# **BMJ Open**

The association of physical therapy, chiropractic care and early opioid prescription for acute low back pain with Workers' Compensation claim duration: An observational cohort study

Journal:	BMJ Open		
Manuscript ID:	: bmjopen-2015-007836		
Article Type:	Research		
Date Submitted by the Author:	30-Jan-2015		
Complete List of Authors:	Busse, Jason; McMaster University, Anesthesia; McMaster University, Clinical Epidemiology & Biostatistics Ebrahim, Shanil; Mcmaster University, Clinical Epidemiology and Biostatistics Heels-Ansdell, Diane; Mcmaster University, Clinical Epidemiology and Biostatistics Wang, Li; McMaster University, Anesthesia Walter, Stephen; Mcmaster University, Clinical Epidemiology and Biostatistics		
<b>Primary Subject Heading</b> :	Rehabilitation medicine		
Secondary Subject Heading:	Complementary medicine, Evidence based practice		
Keywords:	COMPLEMENTARY MEDICINE, Pain management < ANAESTHETICS, OCCUPATIONAL & INDUSTRIAL MEDICINE		
	•		

SCHOLARONE™ Manuscripts

The association of physical therapy, chiropractic care and early opioid prescription for acute low back pain with Workers' Compensation claim duration: An observational cohort study

Jason W. Busse, 1,2,3 \* Shanil Ebrahim, 2,3,4,5 Diane Heels-Ansdell, 1 Li Wang,2 Stephen D. Walter 3

<sup>\*</sup>Corresponding Author: email: bussejw@mcmaster.ca

### **Abstract**

**Objective:** To assess the association between reimbursement for physiotherapy, chiropractic, and early opioid prescription for low back pain (LBP) with disability claim duration.

**Design:** Observational cohort study.

**Setting and participants:** From a random sample of 6,665 claims for acute, uncomplicated LBP approved by the Ontario Workplace Safety and Insurance Board (WSIB) in 2005, we analyzed 1,442 that remained on full benefits at four weeks.

Primary and secondary outcome measures: Our primary outcome was WSIB claim duration.

Results: In our adjusted analysis, older age (e.g. hazard ratio [HR] for age ≥55 versus <25 = 0.52;

99% confidence interval [CI] = 0.36, 0.74) and WSIB-reimbursement for opioid prescription in the first 4 weeks of a claim (HR = 0.69; 99% CI = 0.53, 0.89) were associated with longer claim duration. Higher pre-disability income was associated with longer claim duration, but only among persistent claims (e.g. HR for active claims at 1-year with a pre-disability income

>\$902/week versus ≤\$480 = 0.32; 99% CI = 0.16, 0.63). Missing data for union membership (HR = 1.27; 99% CI = 1.01, 1.60), and working for an employer with a return-to-work program was associated with fewer days on claim (HR = 1.77; 99% CI = 1.45, 2.18). Neither reimbursement for physiotherapy (HR = 0.99; 99% CI = 0.85, 1.16) or chiropractic care (HR = 1.08; 99% CI = 0.91, 1.29) were associated with claim duration.

**Conclusions:** Our analysis found that treatment commonly reimbursed by the Ontario WSIB for disabling acute LBP does not affect, or prolongs, claim duration. Well-designed randomized controlled trials are needed to confirm or refute our findings.

Predicting Workers' Compensation claim duration

**Key Words:** Low back pain; disability; compensation; Workers Compensation Board; physical therapy; chiropractic; opioid; survival analysis

Predicting Workers' Compensation claim duration

#### **Strengths and Limitations**

- A priori creation of our regression model and the anticipated direction of included independent variables, and assessment of the proportional hazards assumption for all independent variables, provide greater confidence in our findings.
- The reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are
  uncertain, and despite our adjustments for potential confounders it remains possible
  that LBP claimants chosen to receive these healthcare interventions were prognostically
  different than those claimants who did not.
- A number of variables that may be important to consider were unavailable (e.g., patient expectations regarding recovery), and some variables were not optimally collected. For example, chiropractic and physiotherapy are professions and not modalities, and there were no details of treatment provided.
- Our primary outcome, claim closure, is a surrogate for patient-important outcomes such as functional restoration or return to work.

### Introduction

Back pain is a common problem among working adults in North America, with a lifetime prevalence of 63% and a point prevalence of 21%. After the common cold, low back pain (LBP) is the most frequent cause of lost time from work. Globally, LBP is the primary cause of years lived with disability.

In Canada, annual medical expenditures for LBP are estimated to be between \$6 and \$12 billion, with additional costs associated with loss in worker productivity from time off work and associated disability payments. Canadian workers who are disabled secondary to a work-related LBP injury are typically eligible for wage replacement benefits through their provincial Workers' Compensation Board.

In 2012, the Ontario Workplace Safety and Insurance Board (WSIB) approved approximately 233,000 claims that were associated with \$2,918 million in payments, most commonly for strains and sprains affecting the low back. <sup>5,6</sup> The WSIB's liability associated with disability claims greatly exceeds their assets, and as of March 31, 2013 the WSIB's unfunded liabilities were \$12.4 billion <sup>7</sup> – more than double their unfunded liability of \$5.9 billion in 2006

Interventions that are commonly reimbursed by WSIB for LBP claims include physiotherapy, chiropractic care and opioids; however, there is limited evidence to inform their effectiveness. There are no randomized controlled trials of these interventions focused on workers disabled by LBP, and many trials that enroll LBP patients use receipt of disability benefits as an exclusion criterion because of concerns that secondary gain (e.g., receipt of financial compensation conditional on disability) will reduce the treatment effect of study

Predicting Workers' Compensation claim duration

interventions. There is indirect evidence for this hypothesis from surgical research: a recent meta-analysis of 129 studies revealed that the odds of an unsatisfactory outcome in surgical patients receiving disability benefits or engaged in litigation was 3.79 times greater (95% confidence interval [CI]: 3.28 to 4.37) than similar patients not receiving disability benefits or pursuing litigation.<sup>9</sup>

Using administrative data from the Ontario WSIB, we evaluated the association between reimbursement for physiotherapy, chiropractic care, or early prescription for opioids for uncomplicated LBP with disability claim duration. We reported our findings in concordance with the STROBE <sup>10</sup> and TRIPOD <sup>11</sup> statements.

### Methods

### Patient characteristics and eligibility criteria

We enrolled an inception cohort of workers with uncomplicated low back injuries (i.e., strain or sprain) who were fully disabled from working and receiving wage replacement benefits from the Ontario WSIB. We excluded workers if they were approved for no-lost-time claims. The prognosis for chronic LBP (duration >12 weeks) is different than the prognosis for acute LBP, <sup>12</sup> and we excluded all claims in which the number of days between accident date and registration date of the claim was greater than 30 days. Most LBP claims resolve within the 1<sup>st</sup> month, <sup>13</sup> and Workers' Compensation Boards are primarily interested in factors that predict claim resolution among claimants that remain disabled after this time. Further, the Ontario WSIB administrative database we used to acquire our model variables consists of scanned forms, and all data had to be manually extracted by reviewers. To increase the feasibility of our study we excluded claims that ended before 4 weeks.

Between January 1 and June 30, 2005, the Ontario WSIB approved 18,974 lost-time claims for an uncomplicated back injury. Using the WSIB's administrative database, we acquired a random sample of 6,665 injured workers from this population; 1,442 remained on full benefits at four weeks and provided data for our analysis. The WSIB database recorded benefit status for two years after the first day of injury. Patient information was anonymized and de-identified prior to analysis. The Health Sciences Research Ethics Board of the University of Toronto approved our study protocol.

#### **Administrative Variables**

The WSIB database consisted of demographic, administrative, and clinical information, which we acquired from forms completed by the worker, their employer, and their primary health-care provider. The employer form (Employer's Report of Injury/Disease Form; Form 7) is mandatory and must be submitted within three days of a work-related injury. The worker may elect to fill out a form (Worker's Report of Injury/Disease; Form 6) on a voluntary basis if they have expenses related to their injury. The healthcare provider can elect to complete a form (Health Professional's Report; Form 8) to support their patient's claim that their injury is work-related, which is a pre-requisite for wage replacement benefits through the WSIB. Healthcare providers are asked to complete and submit a Functional Abilities Form for Planning Early and Safe Return to Work for each claim, and the WSIB provides reimbursement as incentive.

In order to increase confidence in our findings, we constructed our regression model before conducting any analyses. Guided by the results from our ongoing systematic review of observational studies evaluating predictors of recovery in patients receiving disability benefits,<sup>14</sup> feedback from administrators at WSIB, and content experts within our research team, we selected, *a priori*, 11 variables from the WSIB database that we judged may be associated with claim closure and predicted the direction of anticipated effects on claim duration (Table S1 in File S1). To avoid overfitting our models, we required at least 10 observations per variable term for our Cox regression model, for a total of 190 disabled workers.<sup>15</sup>

#### **Data Extraction**

Predicting Workers' Compensation claim duration

The Ontario WSIB's database consists of scanned paper documents, and data must therefore be extracted manually for analysis. Two experienced reviewers extracted data, independently and in duplicate, from the first 100 eligible claims into an Access database. In order to minimize data entry mistakes, we developed data entry forms that included range checks and missing value alerts. The PROC COMPARE procedure in SAS version 9.2 (SAS Institute Inc., Cary, NC) revealed 98% agreement for the initial 100 claims; therefore, to increase feasibility, only a single abstractor completed data entry for the remaining claims used in our analysis.

We screened all data to identify outliers, inconsistencies and missing data by calculating summary statistics, and explored distributions graphically. We worked with WSIB representatives to correct identified outliers and inconsistencies. If inconsistencies could not be corrected, we treated them as missing data. Some WSIB forms are voluntary, and we included "missing data" as a discrete category for independent variables when applicable.

### Statistical analysis

We generated frequencies for all collected data. We reported the mean and standard deviation (SD) of continuous variables, and the number of occurrences with proportions represented as percentages for categorical variables. Neither age nor pre-disability gross income were normally distributed and were therefore entered as categorical variables into our regression model; by decade for age, starting at age 15 and ending at age 65, and by quartiles for pre-disability income.

Our primary outcome was time to claim closure, defined as the duration from disability claim approval until the claim was closed. We performed a time-to-event analysis using a Cox

Predicting Workers' Compensation claim duration

proportional hazards regression model to assess the association between time to claim closure and the independent variables. We treated receipt of WSIB-reimbursed chiropractic care or physiotherapy as time-dependent covariates to account for when treatments were initiated during the course of the disability claim.

We tested for collinearity to assess if an independent variable was highly correlated with another (correlation coefficient r>0.5) using a correlation matrix. If two variables were highly correlated, we removed the variable that was considered to be of lesser importance, as guided by content experts on our team.

For claims that were unresolved when the data was extracted, we used the date of data extraction as our censoring point. In order to be more stringent and minimize the likelihood of spurious findings, an independent variable was considered statistically significant if it had a p-value ≤0.01 in our adjusted model. We calculated adjusted hazard ratios (HRs) for our time-to-event analyses, their associated 99% confidence intervals (CIs), and the associated p-values. We assessed each independent variable in our model to ensure that the proportional hazards assumption was met by entering each variable in the model separately and calculating the interaction with time. We considered a p-value of ≤0.01 for the interaction term as significant to account for multiple comparisons. We reported the HRs for independent variables that violated the proportional hazards assumption at 30-days (baseline), 6-months, and 1-year.

### Pooling data from similar studies

When possible, we pooled the association between physiotherapy, chiropractic care, or early opioid use and claim duration in our sample with similar data from observational studies

Predicting Workers' Compensation claim duration identified through an ongoing systematic review. We considered studies to be similar if they enrolled Workers' Compensation patients with uncomplicated LBP and, in the case of physiotherapy or chiropractic care, entered these variables as time-dependent covariates. When necessary, we converted odds ratios measures to a relative risk (RR), then to a HR, using the following formula: 16

$$RR = OR/(1-P_0 + P_0 \times OR)$$

$$HR = (ln(1-RR \times P_0))/(ln(1-P_0))$$

P<sub>0</sub> is the proportion of patients in the control group who had an event by the follow-up time

We used random-effects meta-analyses, which are conservative in that they take both within-study and between-study variability into account. We examined heterogeneity using both a chi-squared test and the I² statistic, the latter being the percentage of the total variation in outcomes that is associated with between-study variability (i.e., true differences between studies rather than with sampling error (chance)). Heterogeneity of 0% to 40% was considered 'might not be important', 30% to 60% to be 'moderate heterogeneity', 50% to 90% to be 'substantial heterogeneity', and 75% to 100% to be 'considerable heterogeneity'. The Cochrane Collaboration has proposed overlapping categories to convey that there are no strict cut-offs for interpreting heterogeneity, and this decision will depend on the magnitude and direction of effects, and the strength of evidence for heterogeneity.

### Results

Table 1 presents the baseline characteristics for the 1,442 disabled workers included in our analysis. At some point during their claim, the Ontario WSIB reimbursed 786 (55%) claimants for physiotherapy and 391 (27%) for chiropractic care; approximately 9% of claimants (n=136) were reimbursed for an opioid prescription in the first 4 weeks of their claim. 1348 (93.5%) claims were closed prior to 2 years and 94 (6.5%) were censored. Figure 1 presents the Kaplan-Meier curve for time to claim closure for LBP claimants.

We did not identify any significant collinearity among independent variables in our regression model. Our adjusted regression analysis showed that older age (e.g., HR for age ≥55 versus <25 = 0.52; 99% CI = 0.36, 0.74) and opioid prescription reimbursed by the Ontario WSIB in the first 4-weeks of claim (HR = 0.69; 99% CI = 0.53, 0.89) were associated with longer claim duration. In other words, at any point in time, claimants who were reimbursed for opioids within the first 4 weeks of their claim were 31% less likely to resolve their claim within 2 years versus claimants who were not reimbursed for opioids. The association of pre-disability income with claim duration was not proportional over time, and older claims showed a significant association with longer claim duration (e.g. HR for active claims at 1-year with a pre-disability income >\$902/week = 0.32; 99% CI = 0.16, 0.63). Working for an employer that had a return-towork program (HR = 1.77; 99% CI = 1.45, 2.18), and missing data regarding union membership (HR = 1.27, 99%CI = 1.01, 1.60) was associated with shorter claim duration. Contrary to our predictions, neither WSIB-reimbursement for physiotherapy nor chiropractic care were associated with claim duration (Table 2).

Predicting Workers' Compensation claim duration

Our ongoing systematic review <sup>14</sup> identified 3 observational studies that explored early use of opioids for Workers' Compensation claims due to uncomplicated LBP, 2 of which adjusted for injury severity, and all of which reported a significant association with prolonged claim duration (Table S2 in File S1).<sup>20-22</sup> When our results were pooled with the 2 studies that reported measures of association in relative units,<sup>21,22</sup> resulting in a total of 51,069 participants, the association between early opioid use and prolonged claim duration was consistent with our findings (adjusted HR = 0.58, 95%CI (0.48 to 0.70), heterogeneity test p=0.02, I<sup>2</sup>=76%) (Figure 2). The pooled effect was associated with substantial heterogeneity; however, statistical tests of heterogeneity can be misleading when sample sizes are very large and confidence intervals are therefore very narrow.<sup>23</sup> These results provide an excellent example of the phenomenon. The three studies all show a large effect, and the point estimates are very consistent one with the other (0.52 to 0.69) which increases confidence in our findings.

### **Discussion**

Our analysis of the Ontario WSIB's administrative data revealed that older claimants who were disabled due to uncomplicated LBP and who are reimbursed for opioid prescription in the first 4 weeks of their claim are more likely to experience prolonged claim duration. Higher pre-injury income was also associated with prolonged claim duration, but only among persistent claims. Injured workers employed by organizations with a RTW program who are missing information on union affiliation are likely to resolve their claim faster. The WSIB reimburses physiotherapy for 55% of uncomplicated LBP claims, and chiropractic care for 27% of LBP claims, and neither was associated with claim duration.

The strengths of our study included the *a priori* creation of our regression model and the anticipated direction of included independent variables. We were unable to adjust for injury severity, but attempted to enroll patients with similar injuries by restricting our cohort to acute, uncomplicated LBP. Other strengths include limited missing data, correction of identifiable data errors and inconsistencies, and validation checks to ensure the accuracy of the data used to inform our regression model. We assessed the proportional hazards assumption for all independent variables in our regression model.

