3)	48.8 (0.7)	-4.4% (-9.6%, 1.1%)	0.3% (-0.8%, 1.3%)
Mean dispense dose (MME)				
Up-scheduled low-dose codeine	58.4 (3.7)	55.2 (4.1)	-18.5% (-27.7%, -8.0%)§	2.2% (0.8%, 3.6%)§
High-dose codeine	167.2 (6.8)	155.3 (9.3)	-8.9% (-18.0%, 1.2%)	0.6% (-0.7%, 1.9%)
Opioids (excl. codeine)	529.2 (31.6)	519.1 (25.5)	–12.3% (–19.0%, –5.1%)§	0.0% (-1.0%, 1.0%)

*2 years/24 months before the up-scheduling of codeine (February 2016 to January 2018). †1 year/12 months after the up-scheduling of codeine (February 2018 to February 2019). ‡Percentage change (99% CI).

prevalent in those workers whose claims commenced after up-scheduling. However, there was also a significant step decrease in the monthly mean dose per opioid dispensed at the time of up-scheduling.

We did not observe the same significant decreases in low-dose codeine dispenses following up-scheduling in our sample as in other Australian studies.^{8-11 25} This

lack of change in our sample could be related to several factors. First, up-scheduled low-dose codeine products were available over the counter for many years at a relatively low cost, so it is possible that few injured workers claimed reimbursement from the workers' compensation scheme. Second, while claims for prescription medicines are lodged by pharmacists with detailed medicine

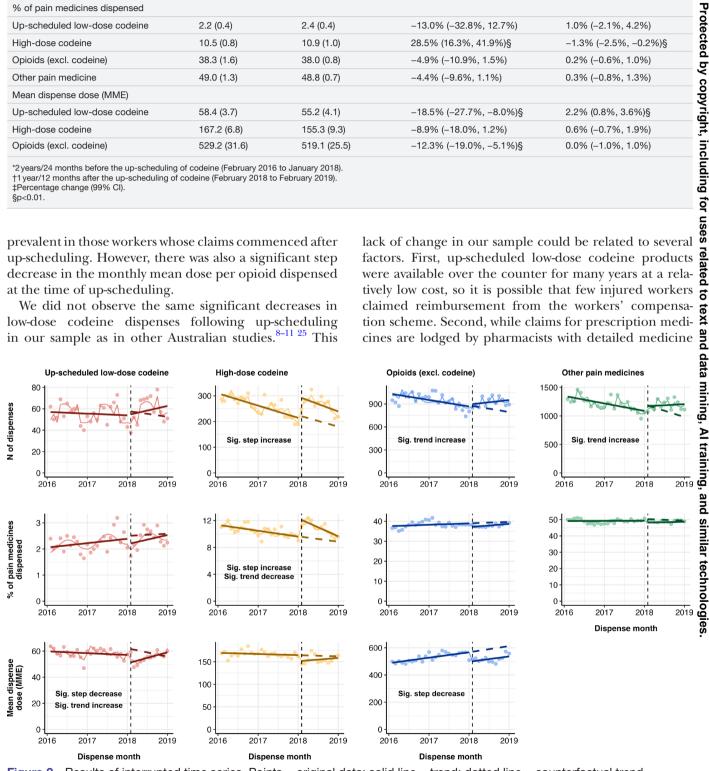


Figure 2 Results of interrupted time series. Points = original data; solid line = trend; dotted line = counterfactual trend.

^{\$}p<0.01.

data

recording forms, over-the-counter medicines are not necessarily recorded in detail and are often reported as 'over-the-counter medicine'; moving to a prescriptiononly medicine may have improved data quality. Finally, we included a sample of specific musculoskeletal conditions as opposed to an entire population. Our sample also represents workers with at least 2weeks of absence from work who likely have more severe injuries.