Our study has several limitations. First, it was a retrospective cohort study in which the reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are uncertain. Thus, despite our adjustments for potential confounders, it remains possible that LBP claimants chosen to receive these healthcare interventions were prognostically different than those claimants who did not. Second, the WSIB database only records physiotherapy or chiropractic treatments that are reimbursed by the WSIB, and it is possible that some patients paid out-of-

Predicting Workers' Compensation claim duration pocket to receive these services. Third, a number of variables that may be important to consider were unavailable (e.g., patient expectations regarding recovery <sup>24</sup>), and some variables were not optimally collected. For example, chiropractic and physiotherapy are professions and not modalities, and there were no details of treatment provided. Fourth, our study focused on workers with LBP receiving disability benefits from the Ontario WSIB, we cannot say whether our findings are generalizable to other Workers' Compensation Boards. Finally, our primary outcome, claim closure, is a surrogate for patient-important outcomes such as functional restoration or return to work.

Our finding that older age is associated with prolonged claim duration is consistent with the literature. We predicted that injured workers employed by companies that had formal return to work programs would resolve their claim faster, and this was supported by our findings. It is unclear why missing data for union membership was also associated with shorter claim duration. Our finding that higher pre-disability income is associated with prolonged claim duration, but only among persistent claims, suggests that injured workers with higher salaries that do not resolve their claim in the initial 6-months may find it more difficult to identify suitable employment that provides similar earnings.

Although there are no randomized controlled trials exploring the effect of physiotherapy, chiropractic care, or early opioid use for workers with uncomplicated LBP receiving compensation benefits, there are observational studies that are relevant to our findings. Wasiak and colleagues found that workers in Florida, USA, with low back injuries who were reimbursed for limited chiropractic care (<30 days) experienced an 8.6% shorter duration of work disability versus Workers' Compensation claimants who were reimbursed for prolonged

Predicting Workers' Compensation claim duration

chiropractic care (>30 days). <sup>26</sup> The authors did exclude severe injuries from their population, but were unable to adjust for injury severity within their sample. Further, chiropractic care was collected after baseline and not treated as time-dependent, and so it is not surprising that claims with longer duration also received more chiropractic care.

Lemstra and Olszynski explored the effect of standard care (which included long waiting lists for physiotherapy) to provision of rapid rehabilitation services on Workers' Compensation claim duration from a company in Saskatchewan, Canada. After adjusting for a number of factors, including age and injury severity, longer claim duration was associated with both chiropractic care (adjusted HR = 2.88, 95% CI = 1.45, 5.73) and physical therapist involvement (adjusted HR = 19.88, 95% CI = 7.95, 39.77). The authors collected healthcare provider utilization data after baseline and did not treat these variables as time-dependent, and an alternative explanation is that claims with longer duration are more likely to involve either chiropractors or physical therapists.

Turner and colleagues followed 1,885 workers from Washington for 1 year after they had been awarded Workers' Compensation benefits for acute LBP. In a comprehensive regression model adjusted for multiple sociodemographic, employment, clinical, healthcare, and administrative factors – including injury severity – they found that workers who attended a chiropractor first, versus a primary care provider, were significantly less likely to remain on disability benefits at 1 year (adjusted OR = 0.41, 95% CI = 0.24, 0.70).<sup>28</sup>

Canada is currently the largest per-capita consumer of opioids in the world; <sup>29</sup> however, prescribing patterns between primary care providers in Ontario show considerable variation <sup>30</sup>. Workers' Compensation data has shown an almost 10 fold difference (5.7% to 52.9%) in the

Predicting Workers' Compensation claim duration

early prescription of opioid medications between various US states, suggesting that local prescribing patterns have significant influence on the use of these analgesics. <sup>31</sup> Findings from a study of Workers' Compensation nonspecific LBP claims revealed that, compared with a no opioid reference group, odds of chronic work loss were six times greater for claimants that used strong opioids and 11-14 times greater for claimants with opioid prescriptions which exceeded 90 days. <sup>32</sup> We found that reimbursement for early opioid use by the Ontario WSIB prolongs claim duration for uncomplicated back pain, and pooling of our data with similar studies <sup>21,22</sup> shows a consistent effect, which increases confidence in our findings.

To manage their growing unfunded liability, the Ontario WSIB has focused on increasing their claim denial rate, decreasing benefits to injured workers, reducing WSIB staff, and raising employer premiums. 33,34 Most employers are obligated to pay WSIB premiums as they are legally bound to provide injury benefits to their employees and the Ontario WSIB is protected by laws prohibiting competition in the marketplace. Another strategy is to optimize clinical management of injured workers. Our findings suggest that therapies for acute, uncomplicated LBP commonly reimbursed by the Ontario WSIB may be ineffective or even prolong claim duration. High quality, randomized controlled trials are urgently needed to confirm or refute our findings.

Predicting Workers' Compensation claim duration

#### **Author affiliations**

- <sup>1</sup> The Michael G. DeGroote Institute for Pain Research and Care, McMaster University, Hamilton, Canada
- <sup>2</sup> Department of Anesthesia, McMaster University, Hamilton, Canada
- <sup>3</sup> Department of Clinical Epidemiology and Biostatistics, McMaster University, Hamilton, Canada
- <sup>4</sup> Stanford Prevention Research Center, Department of Medicine, Stanford University, Stanford, California, USA
- <sup>5</sup> Department of Anaesthesia & Pain Medicine, the Hospital for Sick Children, Toronto, Canada

#### **Acknowledgements**

The authors thank Dr. Gordon Guyatt for helpful discussion.

#### **Contributors**

JWB, SE, D-HA and SDW designed the study. DH-A and LW conducted data analysis and SDW provided statistical advice. JWB, SE, D-HA and SDW were involved in interpreting the data. JWB drafted the manuscript and wrote the final version. All authors critically revised the manuscript, provided comment and approved the final version for publication.

#### **Funding**

This study was funded by research grants from the Ontario Workers Safety and Insurance Board Research Advisory Council and the Ontario Chiropractic Association. Shanil Ebrahim is supported by a MITACS Elevate Postdoctoral Fellowship Award. The funding sources had no role in design

or conduct of the study; the collection, management, analysis, or interpretation of the data; or the preparation, review, or approval of the manuscript.

Predicting Workers' Compensation claim duration

#### **Competing interests**

Jason Busse acts as a consultant to Prisma Health Canada, a private incorporated company funded by employers and insurers that consults on and manages long-term disability claims.

#### **Ethics approval**

The study was approved by the Health Sciences Research Ethics Board of the University of Toronto.

#### Provenance and peer review

Not commissioned; externally peer reviewed.

#### Data sharing statement

No additional data are available.

#### **Open Access**

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work noncommercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://

creativecommons.org/licenses/by-nc/4.0/



Predicting Workers' Compensation claim duration

#### References

- 1. Thiese MS, Hegmann KT, Wood EM, et al.; BackWords Study Team. Low-back pain ratings for lifetime, 1-month period, and point prevalences in a large occupational population. *Hum Factors* 2014; 56: 86-97.
- 2. Deyo RA, Phillips WR. Low Back Pain: A Primary Care Challenge. *Spine* 1996; 21: 2826-2832.
- 3. Hoy D, March L, Brooks P, et al. The global burden of low back pain: estimates from the Global Burden of Disease 2010 study. *Ann Rheum Dis* 2014; 73: 968-974.
- 4. Brown A, Angus D, Chen S, et al. Costs and outcomes of chiropractic treatment for low back pain [Technology report no 56]. Ottawa: Canadian Coordinating Office for Health Technology Assessment. 2005.
- By the Numbers: 2012 WSIB Statistical Report. Schedule 1
   [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S1.pdf] (accessed August 3, 2014).
- By the Numbers: 2012 WSIB Statistical Report. Schedule 2
   [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S2.pdf] (accessed August 3, 2014).
- Workplace Safety and Insurance Board 2013: First Quarter Sufficiency Report to Stakeholders

Predicting Workers' Compensation claim duration [http://www.wsib.on.ca/files/Content/SufficiencySufficiencyReport2013/2013Q1WSIBSufficiencyReport.pdf] (accessed August 3, 2014).

- Ontario WSIB funding fiasco. National Post. November 18, 2008
   [http://www.nationalpost.com/opinion/story.html?id=eb83e348-1660-4771-9804-e47a502aa756] (accessed August 3, 2014).
- 9. Harris I, Mulford J, Solomon M, et al. Association between compensation status and outcome after surgery: a meta-analysis. *JAMA* 2005; 293: 1644–1652.
- von Elm E, Altman DG, Egger M, et al. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) Statement: guidelines for reporting observational studies.
   Ann Intern Med 2007; 147: 573-577.
- Collins GS, Reitsma JB, Altman DG, et al. Transparent Reporting of a multivariable prediction model for Individual Prognosis Or Diagnosis (TRIPOD): The TRIPOD Statement. *Ann Intern Med* 2015; 162: 55-63.
- 12. da C Menezes Costa L, Maher CG, Hancock MJ, et al. The prognosis of acute and persistent low-back pain: a meta-analysis. *CMAJ* 2012; 184: E613-624.

Predicting Workers' Compensation claim duration

- 13. Frank JW, Brooker AS, DeMaio SE, et al. Disability resulting from occupational low back pain. Part II: What do we know about secondary prevention? A review of the scientific evidence on prevention after disability begins. *Spine (Phila Pa 1976)* 1996; 21: 2918-2929.
- 14. [blinded]
- 15. Harrell FE. Multivariate modeling strategies. In: Harrell FE, ed. Regression Modeling Strategies With Applications to Linear Models, Logistic Regression and Survival Analysis.

  New York, NY: Springer; 2001: 53–85.
- 16. Zhang J, Yu KF. What's the Relative Risk? A Method of Correcting the Odds Ratio in Cohort Studies of Common Outcomes. *JAMA* 1998; 280: 1690-1691.
- 17. Montori, V, Ioannidis J, Cook DJ, et al. Fixed-effects and random-effects models. In G. Guyatt, Rennie D, Meade MO, Cook DJ (Ed.), Users' guides to the medical literature: A manual for evidence-based clinical practice (2nd ed., pp. 555-562). USA: McGraw Hill Companies. 2008.
- 18. Higgins JPT, Thompson SG, Deeks JJ, et al. Measuring inconsistency in meta-analyses. *BMJ* 2003; 327: 557-560.

Predicting Workers' Compensation claim duration

19. The Cochrane Collaboration. 9.5.2 Identifying and measuring heterogeneity. In J. P. T. Higgins, Thompson SG (Ed.), Cochrane Handbook for Systematic Reviews of Interventions (Version 5.0.2). Oxford, UK: Cochrane Collaboration. 2009.

- 20. 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: 2127-2132.
- 21. Franklin GM, Stover BD, Turner JA, et al.; Disability Risk Identification Study Cohort. 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.
- 22. Gross DP, Stephens B, Bhambhani Y, et al. Opioid prescriptions in Canadian workers' compensation claimants: prescription trends and associations between early prescription and future recovery. *Spine (Phila Pa 1976)* 2009; 34: 525-531.
- 23. Rücker G, Schwarzer G, Carpenter JR, et al. Undue reliance on I<sup>2</sup> in assessing heterogeneity may mislead. *BMC Med Res Methodol* 2008; 8: 79.
- 24. Cole DC, Mondloch MV, Hogg-Johnson S; Early Claimant Cohort Prognostic Modelling Group. Listening to injured workers: how recovery expectations predict outcomes--a prospective study. *CMAJ* 2002; 166: 749-754.

Predicting Workers' Compensation claim duration

- 25. Hadler NM. The bane of the aging worker. Spine (Phila Pa 1976) 2001; 26: 1309-1312.
- 26. Wasiak R, Kim J, Pransky GS. The association between timing and duration of chiropractic care in work-related low back pain and work-disability outcomes. *J Occup Environ Med* 2007; 49: 1124-1134.
- 27. Lemstra M, Olszynski WP. The effectiveness of standard care, early intervention, and occupational management in worker's compensation claims. *Spine (Phila Pa 1976)* 2003; 28: 299-304.
- 28. Turner JA, Franklin G, Fulton-Kehoe D, et al. ISSLS prize winner: early predictors of chronic work disability: a prospective, population-based study of workers with back injuries. *Spine* (*Phila Pa 1976*) 2008; 33: 2809-2818
- DCAM Consortium Drug Consumption Motion Chart [http://ppsgproduction.heroku.com/chart] (accessed August 3, 2014).
- 30. Dhalla IA, Mamdani MM, Gomes T, et al. Clustering of opioid prescribing and opioid-related mortality among family physicians in Ontario. *Can Fam Physician* 2011; 57: e92–96.

Predicting Workers' Compensation claim duration

31. Webster BS, Cifuentes M, Verma S, et al. Geographic variation in opioid prescribing for acute, work-related, low back pain and associated factors: a multilevel analysis. *Am J Ind Med* 2009; 52: 162-171.

- 32. Volinn E, Fargo JD, Fine PG. Opioid therapy for nonspecific low back pain and the outcome of chronic work loss. *Pain* 2009; 142: 194-201.
- 33. WSIB President gets \$80K bonus on the backs of injured workers. UFCW Canada [http://www.ufcw.ca/index.php?option=com\_content&view=article&id=2760:wsib-president-gets-80k-bonus-on-the-backs-of-injured-workers&Itemid=6&Iang=en] (accessed August 3, 2014).
- 34. WSIB raises its rates to tackle \$12 billion in unfunded liabilities. Machinery and Equipment MRO. November 1, 2010 [http://www.mromagazine.com/news/wsib-raises-its-rates-to-tackle-12-billion-in-unfunded-liabilities/1000391835/?&er=NA] (accessed August 3, 2014).

Predicting Workers' Compensation claim duration

### **Figure Legends**

Figure 1: Kaplan-Meier curve for time to claim closure

Figure 2: The association between early opioid use/prescription and claim duration



## **Supporting Information Legends**

**Table S1: Description of model variables** 

Table S2: Observational studies exploring the association between early opioid use and



Table 1: Baseline Characteristics of WSIB Low Back Pain Claims (n=1,442)

Age, mean (SD)	41.3 (10.5)
Gender, n (%)	
Female	552 (38.3)
Male	890 (61.7)
First language, n (%)	
English	1372 (95.1)
Other	70 (4.9)
Pre-disability income (dollars/week)	
mean (SD)	731.4 (332.5)
Opioid prescription reimbursed by WSIB in the first 4-weeks of	
claim, n (%)	
Yes	136 (9.4)
No	1306 (90.6)
Prior WSIB claim, n (%)	
Yes	1091 (75.7)
No	351 (24.3)
Union membership, n (%)	
Yes	610 (42.3)
No	656 (45.5)
Missing data	176 (12.2)
Employer RTW program, n (%)	
Yes	1042 (72.3)
No	278 (19.3)
Missing data	122 (8.5)
Employer doubts work-relatedness of injury, n (%)	
Yes	195 (13.5)
No	1051 (72.9)
Missing data	196 (13.6)
Chiropractic care reimbursed by WSIB during claim, n (%)	391 (27.1)
Physiotherapy reimbursed by WSIB during claim, n (%)	786 (54.5)

	Adjusted Hazard Batic (00% CI)	and Patio (00% CI)	
A ===	Adjusted Hazard Ratio (99% CI)	p-value	
Age	1.00	<0.001	
15 to <25	1.00		
25 to <35	0.79 (0.58, 1.08)		
35 to <45	0.70 (0.52, 0.95)		
45 to <55	0.67 (0.49, 0.91)		
55 to 65	0.52 (0.36, 0.74)		
Gender		0.45	
Male	1.00		
Female	0.96 (0.82, 1.12)		
First language		0.33	
English	1.00		
Other	0.88 (0.63, 1.23)		
Pre-disability gross income (dollars/week)			
At 30 days:			
≤480	1.00		
481-694	1.08 (0.85, 1.37)	0.396	
695-920	1.05 (0.81, 1.36)	0.614	
>920	1.04 (0.80, 1.35)	0.689	
At 180 days:			
≤480	1.00		
481-694	0.85 (0.65, 1.12)	0.130	
695-920	0.71 (0.52, 0.96)	0.003	
>920	0.61 (0.45, 0.84)	<0.001	
At 365 days:			
≤480	1.00		
481-694	0.63 (0.35, 1.16)	0.051	
695-920	0.44 (0.23, 0.84)	0.001	
>920	0.32 (0.16, 0.63)	<0.001	
Opioid prescription reimbursed by the Ontario WSIB in			
the first 4 weeks of claim		<0.001	
No	1.00		
Yes	0.69 (0.53, 0.89)		
Prior WSIB claim		0.66	
No	1.00		
Yes	1.03 (0.87, 1.22)		
Union membership		0.01	
No	1.00		
Yes	1.14 (0.96, 1.36)		
Missing data	1.27 (1.01, 1.60)		
Employer with a RTW program	( - , ,	<0.001	
No	1.00		
Yes	1.77 (1.45, 2.18)		
Missing data	1.18 (0.86, 1.60)		
Employer doubts work-relatedness of injury		0.12	
No	1.00	0.12	
Yes	0.87 (0.70, 1.08)		
Missing data Chicagrattic care reimbursed by WSIR	1.08 (0.87, 1.33)	0.27	
Chiropractic care reimbursed by WSIB	1.08 (0.91, 1.29)	0.27	
Physiotherapy reimbursed by WSIB  * HR>1 indicates faster claim closure; RTW = return to wo	0.99 (0.85, 1.16)	0.91	

<sup>\*</sup> HR>1 indicates faster claim closure; RTW = return to work

Figure 1: Kaplan-Meier curve for time to claim closure

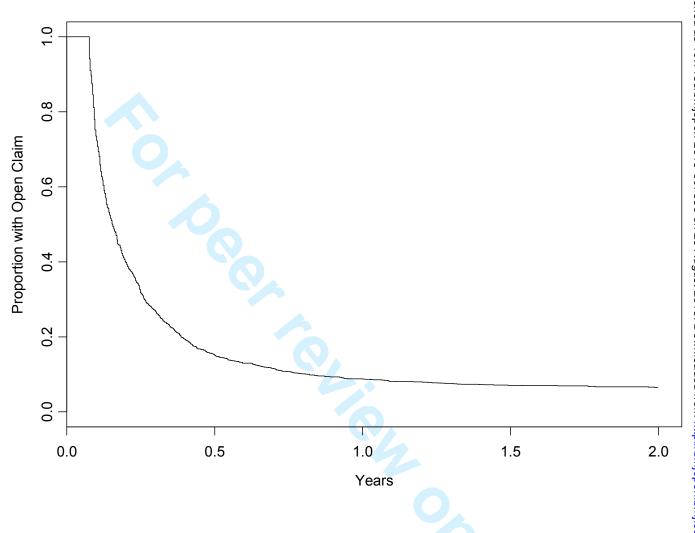
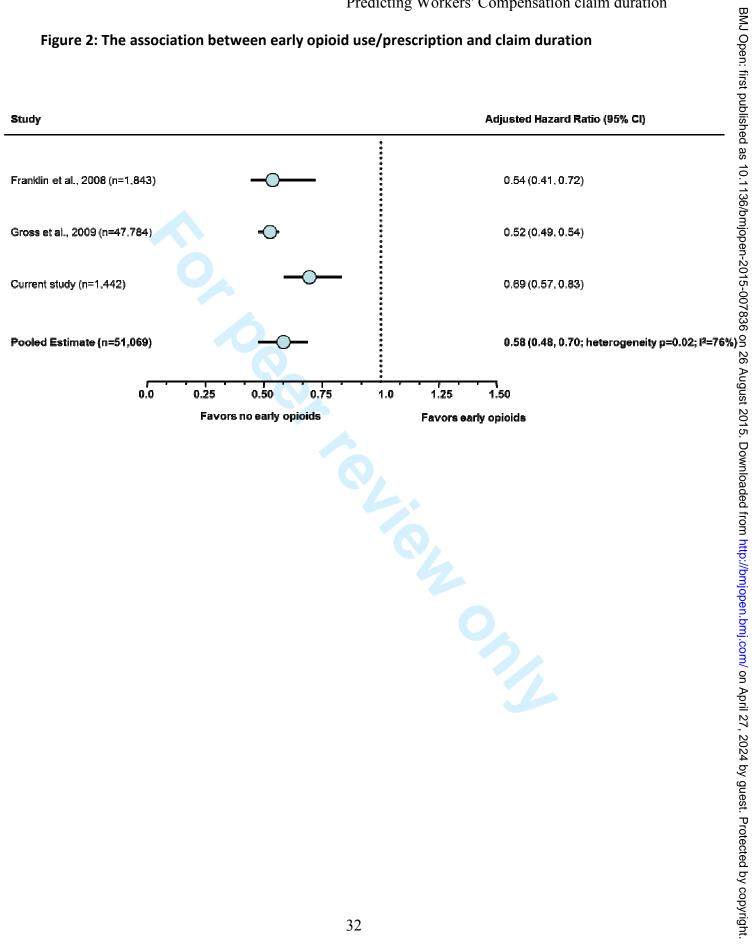


Figure 2: The association between early opioid use/prescription and claim duration



**Table S1: Description of model variables** 

Variable	Anticipated Direction of		
	Effect		
Age	older age: -		
Gender	female: -		
First language	non-English: -		
Pre-disability income	higher income: -		
Reimbursement for opioid	opioid reimbursement: -		
prescription			
Prior claim(s)	prior claim: -		
Union membership	Union member: +		
Employer RTW-program	RTW program: +		
Work-relatedness	work-related: +		
Chiropractic care*	DC care: +		
Physiotherapy*	PT care: +		

- +: associated with faster claim closure
- re -: associated with slower claim closure
- \*: time dependant covariate

Table S2: Observational studies exploring the association between early opioid use and Workers' Compensation claim duration

Study	Population	Opioid variable tested	Adjustments	Dependant variable	Results*
Webster et al., 2007	8,443 American Workers' Compensation claimants with new-onset, disabling LBP	Receipt of opioids within the first 15 days of claim	Injury severity, age, gender, length of job tenure	Change in mean disability duration	1-140mg MEA 5.4 days, 95%Cl = -14.6 to 25.0 141-225mg MEA 21.9 days, 95%Cl = 3.2 to 40.6 226-450mg MEA 43.8 days, 95%Cl = 23.7 to 63.9 >450mg MEA 69.1 days, 95%Cl = 49.3 to 89.0
Franklin et al., 2008	1,843 Washington, US, Workers' Compensation claimants with new-onset, disabling LBP	Reimbursement for opioids within 6 weeks of 1 <sup>st</sup> medical visit for LBP	Age, gender, race, education, injury severity, pain intensity, Roland disability questionnaire	Receipt of wage replacement benefits at 1-year	1-150mg MED  OR = 1.9, 95%CI = 1.2 to 3.1  151-300mg MED  OR = 2.0, 95%CI = 1.2 to 3.3  301-650mg MED  OR = 1.6, 95%CI = 0.9 to 2.6  >650mg MED  OR = 1.9, 95%CI = 1.2 to 2.9
Gross et al., 2009	47,784 Alberta, Canada, Workers' Compensation claimants with new-onset, disabling LBP	Reimbursement for opioids within the first 2 weeks of claim	Age, gender, annual salary, year of claim, number of previous claims	Receipt of wage replacement benefits at 1-year	No early opioids HR = 1.94, 95%CI = 1.86 to 2.02

<sup>\*</sup> The reference group is no early receipt of opioids for Franklin et al., and Webster et al. 95% CI = 95% confidence interval LBP = low back pain

MEA = morphine equivalent amount MED = morphine equivalent dose

OR = odds ratio

HR = hazard ratio



### STROBE Statement—Checklist of items that should be included in reports of *cohort studies*

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the
		abstract (in title on title page)
		(b) Provide in the abstract an informative and balanced summary of what
		was done and what was found (in Abstract)
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being
C		reported (page 1 & 2 in Manuscript)
Objectives	3	State specific objectives, including any prespecified hypotheses ( <b>Appendix</b>
		<b>A</b> )
Methods		
Study design	4	Present key elements of study design early in the paper (In Abstract, and
		page 3-7 in Manuscript)
Setting	5	Describe the setting, locations, and relevant dates, including periods of
		recruitment, exposure, follow-up, and data collection (page 3 in
		Manuscript)
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of
		participants. Describe methods of follow-up (pages 3 & 4 in Manuscript)
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed (not applicable)
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders,
		and effect modifiers. Give diagnostic criteria, if applicable (Appendix A
		and pages 3 & 4 in Manuscript)
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of
		assessment (measurement). Describe comparability of assessment methods
		if there is more than one group (pages 3-5 in Manuscript)
Bias	9	Describe any efforts to address potential sources of bias (pages 4-7 in
		Manuscript)
Study size	10	Explain how the study size was arrived at (page 3 & 4 in Manuscript)
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If
		applicable, describe which groupings were chosen and why (pages 5 & 6 in
		Manuscript)
Statistical methods	12	(a) Describe all statistical methods, including those used to control for
		confounding (pages 3-7 in Manuscript)
		(b) Describe any methods used to examine subgroups and interactions (page
		6 in Manuscript)
		(c) Explain how missing data were addressed (page 5 in Manuscript)
		(d) If applicable, explain how loss to follow-up was addressed ( <b>not</b>
		applicable)
		(e) Describe any sensitivity analyses ( <b>not applicable</b> )
Results	102	
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers
		potentially eligible, examined for eligibility, confirmed eligible, included in
		the study, completing follow-up, and analysed ( <b>Table 1 and page 8 in</b>
		Manuscript)  (b) Give reasons for non-participation at each stage (not applicable)
		(b) Give reasons for non-participation at each stage ( <b>not applicable</b> )

Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical,
		social) and information on exposures and potential confounders ( <b>Table 1</b> )
		(b) Indicate number of participants with missing data for each variable of
		interest (Table 1)
		(c) Summarise follow-up time (eg, average and total amount) (Figure 1)
Outcome data	15*	Report numbers of outcome events or summary measures over time (page 8
		in Manuscript)
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted
		estimates and their precision (eg, 95% confidence interval). Make clear
		which confounders were adjusted for and why they were included (Table 2)
		(b) Report category boundaries when continuous variables were categorized
		(Table 2)
		(c) If relevant, consider translating estimates of relative risk into absolute
		risk for a meaningful time period (not relevant)
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and
,		sensitivity analyses (page 8 in Manuscript)
Discussion		
Key results	18	Summarise key results with reference to study objectives (page 10 in
		Manuscript)
Limitations	19	Discuss limitations of the study, taking into account sources of potential
		bias or imprecision. Discuss both direction and magnitude of any potential
		bias (pages 10 & 11 in Manuscript)
Interpretation	20	Give a cautious overall interpretation of results considering objectives,
		limitations, multiplicity of analyses, results from similar studies, and other
		relevant evidence (page 13 in Manuscript)
Generalisability	21	Discuss the generalisability (external validity) of the study results (page 11
		in Manuscript)
Other information		
Funding	22	Give the source of funding and the role of the funders for the present study
		and, if applicable, for the original study on which the present article is based
		(page 14 in Manuscript)
		1/

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at http://www.strobe-statement.org.

# **BMJ Open**

The association of worker characteristics and early reimbursement for physical therapy, chiropractic and opioid prescriptions with Workers' Compensation claim duration, for cases of acute low back pain: An observational cohort study

Journal:	BMJ Open
Manuscript ID:	bmjopen-2015-007836.R1
Article Type:	Research
Date Submitted by the Author:	21-May-2015
Complete List of Authors:	Busse, Jason; McMaster University, Anesthesia; McMaster University, Clinical Epidemiology & Biostatistics Ebrahim, Shanil; Mcmaster University, Clinical Epidemiology and Biostatistics Heels-Ansdell, Diane; Mcmaster University, Clinical Epidemiology and Biostatistics Wang, Li; McMaster University, Anesthesia Couban, Rachel; McMaster University, Anesthesia Walter, Stephen; Mcmaster University, Clinical Epidemiology and Biostatistics
<b>Primary Subject Heading</b> :	Rehabilitation medicine
Secondary Subject Heading:	Complementary medicine, Evidence based practice
Keywords:	COMPLEMENTARY MEDICINE, Pain management < ANAESTHETICS, OCCUPATIONAL & INDUSTRIAL MEDICINE

SCHOLARONE™ Manuscripts

The association of worker characteristics and early reimbursement for physical therapy, chiropractic and opioid prescriptions with Workers' Compensation claim duration, for cases of acute low back pain: An observational cohort study

Jason W. Busse,<sup>1,2,3\*</sup> Shanil Ebrahim,<sup>2,3,4,5</sup> Diane Heels-Ansdell,<sup>3</sup> Li Wang,<sup>2</sup> Rachel Couban,<sup>2</sup> Stephen D. Walter<sup>3</sup>

\*Corresponding Author: email: bussejw@mcmaster.ca

# **Abstract**

**Objective:** To assess the association between early reimbursement for physiotherapy, chiropractic, and opioid prescription for acute low back pain (LBP) with disability claim duration.

**Design:** Observational cohort study.

**Setting and participants:** From a random sample of 6,665 claims for acute LBP approved by the Ontario Workplace Safety and Insurance Board (WSIB) in 2005, we analyzed 1,442 that remained on benefits at 4-weeks.

**Primary outcome measure:** WSIB claim duration.

Results: Our time-to-event analysis was adjusted for demographic, workplace, and treatment factors, but not injury severity. Older age (e.g. hazard ratio [HR] for age ≥55 versus <25 = 0.52; 99% confidence interval [CI] = 0.36, 0.74) and WSIB-reimbursement for opioid prescription in the first 4-weeks of a claim (HR = 0.68; 99% CI = 0.53, 0.88) were associated with longer claim duration. Our systematic review (n=51,069 workers) confirmed a strong association between early opioid use and prolonged claim duration (HR = 0.57, 95%CI = 0.48 to 0.69). Among persistent claims, higher pre-disability income was associated with longer claim duration (e.g. HR for active claims at 1-year with a pre-disability income >\$902/week versus ≤\$480 = 0.34; 99% CI = 0.17, 0.68). Missing data for union membership (HR = 1.27; 99% CI = 1.01, 1.59), and working for an employer with a return-to-work program was associated with shorter claim duration (HR = 1.78; 99% CI = 1.45, 2.18). Neither reimbursement for physiotherapy (HR = 1.01; 99% CI = 0.86, 1.19) or chiropractic care (HR for active claims at 60 days = 1.15; 99% CI = 0.94, 1.41) within the first 4-weeks were associated with claim duration.

Predicting Workers' Compensation claim duration

**Conclusions:** Our analysis found that early WSIB reimbursement for physiotherapy or chiropractic care is not associated with claim duration, and early reimbursement for opioids predicts prolonged claim duration. Well-designed randomized controlled trials are needed to verify our findings.

**Key Words:** Low back pain; disability; compensation; Workers Compensation Board; physical therapy; chiropractic; opioid; survival analysis

Predicting Workers' Compensation claim duration

#### **Strengths and Limitations**

- A priori creation of our regression model and the anticipated direction of included independent variables, and assessment of the proportional hazards assumption for all independent variables, provide greater confidence in our findings.
- The reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are
  uncertain, and despite our adjustments for potential confounders (but not injury
  severity) it remains possible that LBP claimants who chose to receive these healthcare
  interventions were prognostically different than those claimants who did not.
- A number of variables that may be important to consider were unavailable (e.g., patient
  expectations regarding recovery), and chiropractic and physiotherapy are professions,
  not modalities, and details of the treatment provided could not be obtained for our
  analysis.
- Our primary outcome, time to claim closure, is a surrogate for patient-important outcomes such as functional restoration or return to work.

Predicting Workers' Compensation claim duration

# Introduction

lived with disability.3

Back pain is a common problem among working adults in North America, with a lifetime prevalence of 63% and a point prevalence of 21%. After the common cold, low back pain (LBP) is the most frequent cause of lost time from work.<sup>2</sup> Globally, LBP is the primary cause of years

In Canada, annual medical expenditures for LBP are estimated to be between \$6 and \$12 billion, with additional costs associated with loss in worker productivity from time off work and associated disability payments. 4 Canadian workers who are disabled secondarily to a workrelated LBP injury are typically eligible for wage replacement benefits through their provincial Workers' Compensation Board.

In 2013, the Ontario Workplace Safety and Insurance Board (WSIB) approved approximately 232,000 claims that were associated with \$2,761 million in payments, and 18% of all allowed lost-time claims were for low back injuries. 5, 6 The WSIB's liability for disability claims greatly exceeds their assets, and as of March 31, 2013 the WSIB's unfunded liabilities were \$12.4 billion<sup>7</sup> – more than double their unfunded liability of \$5.9 billion in 2006.<sup>8</sup> Unfunded liability is the amount by which future payment obligations exceed the present value of funds available to pay them. To reduce their unfunded liability, the WSIB has become more aggressive about denying claims, decreasing disability benefits, and increasing employee premiums; 9, 10 however, these measures do not address optimal management of disability claims.

Interventions that are commonly reimbursed by WSIB for LBP claims include physiotherapy, chiropractic care and opioids; however, there is limited evidence about their effectiveness. Our systematic review of the Cochrane Back Review Group trial registry found no randomized controlled trials of these interventions focused on workers disabled by LBP and receiving benefits for lost-time claims (Tables S1 & S2, and Figure S1, in File S1). We also found that many trials that enroll LBP patients use receipt of disability benefits as an exclusion criterion, likely because of concerns that secondary gain (e.g., receipt of financial compensation conditional on disability) will reduce the impact of study interventions. Henshke and colleagues followed a cohort of 973 consecutive primary care patients with non-specific, acute LBP recruited from the clinics of 170 general practitioners, physiotherapists, and chiropractors for 1year. They found that, in an analysis adjusted for age, gender, injury severity and psychological factors, receipt of disability benefits was strongly associated with delayed recovery (hazard ratio [HR]=0.59; 95%CI = 0.47 to 0.74). 11 As compensated acute LBP has a worse prognosis than uncompensated acute LBP, the results of trials that do not enroll patients receiving disability benefits cannot be confidently generalized to patients who are receiving compensation.