Our findings suggest that the up-scheduling of low-dose codeine may have shifted workers already being dispensed low-dose codeine to higher doses of codeine and workers commencing claims after the up-scheduling of codeine to other opioids and other pain medicines. The potential for these shifts was raised as a concern by consumers and healthcare providers prior to up-scheduling.²⁶ First, there was a significant, but temporary, increase in highdose codeine dispenses that were ultimately less prevalent in more recent claims. Some individuals may have been purchasing numerous packs of low-dose codeine medicines per week out of pocket, which would not be detected in our data. It is unlikely that these workers would have attended a doctor multiple times per week for multiple prescriptions and may have been prescribed larger dispenses. This may be one reason behind the temporary significant increase in high-dose codeine dispenses and seems to point to workers who may have been seeking low-dose codeine prior to up-scheduling to be prescribed high-dose codeine or lower doses of other opioids if they are required to obtain a prescription either way. Workers who commenced their claim after low-dose codeine up-scheduling appear to have avoided low- and high-dose codeine products, with significant monthly trend increases and a greater prevalence of workers dispensed opioids and other pain medicines; the latter of which aligns with the findings of Schaffer et al.⁸

Implications for policy and practice

Our findings indicate that a consistent proportion of workers with workers' compensation time loss claims for musculoskeletal conditions continue to be prescribed pain medicines. This is troubling considering the negative disability, health and cost outcomes associated with opioids specifically for compensated workers who likely have long-term pain problems.²⁷⁻³⁰ The changes and trends highlighted in our study indicate that these workers have progressively shifted to a more diverse array of opioids and other pain medicines. The Victoria workers' compensation authority has a clinical panel that conducts internal reviews of healthcare and pharmaceutical treatment of injured workers. However, it is unclear what relation the workers' compensation scheme has with the mandatory prescription-drug monitoring programme (PDMP) that was implemented in April 2020 in Victoria (the state of this study).³¹ Workers' compensation schemes could consider their own systems to flag certain types and doses of medicines dispensed to injured workers to supplement a PDMP if they have not already.

As evidenced by our analysis, the raw data exists to achieve this.

Strengths, limitations and future research

Our study benefited from a large sample of detailed data, enabling us to gain new insights into the impact of a critical policy change. Detailed medicines data linked with detailed worker data allowed us to adjust for numerous important covariates. Limitations of our study should be considered. First, employers must fund the first \$700 (AUD) of medical expenses in the Victorian workers' compensation scheme,¹⁶ which is not recorded in the administrative data. Pain medicines are relatively inexpensive in Australia, and so it is likely that we are missing **J** at least some medicines. Second, we only included medi-8 cines funded by the workers' compensation scheme. It is possible that over-the-counter products like low-dose codeine were purchased out-of-pocket and subsequently not detected in our data. Thirdly, background trends in medicines dispensed may have confounded interrupted time-series analyses. For example, there was an apparent rise in the use of tapentadol in the years before and after ğ the up-scheduling of codeine that may have contributed uses r to observed changes in the mean monthly dose of opioids. Finally, we only included a follow-up period of 1 year since the insurer received the claim, missing potentially longerterm trends in pain medicine dispensing. Workers are eligible for healthcare funding for up to 52 weeks after đ income replacement ends at 130 weeks in the Victorian text and workers' compensation scheme. Future research could, therefore, consider follow-up periods of up to 3.5 years.¹⁶

Conclusions

Up-scheduling codeine did not significantly change the Ξ dispensing of low-dose codeine-containing products to Australian workers with accepted workers' compensation time loss claims for musculoskeletal conditions. A tempo- ≥ rary increase in high-dose codeine, step decrease in mean opioid dispense dose and trend increases in other opioids and other pain medicines appear to indicate a shift away from any dose of codeine to lower dose opioids and a other analgesics, such as pregabalin. Workers' compensasimilar tion schemes could consider utilising detailed medicine data to monitor medicine dispenses as a supplement to prescription drug monitoring programmes.

Acknowledgements WorkSafe Victoria supplied data for use in this study. The views expressed in this paper are those of the authors and are not necessarily the views of WorkSafe Victoria. The authors would like to thank Dr Tyler J Lane for his assistance with the interrupted time series analysis.

Contributors MFDD, SM and AC conceived the study. MFDD cleaned and prepared data with assistance from YGT, SM and CAS. MFDD conducted analyses. MFDD drafted the manuscript. All authors reviewed the manuscript and provided input to the interpretation of results. All authors approved the final manuscript. MFDD is the guarantor.