Using administrative data from the Ontario WSIB, we evaluated the association between receiving early reimbursement for physiotherapy, chiropractic care, or prescription for opioids for uncomplicated LBP and disability claim duration. On the basis of prior observational studies, <sup>12-16</sup> we hypothesized that early reimbursement for opioids would be associated with delayed recovery, and early reimbursement for physiotherapy or chiropractic care would be associated with faster recovery. We reported our findings in concordance with the STROBE <sup>17</sup> and TRIPOD <sup>18</sup> statements.

Predicting Workers' Compensation claim duration

# **Methods**

### Patient characteristics and eligibility criteria

Using WSIB administrative data, we identified an inception cohort of workers with uncomplicated low back injuries (i.e., strain or sprain) who were fully disabled from working and receiving wage replacement benefits from the Ontario WSIB. We excluded workers if they were approved for no-lost-time claims. The prognosis for chronic LBP (duration >12 weeks) is different than for acute LBP, <sup>19</sup> and we excluded all claims in which the number of days between accident date and registration date of the claim was greater than 30 days. Most LBP claims resolve within the 1<sup>st</sup> month, <sup>20</sup> and Workers' Compensation Boards are primarily interested in factors that predict claim resolution among claimants that remain disabled after this time. We therefore excluded claims that ended before 4 weeks.

Between January 1 and June 30, 2005, the Ontario WSIB approved 18,974 lost-time claims for an uncomplicated, acute low back injury. Using the WSIB's administrative database, we acquired a random sample of 6,665 injured workers from this population; 1,442 unique workers remained on full benefits at four weeks and provided data for our analysis. If a worker had more than 1 claim for acute LBP, their first claim was used. The WSIB database recorded benefit status for two years after the first day of injury. Patient information was anonymized and de-identified prior to analysis. The Health Sciences Research Ethics Board of the University of Toronto approved our study protocol.

#### **Administrative Variables**

Our primary outcome was time to claim closure, defined as the duration in days from disability claim approval until the claim was closed. The WSIB database also contained demographic, administrative, and clinical information, which we acquired from forms completed by the worker, their employer, and their primary health-care provider. The employer form (Employer's Report of Injury/Disease Form; Form 7), which is used to indicate whether there is doubt regarding the work-relatedness of an employee's back injury, is mandatory and must be submitted within three days of a work-related injury. The form asks employer's "Do you have any reason to doubt the injury/disease is work-related?" and they can indicate either "no" or "yes".

The worker may elect to fill out a form (Worker's Report of Injury/Disease; Form 6) if they have expenses related to their injury. The healthcare provider can elect to complete a form (Health Professional's Report; Form 8) to support their patient's claim that their injury is work-related, which is a pre-requisite for wage replacement benefits through the WSIB. Healthcare providers are asked to complete and submit a Functional Abilities Form for Planning Early and Safe Return to Work for each claim, and the WSIB provides compensation as incentive.

In order to increase confidence in our findings, we defined our regression model before conducting any analyses. Guided by the results from our ongoing systematic review of observational studies evaluating predictors of recovery in patients receiving disability benefits, <sup>21</sup> feedback from administrators at WSIB, and content experts within our research team, we selected, *a priori*, 11 variables from the WSIB database that we judged may be associated with claim closure; we also specified the direction of anticipated effects on claim

Predicting Workers' Compensation claim duration

duration (Table 1): age, gender, native language, pre-disability income, prior disability claim, union membership, employer with a return to work (RTW) program, employer's doubt that the injury was work-related, and early (1<sup>st</sup> month) receipt of reimbursement for treatment with opioids, physiotherapy, or chiropractic care (Table 1). The Ontario WSIB does not capture any measure of LBP injury severity, and as such we were unable to adjust for this variable. Injured workers may attend a healthcare provider for assessment purposes. We required ≥3 reimbursed visits for physiotherapy or chiropractic, within the first 28-days of claim, in order to qualify as reimbursement for treatment.

We hypothesized that workers represented by a union would resolve their claim faster, as they likely had more support for re-engagement with competitive employment (e.g. graduated work hours) versus workers who were not supported by a union. We also hypothesized that claims due to injuries that employers reported were work-related would resolve faster than injuries in which the employer doubted that the employee was injured at work, as we felt this may be a surrogate for the influence of non-medical factors (e.g. secondary gain). Based on the findings of a recent systematic review that found RTW coordination was associated with faster RTW for disabled employees, <sup>22</sup> we hypothesized that claimants employed by companies with formal RTW programs would resolve their claim faster.

#### **Data Extraction**

The Ontario WSIB's database consists of scanned paper documents, and data must therefore be extracted manually for analysis. Two reviewers extracted data, independently and in duplicate, from the first 100 eligible claims into an Access database (Microsoft Access, Filemaker). In order

to minimize data entry mistakes, we developed data entry forms that included range checks and missing value alerts. The PROC COMPARE procedure in SAS version 9.2 (SAS Institute Inc., Cary, NC) revealed 98% agreement for the initial 100 claims; therefore, to increase feasibility, only a single abstractor completed data entry for the remaining claims used in our analyses.

We screened all data to identify outliers, inconsistencies and missing data by calculating summary statistics, and explored distributions graphically. We worked with WSIB representatives to correct identified outliers and inconsistencies. If inconsistencies could not be corrected, we treated them as missing data. Some WSIB forms are voluntary, and so we included "missing data" as a discrete category for independent variables when applicable.

### Statistical analysis

We generated frequencies for all collected data. We reported the mean and standard deviation (SD) of continuous variables, and the number of occurrences represented as proportions for categorical variables. Age was negatively skewed and pre-disability gross income was positively skewed and were therefore entered as categorical variables into our regression model; by decade for age, starting at age 15 and ending at age 65, and by quartiles for pre-disability income.

We performed a time-to-event analysis using a Cox proportional hazards regression model to assess the association between time to claim closure and all 11 independent variables described in Table 1. To avoid overfitting our models, we required at least 10 observations per variable term for our Cox regression model, for a total of 190 disabled workers. We set a threshold of at least 50 observations per category for each independent factor in our regression

model to provide some reassurance that each variable had sufficient discriminant power to detect an association with claim duration, if such an association existed.

For claims that were unresolved when the data was extracted, we used 2-years after claim approval as a censoring point. In order to be more stringent and minimize the likelihood of spurious findings, an independent variable was considered statistically significant if it had a p-value of ≤0.01 in our adjusted model. We calculated adjusted hazard ratios (HRs) for our time-to-event analyses, their associated 99% confidence intervals (CIs), and the associated p-values. We assessed each independent variable in our model to ensure that the proportional hazards assumption was met, by entering each variable in the model separately and calculating its interaction with time. We considered a p-value of ≤0.05 for the interaction term as significant. We reported the HRs for independent variables that violated the proportional hazards assumption at 60-days, 6-months, and 1-year. We conducted a sensitivity analysis to investigate the impact of entering receipt of WSIB-reimbursed chiropractic care or physiotherapy as time-dependent covariates in a Cox proportional hazards regression model. This approach accounts for when treatments were initiated during the course of the disability claim.

### Pooling data from similar studies

When possible, we pooled the association between early opioid, physiotherapy, or chiropractic care and claim duration in our sample with similar data from observational studies identified through a systematic review (search strategy, Table S4 in File S1). We considered studies to be similar if they enrolled Workers' Compensation patients with uncomplicated LBP and, explored

Predicting Workers' Compensation claim duration

the association of early treatment with opioids, physiotherapy or chiropractic care with claim duration. Using a standardized, pilot-tested forms, 2 reviewers screened, independently and in duplicate, titles and abstracts of identified citations and then full texts of potentially eligible studies. The same reviewers extracted patient characteristics, methodology, and measures of association between early use of opioids, physiotherapy or chiropractic care and disability claim duration from eligible articles.

We used the following criteria to gauge risk of bias: (1) representativeness of the study population (low risk of bias when using random sampling, consecutive sampling, or data collected from national or international cancer registry, high risk of bias when the source of study population was not reported or acquired through convenience sampling); (2) validity of outcome assessment (low risk of bias when claim duration was acquired directly from the benefits administrator); (3) proportion of lost to follow-up (high risk of bias if >20%); and (4) whether or not predictive models were appropriately adjusted (low risk of bias if adjusted for age, gender, and illness severity).

When possible, we pooled measures of association between early opioid, physiotherapy, or chiropractic care and claim duration, and presented the pooled estimate as a HR and the associated 95%CI. When necessary, we converted odds ratios (ORs) to a relative risk (RR), then to a HR, using the following formula: <sup>24</sup>

$$RR = OR/(1-P_0 + P_0 \times OR)$$

$$HR = (In(1-RR \times P_0))/(In(1-P_0))$$

P<sub>0</sub> is the proportion of patients in the control group who had an event by the follow-up time.

Predicting Workers' Compensation claim duration

We used random-effects meta-analyses, which are usually conservative in that they take both within-study and between-study variability into account. We examined heterogeneity using both a chi-squared test and the I<sup>2</sup> statistic, the latter being the percentage of the total variation in outcomes that is associated with between-study variability (i.e., true differences between studies rather than with sampling error (chance)). Heterogeneity of 0% to 40% was considered 'might not be important', 30% to 60% to be 'moderate heterogeneity', 50% to 90% to be 'substantial heterogeneity', and 75% to 100% to be 'considerable heterogeneity'. The Cochrane Collaboration has proposed overlapping categories, to convey that there are no strict cut-offs for interpreting heterogeneity, and that this decision will depend on the magnitude and direction of effects, as well as the strength of evidence for heterogeneity.

We used the GRADE approach to summarize the certainty of evidence for the effect of early opioid use on claim duration as high, moderate, low, or very low. <sup>28</sup> Using GRADE, observational studies begin as moderate certainty but can be rated down due to: (1) risk of bias; (2) inconsistency; (3) indirectness; (4) imprecision; or (5) publication bias. GRADE suggests considering rating up quality of evidence one level when methodologically rigorous observational studies show at least a two-fold reduction or increase in risk, and rating up two levels for at least a five-fold reduction or increase in risk. <sup>29</sup> We assessed publication bias by visually observing asymmetry of funnel plots, but only if there were ≥10 studies eligible for meta-analysis. We performed all statistical analyses using SAS version 9.2 (SAS Institute Inc., Cary, NC, USA). All hypothesis tests were 2-tailed and p≤ 0.05 was considered statistically significant.

We estimated the cumulative proportion of claims closed in our WSIB dataset at 90-days for disabled workers that did, and did not, receive early opioids by using the following formula:

$$P_1 = 1 - (1 - P_0)^{HR}$$

where  $P_1$  is the cumulative proportion of claims closed by 90-days in the early opioid group,  $P_0$  is the cumulative proportion of claims closed by 90-days in the group that did not receive early opioids, and HR is the pooled estimate of the hazard ratio from our meta-analysis.

# **Results**

Table 2 presents the baseline characteristics for the 1,442 disabled workers included in our analysis. The Ontario WSIB reimbursed 786 (55%) claimants for physiotherapy and 391 (27%) for chiropractic care. In the first 4 weeks of their claim, 27% (n=388) were reimbursed for ≥3 physiotherapy treatments, 17% (n=247) were reimbursed for ≥3 chiropractic treatments, and 9% of claimants (n=136) were reimbursed for an opioid prescription. Figure 1 presents the Kaplan-Meier curve for time to claim closure for LBP claimants. Most workers (67%, n=966) had their claim closed by 90 days, 84% (n=1211) by 180 days, and 91% by 1 year (n=1312); 1348 (93.5%) claims were closed prior to 2 years and 94 (6.5%) were censored.

#### **Time-to-Event Analysis**

Our adjusted regression analysis showed that older age (e.g., HR for age ≥55 versus <25 = 0.52; 99% CI = 0.36, 0.74) and opioid prescription reimbursed by the Ontario WSIB in the first 4-weeks of claim (HR = 0.68; 99% CI = 0.53, 0.88) were associated with longer claim duration. The hazard ratios for pre-disability income and receiving reimbursement for early chiropractic care with claim duration were not proportional over time (p=0.001 and 0.031, respectively), and older claims showed a significant association of greater pre-disability income with longer claim duration (e.g. HR for active claims at 1-year with a pre-disability income >\$902/week versus ≤\$480/week = 0.34; 99% CI = 0.17, 0.68). Working for an employer that had a RTW program (HR = 1.78; 99% CI = 1.45, 2.18), and missing data regarding union membership (HR = 1.27, 99%CI = 1.01, 1.59) were associated with shorter claim duration. Contrary to our predictions, neither

early receipt of WSIB-reimbursement for physiotherapy (HR = 1.01; 99% CI = 0.86, 1.19) nor chiropractic care (e.g. HR for active claims at 60-days = 1.15; 99% CI = 0.94, 1.41) were associated with claim duration (Table 3). We found no important differences using alternative analytic methods (Table S3 in File S1). Figures S2-S4 present the Kaplan-Meier curves for time to claim closure for LBP claimants who received reimbursement for early opioid prescription, physiotherapy and chiropractic care.

#### **Systematic Review**

Our systematic review of observational studies identified identified 2998 unique records, of which we retrieved 99 in full text; three were eligible for our review and explored early opioid use (Figure S5 in File S1). All 3 observational studies that explored early use of opioids for Workers' Compensation claims due to uncomplicated LBP reported a significant association with prolonged claim duration, and two studies adjusted for injury severity in their regression models (Table 4). When our results were pooled with the 2 studies that reported measures of association in relative units,  $^{14, 15}$  resulting in a total of 51,069 participants, the association between early opioid use and prolonged claim duration was consistent with our findings (adjusted HR = 0.57, 95% CI = 0.48 to 0.69), heterogeneity test p=0.02,  $I^2$ =75%; moderate certainty evidence) (Figure 2). Applying this effect to our WSIB dataset means that, at 90-days, 69% of workers without reimbursement for early opioids had resolved their disability claim versus 49% of workers who received reimbursement for early opioids.

The pooled effect was associated with substantial heterogeneity; however, statistical tests of heterogeneity can be misleading when sample sizes are very large and confidence

intervals for measures of association are therefore very narrow. 30 These results provide an excellent example of the phenomenon. The three studies all show consistent, large effect estimates (0.52 to 0.69), which increases confidence in our findings.



# **Discussion**

#### **Statement of Principle Findings**

Our analysis of the Ontario WSIB's administrative data revealed that older claimants who were disabled due to uncomplicated LBP and who are reimbursed for opioid prescription in the first 4 weeks of their claim are more likely to experience prolonged claim duration. Higher pre-injury income was also associated with prolonged claim duration, but only among persistent claims. Injured workers employed by organizations with a RTW program and/or missing information on union affiliation are likely to resolve their claim faster. Neither early receipt of reimbursement for physiotherapy or chiropractic care for uncomplicated LBP was associated with claim duration.

#### **Strengths & Weaknesses**

A priori specification of our regression model and stating the anticipated direction of included independent variables, as well as the assessment of the proportional hazards assumption for all independent variables, provide greater confidence in our findings. The Ontario WSIB does not capture any measure of LBP injury severity and we were therefore unable to adjust for this factor, but we attempted to include patients with similar injuries by restricting our cohort to acute, uncomplicated LBP. There may still be important differences in injury severity in our cohort, but nevertheless our findings regarding the association with early opioid use and delayed claim recovery are consistent with other studies that have adjusted for low back injury severity. <sup>13, 14</sup> Other strengths include limited missing data, correction of

Predicting Workers' Compensation claim duration

identifiable data errors and inconsistencies, and validation checks to ensure the accuracy of the data used to inform our regression model.

Our study has several limitations. First, it was a retrospective cohort study in which the reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are uncertain. Thus, despite our adjustments for potential confounders, it remains possible that LBP claimants who chose to receive these healthcare interventions were prognostically different than those claimants who did not. Second, the WSIB database captures only those physiotherapy or chiropractic treatments that are reimbursed by the WSIB, and it is possible that some patients paid out-of-pocket to receive these services. It is highly unlikely that patients would have received opioids outside of WSIB reimbursement (personal communication, Dr. Norman Buckley, Chair of Anesthesiology, McMaster University). Third, a number of variables that may be important to consider were unavailable (e.g., patient expectations regarding recovery<sup>31</sup>), and some variables were not optimally collected. For example, chiropractic and physiotherapy are professions and not modalities, and there were no details of treatment provided. Fourth, our study focused on workers with LBP who were receiving disability benefits from the Ontario WSIB for at least 4-weeks in 2005, and we cannot say whether our findings are generalizable to other disabled workers. We are, however, unaware of any major changes in practices among Ontario chiropractors or physiotherapists since 2005, and there is evidence that both rates of opioids prescriptions and average morphine equivalent dose for non-malignant pain have increased since 2005, which would suggest that our findings regarding early reimbursement for opioids apply to a greater proportion of current WSIB LBP claimants. 32 Finally, our primary outcome, claim closure, is a surrogate for patient-important outcomes such as functional

restoration or return to work; however, claim closure and faster claim resolution is associated with functional recovery among adults disabled by non-severe whiplash injuries, which provides some assurances that patients who resolve their disability claim are also likely to experience clinical improvement.

### Our Findings in the Context of Other Relevant Literature

Our finding that older age is associated with prolonged claim duration is consistent with the literature.<sup>34</sup> We predicted that injured workers employed by companies that had formal return to work programs would resolve their claim faster, and this was supported by our findings. It is unclear why missing data for union membership was associated with shorter claim duration. Similarly, reasons why higher pre-disability income was associated with prolonged claim duration, but only among persistent claims, are uncertain. Possibilities include that injured workers with higher salaries who do not resolve their claim in the initial 6-months may find it more difficult to identify suitable employment at similar earnings levels, or that compared to other workers (i.e. with lower pre-disability income) workers with higher salaries can better afford to live with limited compensatory income for longer periods of time.

Although there are no randomized controlled trials exploring the effect of physiotherapy, chiropractic care, or early opioid use for workers with uncomplicated LBP receiving lost-time compensation benefits (Table S2 in File S1), our systematic review identified 6 observational studies that are relevant to our findings. Turner and colleagues followed 1,885 workers from Washington for 1 year after they had been awarded Workers' Compensation benefits for acute LBP. In a comprehensive regression model adjusted for

Predicting Workers' Compensation claim duration

multiple sociodemographic, employment, clinical, healthcare, and administrative factors – including injury severity – they found that workers who attended a chiropractor first, versus a primary care provider, were significantly less likely to remain on disability benefits at 1 year (adjusted OR = 0.41, 95% CI = 0.24, 0.70).

Wasiak and colleagues found that workers in Florida, USA, with low back injuries who were reimbursed for limited chiropractic care (<30 days) experienced an 8.6% shorter duration of work disability versus Workers' Compensation claimants who were reimbursed for prolonged chiropractic care (>30 days). The authors did exclude severe injuries from their population, but were unable to adjust for injury severity within their sample. Further, chiropractic care was collected after baseline and not treated as time-dependent, and so it is not surprising that claims with longer duration also received more chiropractic care.

Lemstra and Olszynski explored the effect of standard care (which included long waiting lists for physiotherapy) to provision of rapid rehabilitation services on Workers' Compensation claim duration from a company in Saskatchewan, Canada. After adjusting for a number of factors, including age and injury severity, longer claim duration was associated with both chiropractic care (adjusted HR = 2.88, 95% CI = 1.45, 5.73) and physical therapist involvement (adjusted HR = 19.88, 95% CI = 7.95, 39.77). The authors collected healthcare provider utilization data after baseline and did not treat these variables as time-dependent, and so an alternative explanation is that claims with longer duration are simply more likely to involve either chiropractors or physical therapists.

Canada is currently the largest per-capita consumer of opioids in the world; <sup>37</sup> however, prescribing patterns in Ontario show considerable variation between primary care providers. <sup>38</sup>

Predicting Workers' Compensation claim duration

Workers' Compensation data from the US has shown an almost 10-fold range (5.7% to 52.9%) in the early prescription of opioid medications between various states, suggesting that local prescribing patterns have significant influence on the use of these analgesics. Findings from a study of Workers' Compensation nonspecific LBP claims (that did not adjust for injury severity) revealed that, compared with a no opioid reference group, the odds of chronic work loss were six times greater for claimants who used strong opioids and 11-14 times greater for claimants with opioid prescriptions which exceeded 90 days. We found that reimbursement for early opioid use by the Ontario WSIB was associated with prolonged claim duration for uncomplicated back pain, and pooling of our data with similar studies <sup>14, 15</sup> shows a consistent effect, which increases confidence in our findings.

#### **Implications & Future Research**

To manage their growing unfunded liability, the Ontario WSIB has focused on increasing their claim denial rate, decreasing benefits to injured workers, reducing WSIB staff, and raising employer premiums. <sup>9, 10</sup> Most employers are obligated to pay WSIB premiums because they are legally bound to provide injury benefits to their employees, and the Ontario WSIB is protected by laws prohibiting competition in the marketplace. Another strategy is to optimize clinical management of injured workers. Our findings, which were not adjusted for illness severity, suggest that receiving reimbursement from the Ontario WSIB for early chiropractic care or physiotherapy for acute, uncomplicated LBP is not associated with shorter time to claim closure. Receiving reimbursement for early opioids was linked with longer claim duration.

Predicting Workers' Compensation claim duration

However, observational data cannot establish causality and high quality, randomized controlled trials are urgently needed to confirm or refute our findings.

#### **Author affiliations**

- <sup>1</sup> The Michael G. DeGroote Institute for Pain Research and Care, McMaster University,
  Hamilton, Canada
- <sup>2</sup> Department of Anesthesia, McMaster University, Hamilton, Canada
- <sup>3</sup> Department of Clinical Epidemiology and Biostatistics, McMaster University, Hamilton, Canada
- <sup>4</sup> Stanford Prevention Research Center, Department of Medicine, Stanford University, Stanford, California, USA
- <sup>5</sup> Department of Anaesthesia & Pain Medicine, the Hospital for Sick Children, Toronto, Canada

### Acknowledgements

The authors thank Dr. Gordon Guyatt for helpful discussion.

#### **Contributors**

JWB, SE, D-HA and SDW designed the study. DH-A and LW conducted data analysis and SDW provided statistical advice. JWB, SE, D-HA and SDW were involved in interpreting the data. RC designed and conducted all literature searches. JWB drafted the manuscript and wrote the final version. All authors critically revised the manuscript, provided comment and approved the final version for publication.

#### **Funding**

This study was funded by research grants from the Ontario Workers Safety and Insurance Board Research Advisory Council and the Ontario Chiropractic Association. Shanil Ebrahim was

supported by a MITACS Elevate Postdoctoral Fellowship Award. The funding sources had no role in design or conduct of the study; the collection, management, analysis, or interpretation

Predicting Workers' Compensation claim duration

#### **Competing interests**

Jason Busse acts as a consultant to Prisma Health Canada, a private incorporated company funded by employers and insurers that consults on and manages long-term disability claims.

of the data; or the preparation, review, or approval of the manuscript.

#### **Ethics approval**

The study was approved by the Health Sciences Research Ethics Board of the University of Toronto.

#### Provenance and peer review

Not commissioned; externally peer reviewed.

#### Data sharing statement

No additional data are available.

#### **Open Access**

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work noncommercially, and license their derivative works on different terms,

provided the original work is properly cited and the use is non-commercial. See: http:// creativecommons.org/licenses/by-nc/4.0/



Predicting Workers' Compensation claim duration

#### References

- 1. Thiese MS, Hegmann KT, Wood EM, Garg A, Moore JS, Kapellusch JM, et al. Low-back pain ratings for lifetime, 1-month period, and point prevalences in a large occupational population. *Hum Factors* 2014; 56: 86-97.
- 2. Deyo RA, Phillips WR. Low back pain. A primary care challenge. *Spine (Phila Pa 1976)* 1996; 21: 2826-2832.
- 3. Hoy D, March L, Brooks P, Blyth F, Woolf A, Bain C, et al. The global burden of low back pain: estimates from the Global Burden of Disease 2010 study. *Ann Rheum Dis* 2014; 73: 968-974.
- Brown A, Angus D, Chen S, Tang Z, Milne S, Pfaff J, et al. Costs and outcomes of chiropractic treatment for low back pain [Technology report no 56]. Ottawa: Canadian Coordinating Office for Health Technology Assessment, 2005.
- 5. WSIB-CSPAAT Ontario. By the Numbers: 2012 WSIB Statistical Report. Schedule 1 Toronto, ON: WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2014. [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S1.pdf] (Accessed August 3, 2014).
- 6. WSIB-CSPAAT Ontario. By the Numbers: 2012 WSIB Statistical Report. Schedule 2 WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2014.
  [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S2.pdf] (Accessed August 3, 2014).

Predicting Workers' Compensation claim duration

- 7. WSIB-CSPAAT Ontario. First Quarter Sufficiency Report to Stakeholders WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2013.
  [http://www.wsib.on.ca/files/Content/SufficiencySufficiencyReport2013/2013Q1WSIBSufficiencyReport.pdf] (Accessed August 3, 2014).
- 8. Ontario WSIB funding fiasco [Editorial] (November 18, 2008). National Post. 2008. [http://www.nationalpost.com/opinion/story.html?id=eb83e348-1660-4771-9804-e47a502aa756] (Accessed August 3, 2014).
- 9. UFCW Canada. WSIB President gets \$80K bonus on the backs of injured workers. Media & News Canada's Best Labour and Social Justice News. 2012.
  [http://www.ufcw.ca/index.php?option=com\_content&view=article&id=2760:wsib-president-gets-80k-bonus-on-the-backs-of-injured-workers&Itemid=6&lang=en]
  (Accessed August 3, 2014).
- 10. WSIB raises its rates to tackle \$12 billion in unfunded liabilities. Machinery and Equipment MRO Maintenance, Repair and Operations. 2010.
  [http://www.mromagazine.com/news/wsib-raises-its-rates-to-tackle-12-billion-in-unfunded-liabilities/1000391835/?&er=NA] (Accessed August 3, 2014).
- 11. Henschke N, Maher CG, Refshauge KM, Herbert RD, Cumming RG, Bleasel J, et al.

  Prognosis in patients with recent onset low back pain in Australian primary care:
  inception cohort study. *BMJ* 2008; 337: a171.

- 12. Childs JD, Fritz JM, Wu SS, Flynn TW, Wainner RS, Robertson EK, et al. Implications of early and guideline adherent physical therapy for low back pain on utilization and costs. *BMC*Health Serv Res 2015; 15: 150.
- 13. 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: 2127-2132.
- 14. Franklin GM, Stover BD, Turner JA, Fulton-Kehoe D, Wickizer TM. 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.
- 15. Gross DP, Stephens B, Bhambhani Y, Haykowsky M, Bostick GP, Rashiq S. Opioid prescriptions in canadian workers' compensation claimants: prescription trends and associations between early prescription and future recovery. *Spine (Phila Pa 1976)* 2009; 34: 525-531.
- 16. Turner JA, Franklin G, Fulton-Kehoe D, Sheppard L, Stover B, Wu R, et al. ISSLS prize winner: early predictors of chronic work disability: a prospective, population-based study of workers with back injuries. *Spine (Phila Pa 1976)* 2008; 33: 2809-2818.
- von Elm E, Altman DG, Egger M, Pocock SJ, Gotzsche PC, Vandenbroucke JP. The
   Strengthening the Reporting of Observational Studies in Epidemiology (STROBE)
   Statement: guidelines for reporting observational studies. *Int J Surg* 2014; 12: 1495-1499.
- 18. Collins GS, Reitsma JB, Altman DG, Moons KG. Transparent Reporting of a multivariable prediction model for Individual Prognosis Or Diagnosis (TRIPOD): the TRIPOD Statement.

  Br J Surg 2015; 102: 148-158.

- 19. da CMCL, Maher CG, Hancock MJ, McAuley JH, Herbert RD, Costa LO. The prognosis of acute and persistent low-back pain: a meta-analysis. *CMAJ* 2012; 184(11): E613-E624.
- 20. Frank JW, Brooker AS, DeMaio SE, Kerr MS, Maetzel A, Shannon HS, et al. Disability resulting from occupational low back pain. Part II: What do we know about secondary prevention? A review of the scientific evidence on prevention after disability begins. *Spine* (*Phila Pa 1976*) 1996; 21: 2918-2929.
- 21. Busse J, Steenstra I, Riva J, Ebrahim S, de Bruin L, Guyatt G. Predictors of prolonged recovery following acceptance for disability benefits: a systematic review of observational studies. *Occup Environ Med* 2011; 68(Suppl 1): A97.
- 22. Schandelmaier S, Ebrahim S, Burkhardt SC, de Boer WE, Zumbrunn T, Guyatt GH, et al.

  Return to work coordination programmes for work disability: a meta-analysis of randomised controlled trials. *PLoS ONE* 2012; 7: e49760.
- 23. Harrell FE. Multivariate modeling strategies. In: Harrell FE, editor. Regression Modeling Strategies With Applications to Linear Models, Logistic Regression and Survival Analysis. New York, NY: Springer; 2001. p. 53-85.
- 24. Zhang J, Yu KF. What's the relative risk? A method of correcting the odds ratio in cohort studies of common outcomes. *JAMA* 1998; 280: 1690-1691.
- 25. Montori V, Ioannidis J, Cook DJ. Fixed-effects and random-effects models. In: Guyatt GH, Rennie D, Meade MO, Cook DJ, editors. Users' guides to the medical literature: A manual for evidence-based clinical practice (2nd ed). USA: McGraw Hill; 2008. p. 555-62.
- 26. Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. BMJ 2003; 327: 557-560.

- 27. Higgins JPT, Green Se. Cochrane Handbook for Systematic Reviews of Interventions

  Version 5.1.0 [updated March 2011]: The Cochrane Collaboration; 2011.
- 28. Atkins D, Best D, Briss PA, Eccles M, Falck-Ytter Y, Flottorp S, et al. Grading quality of evidence and strength of recommendations. *BMJ* 2004; 328: 1490.
- 29. Guyatt GH, Oxman AD, Sultan S, Glasziou P, Akl EA, Alonso-Coello P, et al. GRADE guidelines: 9. Rating up the quality of evidence. *J Clin Epidemiol* 2011; 64: 1311-1316.
- 30. Rucker G, Schwarzer G, Carpenter JR, Schumacher M. Undue reliance on I(2) in assessing heterogeneity may mislead. *BMC Med Res Methodol* 2008; 8: 79.
- 31. Cole DC, Mondloch MV, Hogg-Johnson S. Listening to injured workers: how recovery expectations predict outcomes--a prospective study. *CMAJ* 2002; 166: 749-754.
- 32. Gomes T, Mamdani MM, Paterson JM, Dhalla IA, Juurlink DN. Trends in high-dose opioid prescribing in Canada. *Can Fam Physician* 2014; 60: 826-832.
- 33. Cote P, Hogg-Johnson S, Cassidy JD, Carroll L, Frank JW. The association between neck pain intensity, physical functioning, depressive symptomatology and time-to-claim-closure after whiplash. *J Clin Epidemiol* 2001; 54: 275-286.
- 34. Hadler NM. The bane of the aging worker. *Spine (Phila Pa 1976)* 2001; 26: 1309-1312.
- 35. Wasiak R, Kim J, Pransky GS. The association between timing and duration of chiropractic care in work-related low back pain and work-disability outcomes. *J Occup Environ Med* 2007; 49: 1124-34.
- 36. Lemstra M, Olszynski WP. The effectiveness of standard care, early intervention, and occupational management in worker's compensation claims. *Spine (Phila Pa 1976)* 2003; 28: 299-304.

Predicting Workers' Compensation claim duration

 Pain and Policy Studies Group. PPSG Opioid Consumption Motion Chart. University of Wisconsin Paul P Carbone Comprehensive Cancer Center Madison, Wisconsin. 2011 [https://ppsg.medicine.wisc.edu/chart] (Accessed May 15, 2015).

- 38. Dhalla IA, Mamdani MM, Gomes T, Juurlink DN. Clustering of opioid prescribing and opioid-related mortality among family physicians in Ontario. *Can Fam Physician* 2011; 57: e92-e96.
- 39. Webster BS, Cifuentes M, Verma S, Pransky G. Geographic variation in opioid prescribing for acute, work-related, low back pain and associated factors: a multilevel analysis. *Am J Ind Med* 2009; 52: 162-171.
- 40. Volinn E, Fargo JD, Fine PG. Opioid therapy for nonspecific low back pain and the outcome of chronic work loss. *Pain* 2009; 142: 194-201.