Funding This study was supported by the National Health and Medical Research Council (NHMRC) Centre for Research Excellence on Low Back Pain (ANZBACK) (APP1171459). MDD is supported by a post-doctoral fellowship from ANZBACK and

X Giovanni E Ferreira @giovanni_ef, Christopher Maher @CGMMaher and Alex Collie @axcollie

a project grant from the New South Wales State Insurance Regulatory Authority. GF is supported by an Emerging Leadership Investigator Fellowship from the National Health and Medical Research Council (APP2009808). AC is supported by an Australian Research Council Future Fellowship (FT19010218). CM is supported by a National Health and Medical Research Council (NHMRC) Investigator Fellowship (APP1194283).

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

Patient consent for publication Not applicable.

Ethics approval This study received ethics approval from the Monash University Human Research Ethics Committee (Project ID 30718). Participant data were de-identified. Workers consent to use of data for research purposes when entering each of the workers' compensation schemes.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data may be obtained from a third party and are not publicly available. WorkSafe Victoria supplied data for use in this study. Data used in this paper are not available for distribution by the authors. The R analysis code used in analyses is available upon request.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/ licenses/by/4.0/.

ORCID iDs

Michael F Di Donato http://orcid.org/0000-0002-6531-5949 Giovanni E Ferreira http://orcid.org/0000-0002-8534-195X Ting Xia http://orcid.org/0000-0001-5033-6248 Christina Abdel Shaheed http://orcid.org/0000-0003-1258-5125 Christopher Maher http://orcid.org/0000-0002-1628-7857 Alex Collie http://orcid.org/0000-0003-2617-9339

REFERENCES

- Australian Commission on Safety and Quality in Health Care (ACSQHC). *Low back pain clinical care standard*. ACSQHC: Sydney, Australia, 2022.
- 2 Dowell D, Haegerich TM, Chou R. CDC Guideline for Prescribing Opioids for Chronic Pain - United States, 2016. *MMWR Recomm Rep* 2016;65:1–49.
- 3 Australian Government Department of Health. National real time prescription monitoring. 2023. Available: https://www.health.gov.au/ our-work/national-real-time-prescription-monitoring-rtpm
- 4 Australian government department of health therapeutic goods administration. Final decisions and reasons for decisions by delegates of the secretary to the department of health. Canberra, Australia Australian Government Department of Health; 2017.
- 5 Therapeutic Goods Administration (TGA). Regulation impact statement: codeine re-scheduling. Canberra, Australia Therapeutic Goods Administration; 2016.
- 6 Safe Work Australia (SWA). Key work health and safety statistics Australia 2023. Canberra, Australia Safe Work Australia; 2023.
- Collie A, Di Donato M, Iles R. Work Disability in Australia, 2023
 Collie A, Di Donato M, Iles R. Work Disability in Australia: An Overview of Prevalence, Expenditure, Support Systems and Services. J Occup Rehabil 2019;29:526–39.
 Schaffer AL Coirce P. Berger M. Schaffer AL Coirce P. Destruction of the State Science Sci
- 8 Schaffer AL, Cairns R, Brown JA, *et al*. Changes in sales of analgesics to pharmacies after codeine was rescheduled as a prescription only medicine. *Med J Aust* 2020;212:321–7.