Predicting Workers' Compensation claim duration

# **Figure Legends**

Figure 1: Kaplan-Meier curve for time to claim closure

Figure 2: The association between early opioid use/prescription and claim duration

## **Supporting Information Legends**

- Table S1: Search strategy for the Cochrane Back Group trials registry
- Table S2: Randomized controlled trials of acute low back pain assessing the effect of opioids, chiropractic care, or physical therapy, and their generalizability to injured workers receiving disability benefits
- Table S3: Sensitivity analysis of factors associated with time to claim closure, entering chiropractic care and physiotherapy as time-dependant co-variates (n=1,442)
- Table S4: Search strategy for observational studies exploring the association of early opioid, physiotherapy, or chiropractic care for workers with acute low back pain with claim duration
- Figure S1: A flow diagram of the literature search process for randomized controlled trials assessing the effect of opioids, physiotherapy, or chiropractic care for acute low back pain.
- Figure S2: Kaplan-Meier curve for early reimbursement for Opioid prescription
- Figure S3: Kaplan-Meier curve for early reimbursement for physiotherapy
- Figure S4: Kaplan-Meier curve for early reimbursement for chiropractic care
- Figure S5: A flow diagram of the literature search process for observational studies assessing the effect of early opioids, physiotherapy, or chiropractic care for acute low back pain.

#### Table 1: Description of model variables

Variable	Anticipated Direction of Effect
Age (in decades)	older age: -
Gender	female: -
First language	non-English: -
Pre-disability income	higher income: -
Reimbursement for opioid prescription	opioid reimbursement: -
in the first 4-weeks of a claim	
Prior claim(s)	prior claim: -
Union membership	Union member: +
Employer RTW-program	RTW program: +
Work-relatedness	work-related: +
Reimbursement for ≥3 chiropractic	DC care: +
treatments in the first 4-weeks of a	
claim	
Reimbursement for ≥3 physiotherapy	PT care: +
treatments in the first 4-weeks of a	
claim	

- +: associated with faster claim closure
- -: associated with slower claim closure

Predicting Workers' Compensation claim duration

Age in years, mean (SD)	41.3 (10.5)
Gender, n (%)	
Female	552 (38.3)
Male	890 (61.7)
First language, n (%)	
English	1372 (95.1)
Other	70 (4.9)
Pre-disability income (dollars/week)	
mean (SD)	731.4 (332.5)
Opioid prescription reimbursed by WSIB in the first 4-weeks of	
claim, n (%)	
Yes	136 (9.4)
No	1306 (90.6)
Prior WSIB claim, n (%)	
Yes	1091 (75.7)
No	351 (24.3)
Union membership, n (%)	
Yes	610 (42.3)
No	656 (45.5)
Missing data	176 (12.2)
Employer RTW program, n (%)	
Yes	1042 (72.3)
No	278 (19.3)
Missing data	122 (8.5)
Employer doubts work-relatedness of injury, n (%)	
Yes	195 (13.5)
No	1051 (72.9)
Missing data	196 (13.6)
Chiropractic care reimbursed by WSIB during claim, n (%)	391 (27.1)
Early Chiropractic care (≥3 treatments received within the first 28	21-11-1
days), n (%)	247 (17.1)
Dhusiathanan raisah rusad hu WCID duning alaina ra (0/)	700 (54.5)
Physiotherapy reimbursed by WSIB during claim, n (%)	786 (54.5)
Early Physiotherapy (≥3 treatments received within the first 28 days), n (%)	388 (26.9)

Table 3: Factors associated with time to claim closure (n=1,442)\*

Univariate			Multivariable		
	Hazard Ratio (99% CI)	p-value	Adjusted Hazard Ratio (99% CI)	p-value	
Baseline predictors					
Age categories in years		<0.001		<0.001	
15 to <25	1.00		1.00		
25 to <35	0.88 (0.65, 1.19)		0.79 (0.58, 1.09)		
35 to <45	0.78 (0.59, 1.04)		0.70 (0.52, 0.95)		
45 to <55	0.76 (0.56, 1.02)		0.67 (0.49, 0.91)		
55 to 65	0.56 (0.40, 0.80)		0.52 (0.36, 0.74)		
Gender		0.114		0.446	
Females	1.09 (0.95, 1.26)		0.96 (0.82, 1.12)		
Males	1.00		1.00		
First language		0.137		0.312	
English	1.00		1.00		
Other	0.83 (0.59, 1.15)		0.88 (0.63, 1.23)		
Pre-disability income in					
dollars per week					
At 60 days:					
<u>,</u> ≤480	1.00	-	1.00	-	
481-694	1.09 (0.89, 1.34)	0.273	1.03 (0.83, 1.27)	0.749	
695-920	1.13 (0.92, 1.39)	0.137	0.96 (0.77, 1.21)	0.678	
>920	1.08 (0.88, 1.33)	0.326	0.93 (0.74, 1.18)	0.443	
At 180 days:	, , ,				
<b>≤</b> 480	1.00	-	1.00	_	
481-694	0.89 (0.68, 1.17)	0.267	0.89 (0.67, 1.17)	0.266	
695-920	0.84 (0.63, 1.12)	0.115	0.73 (0.54, 0.98)	0.006	
>920	0.72 (0.54, 0.97)	0.004	0.63 (0.46, 0.86)	<0.001	
At 365 days:			, , ,		
≤480	1.00	_	1.00	_	
481-694	0.65 (0.36, 1.18)	0.064	0.71 (0.38, 1.31)	0.147	
695-920	0.53 (0.28, 1.02)	0.012	0.47 (0.24, 0.91)	0.003	
>920	0.39 (0.20, 0.75)	<0.001	0.34 (0.17, 0.68)	<0.001	
Early Reimbursement for		<0.001		<0.001	
Opioid prescription (within					
the first 28 days)					
Yes	0.62 (0.48, 0.80)		0.68 (0.53, 0.88)		
No	1.00		1.00		
Prior claims		0.306		0.709	
Yes	1.07 (0.91, 1.26)		1.03 (0.86, 1.22)		
No	1.00		1.00		
Union membership		<0.001		0.016	
Yes	1.29 (1.11, 1.50)		1.14 (0.96, 1.35)		
No	1.00		1.00		
Missing	1.34 (1.07, 1.68)		1.27 (1.01, 1.59)		
Employer RTW program	2.0 . (2.07, 2.00)	<0.001		<0.001	
Yes	1.73 (1.43, 2.09)	10.001	1.78 (1.45, 2.18)	10.001	
No	1.00		1.00		
Missing	1.17 (0.87, 1.58)		1.17 (0.86, 1.59)		
Doubt Work-relatedness	1.17 (0.07, 1.30)	0.174	1.17 (0.00, 1.00)	0.138	
Yes	0.94 (0.76, 1.16)	0.1/4	0.88 (0.71, 1.08)	0.130	

No	1.00		1.00	
Missing	1.13 (0.92, 1.39)		1.08 (0.87, 1.33)	
Early Reimbursement for				
Chiropractic care				
At 60 days:	1.19 (0.99, 1.45)	0.017	1.15 (0.94, 1.41)	0.067
At 180 days:	0.91 (0.65, 1.24)	0.437	0.90 (0.65, 1.24)	0.392
At 365 days:	0.60 (0.29, 1.23)	0.067	0.61 (0.29, 1.29)	0.089
Early Reimbursement for	0.98 (0.84, 1.15)	0.726	1.01 (0.86, 1.19)	0.848
Physiotherapy				

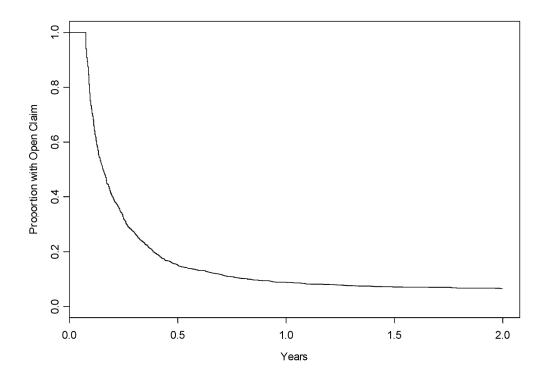
<sup>\*</sup> HR>1 indicates faster claim closure; RTW = return to work; Early reimbursement for chiropractic or physiotherapy = 3 or more treatments received within the first 28 days 3 01 mm.s

Table 4: Observational studies exploring the association between early opioid use and Workers' Compensation claim duration

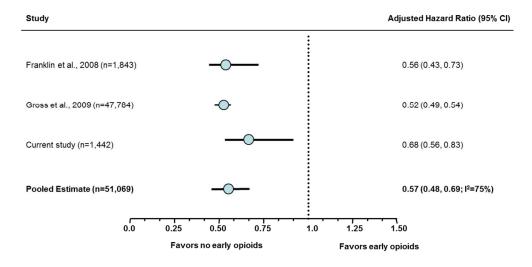
Study	Population	Opioid variable tested	Adjustments	Dependent variable	Results*
Webster et al.,	8,443 American	Receipt of	Injury severity,	Change in	1-140mg MEA
2007	Workers'	opioids within	age, gender,	mean	5.4 days, 95%CI = -14.6 to 25.0
	Compensation	the first 15 days	length of job	disability	141-225mg MEA
	claimants with	of claim	tenure	duration	21.9 days, 95%CI = 3.2 to 40.6
	new-onset,				226-450mg MEA
	disabling LBP	100			43.8 days, 95%CI = 23.7 to 63.9
	_				>450mg MEA
					69.1 days, 95%CI = 49.3 to 89.0
Franklin et al.,	1,843 Washington,	Reimbursement	Age, gender,	Receipt of	1-150mg MED
2008	US, Workers'	for opioids	race,	wage	OR = 1.9, 95%CI = 1.2 to 3.1
	Compensation	within 6 weeks	education,	replacement	151-300mg MED
	claimants with	of 1 <sup>st</sup> medical	injury severity,	benefits at 1-	OR = 2.0, 95%Cl = 1.2 to 3.3
	new-onset,	visit for LBP	pain intensity,	year	301-650mg MED
	disabling LBP		Roland		OR = 1.6, 95%CI = 0.9 to 2.6
	_		disability		>650mg MED
			questionnaire		OR = 1.9, 95%CI = 1.2 to 2.9
Gross et al.,	47,784 Alberta,	Reimbursement	Age, gender,	Receipt of	No early opioids
2009	Canada, Workers'	for opioids	annual salary,	wage	HR = 1.94, 95%CI = 1.86 to 2.02
	Compensation	within the first 2	year of claim,	replacement	
	claimants with	weeks of claim	number of	benefits at 1-	
	new-onset,		previous	year	
	disabling LBP		claims		

<sup>\*</sup> The reference group is no early receipt of opioids for Franklin et al., and Webster et al. 95% CI = 95% confidence interval; LBP = low back pain; MEA = morphine equivalent amount; MED = morphine equivalent dose; OR = odds ratio; HR = hazard ratio

Predicting Workers' Compensation claim duration



Kaplan-Meier curve for time to claim closure 181x130mm (300 x 300 DPI)



The association between early opioid use/prescription and claim duration 177x88mm (300 x 300 DPI)

#### Table S1: Search strategy for the Cochrane Back Group trials registry\*

Search Strategy in Cochrane Library:

\*:ti,ab,kw in Trials, with Back Group in Review Groups 6476

Endnote search strategy:

Any field contains: acute AND Any field contains chiropractic (43) Any field contains: acute AND Any field contains physiotherapy\* (87)

Any field contains: acute AND Any field contains opioid (30)

Any field contains: opioid NOT Any field contains: surgery NOT Any field contains:

chronic (28)

Any field contains: emergency (57) NOT Any field contains: whiplash (18)

<sup>\*</sup>On May 1, 2015 we used the Cochrane Library to search the Cochrane Back Review Group (CBRG) Trials Register, which we exported to Endnote. We also screened the included studies lists of recent CBRG Reviews of chiropractic and physiotherapy interventions for acute low back pain (Franke et al., 2015, Rubinstein et al., 2012).

Table S2: Randomized controlled trials of acute low back pain assessing the effect of opioids, chiropractic care, or physical therapy, and their generalizability to injured workers receiving disability benefits

Study	Participants and Interventions	Representation of injured workers receiving lost-time disability benefits for acute low back pain
Glover 1974	84 patients with unilateral low back pain (LBP), randomized to manipulation or control (sham diathermy)	No mention regarding receipt of lost-time disability benefits
Bergquist- Ullman 1977	217 patients consulting a workplace health centre with acute or subacute LBP randomized to back school, physiotherapy or placebo	At least 88% of patients were enrolled with acute LBP, and of the 217, 184 were "sick-listed" for a median of 21 days during the study, but there was no mention regarding receipt of lost-time disability benefits
Rasmussen 1979	26 patients with LBP duration <3weeks, randomized to manipulation or diathermy	No mention regarding receipt of lost-time disability benefits
Hoehler 1981	95 patients with palpatory cues indicating hyperalgesia or a restricted or painful range of vertebral motion, randomized to rotational manipulation of the trunk or massage	No mention regarding receipt of lost-time disability benefits
Farrell 1982	48 subjects with acute LBP duration <3 weeks, randomized to passive mobilization and manipulation or combination of diathermy, exercises and ergonomic advice	No mention regarding receipt of lost-time disability benefits
Gilbert 1985	270 patients presenting with LBP, randomized to bed rest or physiotherapy	The authors reported that "people who were receiving workman's compensation were also slower to recover." p.794, but they did not report any associated data (e.g. how many patients were receiving disability benefits, or the quantitative results for this subgroup).
Waterworth 1985	112 patients with acute mechanical LBP, randomized to ergonomic advice/ Diflunisal or ergonomic advice/ conservative physiotherapy (ultrasound and exercise) or ergonomic advice/manipulation	The authors enrolled a mixed group of patients, that may include up to 54% receiving lost-time claim benefits, but that the proportion is unequal between study group (ranging from 47% to 63%) and the results are provided for the total population which precludes confident generalizability to only those who were receiving lost-time claim benefits.
Hadler 1987	54 subjects with acute LBP, randomized to mobilization or manipulation	Patients who were receiving disability benefits were explicitly excluded from this trial: "neither workers' compensation nor disability insurance should be at issued [sic] and the acute low-back pain must not be considered work-related."pg 703
MacDonald 1990	95 subjects with LBP duration <4 weeks, randomized to osteopathic manipulation or control (advice to rest and resume activities gradually)	"Less than 30% [of the study group] suffered loss of income because of disability" pg. 366

Cramer 1993	36 subjects with mechanical LBP less than two weeks duration randomized to side-lying manipulation, electrical stimulation and cold packs or control (detuned ultrasound, cold packs and 15-30 sec. gentle massage)	Patients who were receiving disability benefits were explicitly excluded from this trial. Inclusion criteria stipulate "no litigation or workers' compensation" (as per the review by Rubinstein 2012, pg. 52)
Skargren 1997	323 patients with back and neck pain of mixed duration, randomized to chiropractic or physiotherapy.	51% of patients (166 of323) were on "sick-leave" when enrolled (Table 3, pg. 2170), but there was no mention regarding receipt of lost-time disability benefits.
Innes 1998	123 patients with acute LBP, randomized to ketorolac or acetaminophen/codeine	Patients who were receiving disability benefits were limited during enrollment: "Because Workers Compensation status might influence response to therapy, we limited each site to 10 work-related back injuries, hoping to limit such cases to no more than half the total study enrollment." pg. 550
Cherkin 1998	321 adults with acute LBP randomized to the McKenzie method of physical therapy, chiropractic manipulation, or provision of an educational booklet	patients who were receiving disability benefits were explicitly excluded from this trial: "Subjects who were involved in claims for compensation or litigation because of the back injurywere also excluded"; pg. 1022
Seferlis 1998	180 patients sick-listed for < 2 weeks for LBP randomized to General Practitioner Program (rest, sick-leave, analgesics etc.) or Manual Therapy Program (autotraction, manipulation, mobilization etc.) or Intensive Training Program (information, muscle training and general condition training 3x/week for 8 weeks)	All enrolled patients were "sick listed for acute low-back pain for up to 2 weeks", but there is no mention regarding receipt of lost-time disability benefits.
Morton 1999	29 patients with acute mechanical LBP, randomized to manipulation/ exercise or exercise alone	Patients who were receiving disability benefits were explicitly excluded from this trial: "Exclusion criteria were third-party, public liability or workers' compensation claimants"; pg. 185
Veenema 2000	155 patients with musculoskeletal LBP, randomized to meperidine or ketorolac	No mention regarding receipt of lost-time disability benefits
Metscher 2001	192 Patients with acute LBP randomized to dexketoprofen-trometamol or tramadolhydrochloride	Abstract in English, paper in German. No mention in the abstract about receipt of lost-time disability benefits
Palangio 2002	147 patients with acute LBP, first episode or exacerbation of chronic condition with onset <48 hours before enrolment, randomized to combination hydrocodone 7.5 mg and ibuprofen 200 mg (HC/IB) or combination oxycodone 5 mg and acetaminophen 325 mg (OX/AC)	No mention regarding receipt of lost-time disability benefits
Hofstee 2002	250 patients with sciatica of less than 1 months duration randomized to bed rest, physiotherapy or continuation of activities of daily living	No mention regarding receipt of lost-time disability benefits
Johnstone 2002	12 patients with acute LBP with signs of psychological distress (DRAM score Modified Zung score >17) randomized to cognitive behavioral therapy and conventional physiotherapy or conventional physiotherapy alone	Patients with "ongoing medico legal issues" were excluded (pg.183).  No mention regarding receipt of lost-time disability benefits

Childs 2004	131 patients with LBP of median duration of 27 days, randomized to manipulation and exercise or exercise alone	39.8% of patients had missed work due to LBP, Table 2 pg. 925, but there was no mention regarding receipt of lost-time disability benefits
Hoiriis 2004	192 patients with LBP of 3 to 6 weeks duration randomized to chiropractic adjustments with placebo medicine, muscle relaxants with sham adjustments, or placebo medicine with sham adjustments	No mention regarding receipt of lost-time disability benefits
Salvador 2005	28 subjects randomly allocated to a muscle energy technique or transcutaneous electrical nerve stimulation (TENS)	Abstract in English, paper in Portuguese. No mention in the abstract about receipt of lost-time disability benefits
Brennan 2006	123 patients referred to physiotherapy for LBP less than 90 days duration, randomized to manipulation or specific exercise or stabilization	No mention regarding receipt of lost-time disability benefits
Santilli 2006	102 patients with acute moderate to severe radiating LBP of duration <10 days with MRI evidence of disc protrusion, randomized to manipulation or simulated manipulation	No mention regarding receipt of lost-time disability benefits
Hancock 2007	240 subjects with acute LBP duration < 6 weeks, randomized to four groups: control (placebo drug and placebo manipulation) or NSAIDs (diclofenac and placebo manipulation) or manipulation (placebo drug and active manipulation) or manipulation and NSAIDs (diclofenac and active manipulation)	No mention regarding receipt of lost-time disability benefits
Lee 2008	Study of 78 musculoskeletal pain patients, 67% with LBP, randomized to tramadol/paracetemol (n=28 with LBP) or ketorolac/paracetemol (n=24 with LBP)	No mention regarding receipt of lost-time disability benefits
Lau 2008	110 patients with acute LBP, randomized to immediate intervention (advice to stay active, Back Care booklet, reassurance, advice, interferential current therapy) or control (walking training and prescription of walking aids as indicated) followed by outpatient physiotherapy (for both groups)	12% of patients (13 of 110) had work-related injuries (see Table 1), but no mention regarding receipt of lost-time disability benefits
Selkow 2009	20 subjects with acute LBP, randomly allocated to muscle energy technique or sham manual treatment	No mention regarding receipt of lost-time disability benefits
Cleland 2009	112 subjects with LBP, that met 4 out of 5 criteria for a clinical prediction rule for LBP likely to respond to manipulation, randomized to supine thrust manipulation, side-lying thrust manipulation or non-thrust manipulation	Only 6% of patients were unable to work due to LBP. Table 2, pg.2724.
Hallegraef 2009	64 patients with acute nonspecific LBP duration <16 days, randomized to manipulative therapy plus physical therapy or physical therapy alone	No mention regarding receipt of lost-time disability benefits

Sutlive 2009	60 subjects with LBP meeting 3 out of 5 criteria for a clinical prediction rule for LBP likely to respond to manipulation, randomized to lumbopelvic manipulation or neutral gap manipulation	Patients "with litigation pending for their LBP" were excluded.  No mention regarding receipt of lost-time disability benefits
Juni 2009	104 patients with acute LBP duration < 4 weeks, randomized to standard care with manipulation or standard care alone	No mention regarding receipt of lost-time disability benefits
Machado 2010	148 adults with acute LBP duration <6 weeks, randomized to the McKenzie method and first-line care (advice, reassurance and time-contingent acetaminophen) or first-line care alone	Only 3% of participants (4 of 146) were receiving disability benefits for their injury (see Table 1)
Lewis 2011	89 patients with acute LBP duration < 3 months, randomized to strain-counterstrain manual therapy/exercise or exercise alone	No mention regarding receipt of lost-time disability benefits
Biondi 2013	1664 patients with acute LBP, randomized to tapentadol or oxycodone	No mention regarding receipt of lost-time disability benefits
Goertz 2013	91 patients with acute LBP, duration < 4 weeks, randomized to standard medical care and chiropractic manipulation or standard care alone	No mention regarding receipt of lost-time disability benefits
Behrbalk 2014	65 adults with acute LBP, randomized to morphine or morphine/promethazine	No mention regarding receipt of lost-time disability benefits
Eken 2014	137 patients with moderate or severe acute LBP, randomized to paracetemol, morphine or dexketoprophen	No mention regarding receipt of lost-time disability benefits
Tanen 2014	44 patients with acute radicular LBP, randomized to lidocaine or ketorolac	No mention regarding receipt of lost-time disability benefits

#### References for Tables S1 and S2

- 1. Franke H, Fryer G, Ostelo RW, Kamper SJ. Muscle energy technique for non-specific low-back pain. *Cochrane Database Syst Rev* 2015; 2: CD009852.
- 2. Rubinstein SM, Terwee CB, Assendelft WJ, de Boer MR, van Tulder MW. Spinal manipulative therapy for acute low-back pain. *Cochrane Database Syst Rev* 2012; 9: CD008880.
- 3. Glover JR, Morris JG, Khosla T. Back pain: a randomized clinical trial of rotational manipulation of the trunk. *British journal of industrial medicine* 1974; 31: 59-64.
- 4. Bergquist-Ullman M, Larsson U. Acute low back pain in industry. A controlled prospective study with special reference to therapy and confounding factors. *Acta orthopaedica Scandinavica* 1977; 48(Suppl. 170): 1-117.
- 5. Rasmussen GG. Manipulation in treatment of low back pain (a randomized clinical trial. *Man-Med* 1979; 17: 8-10.
- 6. Hoehler FK, Tobis JS, Buerger AA. Spinal manipulation for low back pain. *JAMA* 1981; 245: 1835-1838.
- 7. Farrell JP, Twomey LT. Acute low back pain. Comparison of two conservative treatment approaches. *Medical journal of Australia* 1982; 1: 160-164.
- 8. Gilbert JR, Taylor DW, Hildebrand A, Evans C. Clinical practice of common treatments for low back pain in family practice. *British Medical Journal Clinical Research Ed* 1985; 291: 791-794.
- 9. Waterworth RF, Hunter IA. An open study of diflunisal, conservative and manipulative therapy in the management of acute mechanical low back pain. *New Zealand medical journal* 1985; 98: 372-375.
- 10. Hadler NM, Curtis P, Gillings DB, Stinnett S. A benefit of spinal manipulation as adjunctive therapy for acute low-back pain: a stratified controlled trial. *Spine* 1987; 12: 702-706.
- 11. MacDonald RS, Bell CMJ. An open controlled assessment of osteopathic manipulation in nonspecific low-back pain. *Spine* 1990; 15: 364-370.
- 12. Cramer GD, Humphreys CR, Hondras MA, McGregor M, Triano JJ. The Hmax/Mmax ratio as an outcome measure for acute low back pain. *Journal of manipulative and physiological therapeutics* 1993; 16: 7-13.
- 13. Skargren EI, Oberg BE, Carlsson PG, Gade M. Cost and effectiveness analysis of chiropractic and physiotherapy treatment for low back and neck pain. Six-month follow-up. *Spine* 1997; 22: 2167-2177.
- 14. Innes GD, Croskerry P, Worthington J, Beveridge R, Jones D. Ketorolac versus acetaminophen-codeine in the emergency department treatment of acute low back pain. *Journal of emergency medicine* 1998; 16: 549-556.
- 15. Cherkin DC, Deyo RA, Battie M, Street J, Barlow W. A comparison of physical therapy, chiropractic manipulation, and provision of an educational booklet for the treatment of patients with low back pain. *N Engl J Med* 1998; 339: 1021-1029.
- 16. Seferlis T, Németh G, Carlsson AM, Gillström P. Conservative treatment in patients sicklisted for acute low-back pain: a prospective randomised study with 12 months' follow-up. *European spine journal* 1998; 7: 461-470.
- 17. Morton JE. Manipulation in the treatment of acute low back pain. *Journal of manual & manipulative therapy* 1999; 7: 182-189.
- 18. Veenema KR, Leahey N, Schneider S. Ketorolac versus meperidine: ED treatment of severe musculoskeletal low back pain. *American journal of emergency medicine* 2000; 18: 404-407.

- 19. Metscher B, Kübler U, Jahnel-Kracht H. Dexketoprofen-trometamol and tramadol in acute lumbago [German]. *Fortschritte der Medizin Originalien* 2001; 118: 147-151.
- 20. Palangio M, Morris E, Doyle RT, Dornseif BE, Valente TJ. Combination hydrocodone and ibuprofen versus combination oxycodone and acetaminophen in the treatment of moderate or severe acute low back pain. *Clinical therapeutics* 2002; 24: 87-99.
- 21. Hofstee DJ, Gijtenbeek JM, Hoogland PH, Houwelingen HC, Kloet A, Lötters F, et al. Westeinde sciatica trial: randomized controlled study of bed rest and physiotherapy for acute sciatica. *Journal of neurosurgery* 2002; 96(1 Suppl): 45-49.
- 22. Johnstone R, Donaghy M, Martin D. A pilot study of a cognitive-behavioural therapy approach to physiotherapy, for acute low back pain patients, who show signs of developing chronic pain. *Advances in physiotherapy* 2002; 4: 182-188.
- 23. Childs JD, Fritz JM, Flynn TW, Irrgang JJ, Johnson KK, Majkowski GR, et al. A clinical prediction rule to identify patients with low back pain most likely to benefit from spinal manipulation: a validation study. *Annals of internal medicine* 2004; 141: 920-928.
- 24. Hoiriis KT, Pfleger B, McDuffie FC, Cotsonis G, Elsangak O, Hinson R, et al. A Randomized Clinical Trial Compring Chiropractic Adjustments to Muscle Relaxants for Subacute Low Back Pain. *Journal of manipulative and physiological therapeutics* 2004; 27: 388-398.
- 25. Salvador D, Neto PD, Ferrari FP. Application of muscle energy technique in garbage collectors with acute mechanical lumbar pain [Portuguese]. *Fisioterapia e Pesquisa* 2005; 12: 20-27.
- 26. Brennan GP, Fritz JM, Hunter SJ, Thackeray A, Delitto A, Erhard RE. Identifying subgroups of patients with acute/subacute "nonspecific" low back pain: results of a randomized clinical trial. *Spine* 2006; 31: 623-631.
- 27. Santilli V, Beghi E, Finucci S. Chiropractic manipulation in the treatment of acute back pain and sciatica with disc protrusion: a randomized double-blind clinical trial of active and simulated spinal manipulations. *Spine journal* 2006; 6: 131-137.
- 28. Hancock MJ, Maher CG, Latimer J, McLachlan AJ, Cooper CW, Day RO, et al. Assessment of diclofenac or spinal manipulative therapy, or both, in addition to recommended first-line treatment for acute low back pain: a randomised controlled trial. *Lancet* 2007; 370: 1638-1643.
- 29. Lee HKH, Ting SM, Lau FL. A randomised control trial comparing the efficacy of tramadol and paracetamol against ketorolac and paracetamol in the management of musculoskeletal pain in the emergency department. Hong Kong Journal of Emergency Medicine [Internet]. 2008; 15(1):[5-11 pp.]
- 30. Lau PM, Chow DH, Pope MH. Early physiotherapy intervention in an Accident and Emergency Department reduces pain and improves satisfaction for patients with acute low back pain: a randomised trial. *Australian journal of physiotherapy* 2008; 54: 243-249.
- 31. Selkow N, Grindstaff T, Cross K, Pugh K, Hertel J, Saliba S. Short-Term Effect of Muscle Energy Technique on Pain in Individuals with Non-Specific Lumbopelvic Pain: A Pilot Study. *Journal of Manual & Manipulative Therapy* 2009; 17: 14-18.
- 32. Cleland JA, Fritz JM, Kulig K, Davenport TE, Eberhart S, Magel J, et al. Comparison of the effectiveness of three manual physical therapy techniques in a subgroup of patients with low back pain who satisfy a clinical prediction rule: a randomized clinical trial. *Spine (Phila Pa 1976)* 2009; 34: 2720-2729.
- 33. Hallegraeff JM, Hallegraeff HJ, Greef M, Winters JC, Lucas C. Manipulative therapy and clinical prediction criteria in treatment of acute nonspecific low back pain. *Perceptual and motor skills* 2009; 108: 196-208.

34. Sutlive TG, Mabry LM, Easterling EJ, Durbin JD, Hanson SL, Wainner RS, et al. Comparison of short-term response to two spinal manipulation techniques for patients with low back pain in a military beneficiary population. *Military medicine* 2009; 174: 750-756.

- 35. Jüni P, Battaglia M, Nüesch E, Hämmerle G, Eser P, Beers R, et al. A randomised controlled trial of spinal manipulative therapy in acute low back pain. *Annals of the rheumatic diseases* 2009; 68: 1420-1427.
- 36. Machado LA, Maher CG, Herbert RD, Clare H, McAuley JH. The effectiveness of the McKenzie method in addition to first-line care for acute low back pain: a randomized controlled trial. *BMC Med* 2010; 8: 10.
- 37. Lewis C, Souvlis T, Sterling M. Strain-Counterstrain therapy combined with exercise is not more effective than exercise alone on pain and disability in people with acute low back pain: a randomised trial. *Journal of physiotherapy* 2011; 57: 91-98.
- 38. Biondi D, Xiang J, Benson C, Etropolski M, Moskovitz B, Rauschkolb C. Tapentadol immediate release versus oxycodone immediate release for treatment of acute low back pain. *Pain physician* 2013; 16: E237-E246.
- 39. Goertz CM, Long CR, Hondras MA, Petri R, Delgado R, Lawrence DJ, et al. Adding chiropractic manipulative therapy to standard medical care for patients with acute low back pain: results of a pragmatic randomized comparative effectiveness study. *Spine* 2013; 38: 627-634.
- 40. Behrbalk E, Halpern P, Boszczyk BM, Parks RM, Chechik O, Rosen N, et al. Anxiolytic medication as an adjunct to morphine analgesia for acute low back pain management in the emergency department: A prospective randomized trial. *Spine* 2014; 39: 17-22.
- 41. Eken C, Serinken M, Elicabuk H, Uyanik E, Erdal M. Intravenous paracetamol versus dexketoprofen versus morphine in acute mechanical low back pain in the emergency department: A randomised double-blind controlled trial. *Emergency medicine journal* 2014; 31: 177-181.
- 42. Tanen DA, Shimada M, Danish DC, Santos FD, Makela M, Riffenburgh RH. Intravenous lidocaine for the Emergency Department treatment of acute radicular low back pain, a randomized controlled trial. *Journal of emergency medicine* 2014; 47: 119-124.