- 9 Yu Y, Wilson M, King CE, et al. Up-scheduling and codeine supply in Australia: analysing the intervention and outliers. Addiction 2021;116:3463–72.
- Cairns R, Schaffer AL, Brown JA, *et al*. Codeine use and harms in Australia: evaluating the effects of re-scheduling. *Addiction* 2020;115:451–9.
 McCont L Nickson C. Control Contr
- McCoy J, Nielsen S, Bruno R. A prospective cohort study evaluating the impact of upscheduling codeine in Australia among frequent users of codeine. *Addiction* 2022;117:677–86.
 Elebertor DA C
- 12 Elphinston RA, Connor JP, de Andrade D, *et al.* Impact of a policy change restricting access to codeine on prescription opioid-related emergency department presentations: an interrupted time series analysis. *Pain* 2021;162:1095–103.
- Collie A, Sheehan L, Di Donato M. Variation in General Practice Services Provided to Australian Workers with Low Back Pain: A Cross-Jurisdictional Comparative Study. *J Occup Rehabil* 2022;32:203–14.
- Di Donato M, Iles R, Lane T, *et al*. The impact of income support systems on healthcare quality and functional capacity in workers with low back pain: a realist review. *Pain* 2020;161:2690–709.
- Australian Bureau of Statistics (ABS). 6202.0 labour force, Australia, Feb 2018. Canberra Australian Bureau of Statistics; 2023.
 Statistics (ABS). 6202.0 - labour force, Australia, Feb 2018. Canberra Australian Bureau of Statistics; 2023.
- 16 Safe Work Australia (SWA). Comparison of workers' compensation arrangements in Australia and New Zealand. Canberra, Australia, 2021.
- 17 Prang K-H, Hassani-Mahmooei B, Collie A. Compensation Research Database: population-based injury data for surveillance, linkage and mining. *BMC Res Notes* 2016;9:456.
- 18 Australian Bureau of Statistics (ABS). Socio-economic indexes for areas. 2023. Available: https://www.abs.gov.au/websitedbs/ censushome.nsf/home/seifa
- 19 Australian Bureau of Statistics (ABS). Remoteness areas Australian statistical geography standard (ASGS) edition 3. 2023. Available: https://www.abs.gov.au/statistics/standards/australian-statisticalgeography-standard-asgs-edition-3/jul2021-jun2026/remotenessstructure/remoteness-areas
- 20 World Health Organisation (WHO) Collaborating Centre for Drug Statistics Methodology. ATC / DDD index 2020. 2022. Available: https://www.whocc.no/atc_ddd_index
- 21 Australian Safety and Compensation Council (ASCC). *Type of occurrence classification system (revision one)*. 3rd edn. Canberra, Australia: ASCC, 2008.
- 22 Australian Bureau of Statistics (ABS). *1220.0 ASCO Australian standard classification of occupations, 1997.* 2nd edn. Canberra, Australia, 1997.
- 23 Schaffer AL, Dobbins TA, Pearson S-A. Interrupted time series analysis using autoregressive integrated moving average (ARIMA) models: a guide for evaluating large-scale health interventions. *BMC Med Res Methodol* 2021;21:58.
- 24 Lane TJ, Gray SE, Sheehan L, et al. Increased Benefit Generosity and the Impact on Workers' Compensation Claiming Behavior: An Interrupted Time Series Study in Victoria, Australia. J Occup Environ Med 2019;61:e82–90.
- 25 Middleton M, Nielsen S. Changes in Australian prescription opioid use following codeine rescheduling: A retrospective study using pharmaceutical benefits data. *Int J Drug Policy* 2019;74:170–3.
- McCoy J, Bruno R, Nielsen S. Attitudes in Australia on the upscheduling of over-the-counter codeine to a prescription-only medication. *Drug Alcohol Rev* 2018;37:257–61.
- 27 Di Donato M, Xia T, Iles R, *et al.* Patterns of opioid dispensing and associated wage replacement duration in workers with accepted claims for low back pain: a retrospective cohort study. *Pain* 2022;163:e942–52.
- 28 Franklin GM, Rahman EA, Turner JA, et al. Opioid use for chronic low back pain: A prospective, population-based study among injured workers in Washington state, 2002-2005. *Clin J Pain* 2009;25:743–51.
- 29 Franklin GM, Stover BD, Turner JA, et al. Early opioid prescription and subsequent disability among workers with back injuries: the Disability Risk Identification Study Cohort. *Spine (Phila Pa 1976)* 2008;33:199–204.
- Webster BS, Verma SK, Gatchel RJ. Relationship between early opioid prescribing for acute occupational low back pain and disability duration, medical costs, subsequent surgery and late opioid use. *Spine (Phila Pa 1976)* 2007;32:2127–32.
- use. Spirie (rmia ra 19/6) 2007;32:2127-32.
 Victorian Government Department of Health & Human Services. SafeScript for patients and familes. 2020. Available: https://www2. health.vic.gov.au/public-health/drugs-and-poisons/safescript/ consumers-and-families

for uses related to text and data mining, AI training, and similar technologies

Protected by copyright, including