Table S3: Sensitivity analysis of factors associated with time to claim closure, entering chiropractic care and physiotherapy as time-dependant co-variates (n=1,442)

	Univariate		Multivariable	
	Hazard Ratio (99% CI)	p-value	p-value Adjusted Hazard Ratio (99%	
			CI)	-
Baseline predictors				
Age categories in years		<0.001		<0.001
15 to <25	1.00		1.00	
25 to <35	0.88 (0.65, 1.19)		0.79 (0.58, 1.08)	
35 to <45	0.78 (0.59, 1.04)		0.70 (0.52, 0.95)	
45 to <55	0.76 (0.56, 1.02)		0.67 (0.49, 0.91)	
55 to 65	0.56 (0.40, 0.80)		0.52 (0.36, 0.74)	
Gender		0.114		0.451
Females	1.09 (0.95, 1.26)		0.96 (0.82, 1.12)	
Males	1.00		1.00	
First language		0.137		0.327
English	1.00		1.00	
Other	0.83 (0.59, 1.15)		0.88 (0.63, 1.23)	
Pre-disability income in dollars				
per week				
At 30 days:				
≤480	1.00	_	1.00	_
481-694	1.15 (0.91, 1.45)	0.125	1.08 (0.85, 1.37)	0.396
695-920	1.21 (0.96, 1.54)	0.034	1.05 (0.81, 1.36)	0.614
>920	1.20 (0.94, 1.52)	0.050	1.04 (0.80, 1.35)	0.689
At 60 days:	1.20 (0.34, 1.32)	0.030	1.04 (0.80, 1.33)	0.003
480 ≤480	1.00		1.00	_
481-694	1.09 (0.89, 1.34)	0.273	1.03 (0.83, 1.27)	0.710
695-920	1.13 (0.92, 1.39)	0.273	0.97, (0.77, 1.22)	0.710
>920	1.08 (0.88, 1.33)	0.137	0.94 (0.74, 1.19)	0.746
	1.00 (0.00, 1.55)	0.320	0.94 (0.74, 1.19)	0.477
<b>At 180 days:</b> ≤480	1.00	_	1.00	
				0.120
481-694	0.89 (0.68, 1.17)	0.267 0.115	0.85 (0.65, 1.12) 0.71 (0.52, 0.96)	0.130 0.003
695-920 >920	0.84 (0.63, 1.12)	0.115	,	
	0.72 (0.54, 0.97)	0.004	0.61 (0.45, 0.84)	<0.001
At 365 days:	1.00		1.00	
≤480 481 CO4	1.00	0.064	1.00	0.051
481-694	0.65 (0.36, 1.18)	0.064	0.63 (0.35, 1.16)	0.051
695-920	0.53 (0.28, 1.02)	0.012	0.44 (0.23, 0.84)	0.001
>920	0.39 (0.20, 0.75)	<0.001	0.32 (0.16, 0.63)	<0.001
Opioid prescription	0.52 (0.40, 0.00)	<0.001	0.50 (0.50, 0.00)	<0.001
Yes	0.62 (0.48, 0.80)		0.69 (0.53, 0.89)	
No	1.00		1.00	
Prior claims		0.306		0.661
Yes	1.07 (0.91, 1.26)		1.03 (0.87, 1.22)	
No	1.00		1.00	
Union membership		<0.001		0.014
Yes	1.29 (1.11, 1.50)		1.14 (0.96, 1.36)	
No	1.00		1.00	
Missing	1.34 (1.07, 1.68)		1.27 (1.01, 1.60)	
Employer RTW program		<0.001		<0.001
Yes	1.73 (1.43, 2.09)		1.77 (1.45, 2.18)	

No	1.00		1.00	
Missing	1.17 (0.87, 1.58)		1.18 (0.86, 1.60)	
Doubt Work-relatedness		0.174		0.119
Yes	0.94 (0.76, 1.16)		0.87 (0.70, 1.08)	
No	1.00		1.00	
Missing	1.13 (0.92, 1.39)		1.08 (0.87, 1.33)	
Time-dependent predictors*				
Chiropractic care received	1.11 (0.95, 1.30)	0.096	1.08 (0.91, 1.29)	0.268
after the accident prior to				
claim closure				
Physiotherapy after the	0.96 (0.83, 1.10)	0.420	0.99 (0.85, 1.16)	0.913
accident prior to claim closure				

HR>1 indicates faster claim closure; RTW = return to work

<sup>\*</sup> The time-dependent predictors are "turned on" once the claimant has received their first service after their accident.

2

4

5

6 7 8

9

10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26

27 28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52 53

54

55

56

57

58

59 60

# Table S4: Search strategy for observational studies exploring the association of early opioid, physiotherapy, or chiropractic care with Workers' Compensation claim duration, for cases of acute low back pain

#### MEDLINE (OvidSP)

- 1 exp Whiplash Injuries/
- 2 exp Soft Tissue Injuries/
- 3 repetitive strain injur\$.mp.
- 4 carpal tunnel syndrome.mp.
- 5 exp Cumulative Trauma Disorders/
- 6 exp Back pain/ or exp pain/ or chronic pain.tw.
- 7 exp Anxiety/
- 8 exp Depression/
- 9 exp Neck Pain/
- 10 exp Depressive Disorder/
- 11 exp Back Injuries/
- 12 injured worker\$.mp.
- 13 musculoskeletal injur\$.mp.
- 14 or/1-13
- 15 exp "Wounds and Injuries"/
- 16 Musculoskeletal System/ or Musculoskeletal Diseases/
- 17 15 and 16
- 18 14 or 17
- 19 exp insurance claim reporting/ or exp "insurance claim review"/ or exp insurance, disability/ or insurance, liability/
- 20 Insurance, Accident/
- 21 ((worker\$ or workman\$ or workmen&) adj compensation).mp. or exp Workers' compensation/
- 22 claim.mp.
- 23 claimant.mp.
- 24 or/19-23
- 25 prognosis.mp. or exp Prognosis/
- 26 Time/ or exp Time Factors/
- 27 exp "Recovery of Function"/
- 28 "Severity of Illness Index"/
- 29 exp Trauma Severity Indices/
- 30 (recovery or prognostic).mp.
- 31 or/25-30
- 32 18 and 24 and 31
- 33 exp Disability Evaluation/
- 34 24 and 31 and 33
- 35 exp Occupational Diseases/ or exp Accidents, Occupational/ or (occupational injur: or occupational accident:).mp.
- 36 24 and 31 and 35
- 37 exp Accidents, Traffic/
- 38 24 and 31 and 37
- 39 "Compensation and Redress"/
- 40 18 and 31 and 39

- 41 exp Work Capacity Evaluation/ or exp workload/ or workload.mp.
- 42 (18 or 35) and 24 and 41
- 43 32 or 34 or 36 or 38 or 40 or 42
- 44 18 or 33

2

4

5

6 7

8

9

10

11 12 13

14 15

16

17

18

19

20 21

22

23

24

25

26 27

28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52

53 54

55

56

57

58

59 60

- 45 31 or 41
- 46 35 and 44 and 45
- 47 46 or 43

#### EMBASE (OvidSP)

- 1 whiplash injur\$.mp. or exp whiplash injury/
- 2 exp soft tissue injury/
- 3 soft tissue injur\$.mp.
- 4 repetitive strain injur\$.mp.
- 5 carpal tunnel syndrome.mp. or exp carpal tunnel syndrome/
- 6 exp cumulative trauma disorder/
- 7 back pain.mp. or exp backache/
- 8 backpain.mp.
- 9 chronic pain.mp. or exp chronic pain/
- 10 exp pain/
- 11 anxiety/
- 12 exp depression/
- 13 neck pain.mp. or exp neck pain/
- 14 back injur\$.mp.
- 15 low back injury/
- 16 injured worker\$.mp.
- 17 musculoskeletal injury/
- 18 occupational injuries.mp. or exp occupational accident/
- 19 occupational accidents.mp.
- 20 occupational diseases.mp. or exp occupational disease/
- 21 or/1-20
- 22 insurance/ or exp compensation/ or exp workman compensation/ or exp health insurance/ or
- exp "health plan employer data and information set"/
- 23 accident insurance.mp.
- 24 exp workman compensation/
- 25 ((worker\$ or workman\$ or workmen&) adj compensation).mp.
- 26 (claim or claimant).mp.
- 27 or/22-26
- 28 prognosis.mp. or prognosis/
- 29 exp time/
- 30 recovery of function.mp. or convalescence/
- 31 disease severity/
- 32 exp injury scale/
- 33 (recovery or prognostic).mp.
- 34 workload.mp. or exp workload/
- 35 exp work capacity/
- 36 exp work resumption/ or return to work.mp.
- 37 or/28-36

2

4

5

6 7

8

9 10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26 27

28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52 53

54

55

56

57

58

59 60 39 or/29-38

40 20 and 28 and 39

```
38 21 and 27 and 37
39 or/1-17
40 or/18-20
41 37 and 39 and 40
42 38 or 41
PsycInfo (OvidSP)
1
    exp Whiplash/
2
    whiplash injur:.mp.
3
    soft tissue injur$.mp.
4
    cumulative trauma disorder$.mp.
5
    repetitive strain injur$.mp.
6
    carpal tunnel syndrome.mp.
7
    back pain.mp. or exp Back Pain/
8
    (backpain or backache).mp.
9
    chronic pain.mp. or exp Chronic Pain/
10 exp Musculoskeletal Disorders/ or exp Fibromyalgia/ or fibromyalgia.mp.
11
    exp Anxiety/
12 exp "Depression (Emotion)"/ or exp Major Depression/
13 neck pain.mp.
14 back injur:.mp.
15 musculoskeletal injur$.mp.
16 exp Industrial Accidents/
17 exp Occupational Safety/
18 (occupational injur: or occupational accident:).mp.
19 exp Work Related Illnesses/
20 or/1-19
21 exp Workers' Compensation Insurance/
22 exp Employee Health Insurance/
23 exp Insurance/
24 disability insurance.mp.
25 (claim or claimant).mp.
26 ((worker: or workman: or workmen:) adj compensation).mp.
27 accident insurance.mp.
28 or/21-27
29 prognosis.mp. or exp Prognosis/
30 exp Time/
31 time factors.mp.
32 exp "Recovery (Disorders)"/
33 recovery of function.mp.
34 exp "Severity (Disorders)"/ or severity of illness.mp.
35 (recovery or prognostic).mp.
36 exp Work Load/ or workload.mp. or exp Job Performance/
37 exp Vocational Evaluation/ or exp Disability Evaluation/ or work capacity evaluation.mp. or
exp Reemployment/
38 work resumption.mp.
```

2

4

5

6

7 8

9 10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26

27 28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46

47 48

49

50

51

52

53 54

55

56

57

58

```
41 16 or 17 or 18 or 19
42 or/1-15
43 39 and 41 and 42
44 28 and 41
45 40 or 43 or 44
CINAHL (Ebsco)
49
    S46 or S48
S48 S22 and S45 and S47
S47 S23 or S24 or S25 or S26
S46 S27 and S33 and S45
S45 S34 or S35 or S36 or S37 or S38 or S39 or S40 or S41 or S42 or S43 or S44
S44 (MH "Disability Evaluation+")
S43 "work resumption"
S42 "return to work" OR (MH "Job Re-Entry")
S41 (MH "Work Capacity Evaluation")
S40 (MH "Workload Measurement") OR (MH "Workload") OR "workload"
S39 (recovery or prognostic)
S38 (MH "Severity of Illness") OR (MH "Severity of Illness Indices+")
S37 (MH "Recovery")
S36 "recovery of function"
S35 (MH "Time+") OR (MH "Time Factors")
S34 (MH "Prognosis+") OR "prognosis"
S33 S28 or S29 or S30 or S31 or S32
S32 "accident insurance"
S31 (worker* N2 compensation) OR (workman* N2 compensation) OR (workmen* N2
compensation)
S30 (Claim or claimant)
S29 (MH "Insurance") OR (MH "Insurance, Disability+")
S28 (MH "Worker's Compensation")
S27 S22 or S23 or S24 or S25 or S26
S26 "occupational accident" OR (MH "Accidents, Occupational+")
S25 "occupational inju*"
S24 (MH "Occupational-Related Injuries")
S23 (MH "Occupational Diseases+")
S22 S18 or S21
S21 S19 and S20
S20 (MH "Wounds and Injuries+") OR (MH "Occupational-Related Injuries")
S19 (MH "Musculoskeletal Diseases+") or (MH "Musculoskeletal System+")
S18 S1 or S2 or S3 or S4 or S5 or S6 or S7 or S8 or S9 or S10 or S11 or S12 or S13 or S14 or
S15 or S16 or S17
S17 "musculoskeletal injur*"
S16 "back injur*"
S15 (MH "Back Injuries+")
S14 (MH "Neck Pain") OR "neck pain"
S13 (MH "Depression+")
S12 (MH "Anxiety+")
S11 (MH "Pain+")
```

- S10 (MH "Fibromyalgia") OR "fibromyalgia"
- S9 (MH "Chronic Pain") OR "chronic pain"
- S8 (backpain or backache)
- S7 (MH "Back Pain") OR "back pain"
- S6 (MH "Carpal Tunnel Syndrome") OR "carpal tunnel syndrome"
- S5 "repetitive strain injur\*"
- S4 (MH "Cumulative Trauma Disorders+")
- S3 "soft tissue injur\*"
- S2 (MH "Soft Tissue Injuries")
- S1 "whiplash injur\*" OR (MH "Whiplash Injuries")

Figure S1: Flow diagram of the literature search process for randomized controlled trials assessing the effect of opioids, physiotherapy, or chiropractic care for acute low back pain.

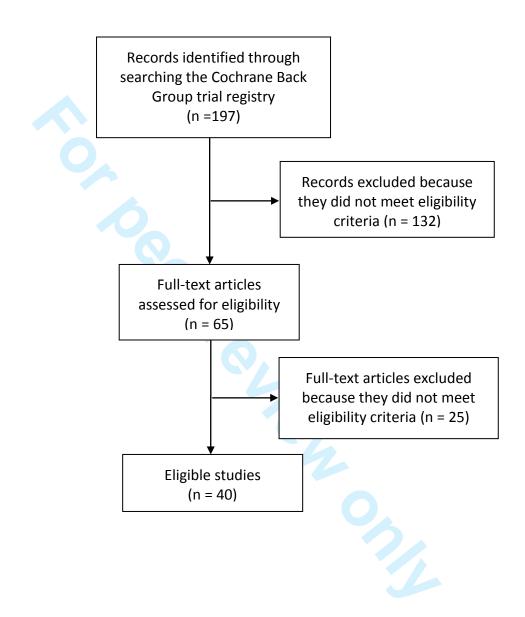


Figure S2: Kaplan-Meier curve for early reimbursement for Opioid prescription



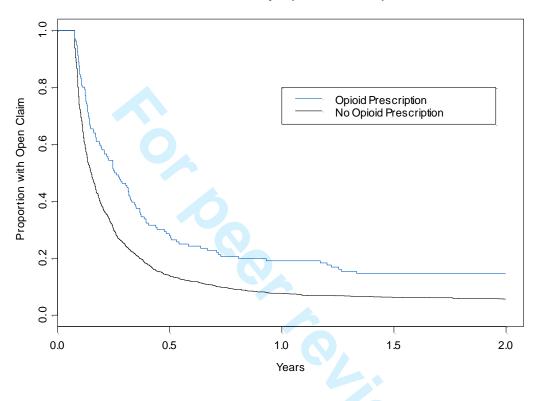


Figure S3: Kaplan-Meier curve for early reimbursement for physiotherapy

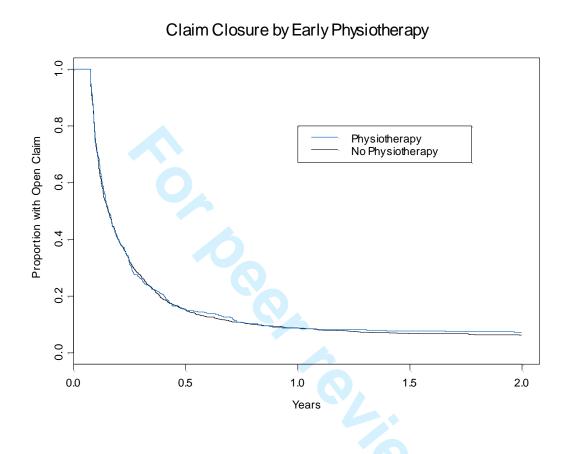
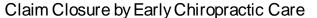


Figure S4: Kaplan-Meier curve for early reimbursement for chiropractic care



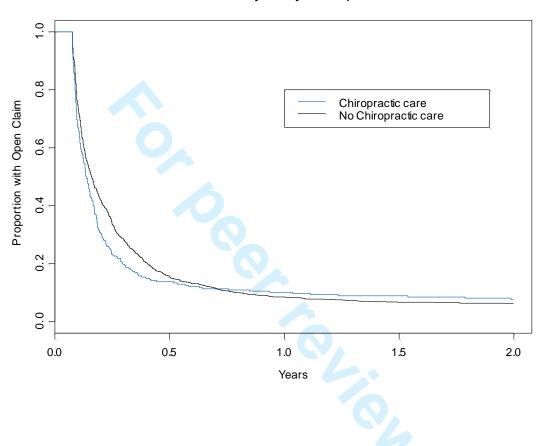
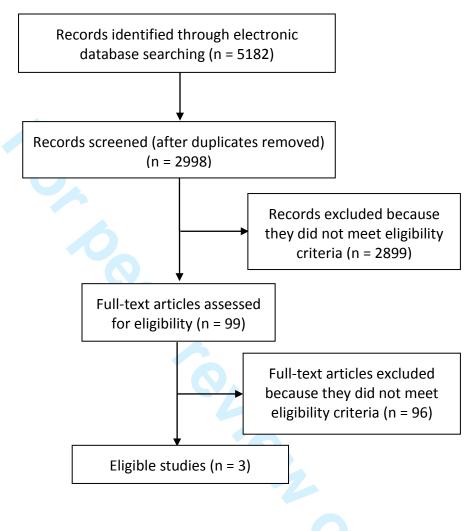


Figure S5: Flow diagram of the literature search process for observational studies assessing the effect of early opioids, physiotherapy, or chiropractic care for acute low back pain.



STROBE Statement—Checklist of items that should be included in reports of *cohort studies* 

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the
		abstract (in title on title page)
		(b) Provide in the abstract an informative and balanced summary of what
		was done and what was found (in Abstract)
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being
		reported (page 1 & 2 in Manuscript)
Objectives	3	State specific objectives, including any prespecified hypotheses (Appendix
J		A)
Methods		,
Study design	4	Present key elements of study design early in the paper (In Abstract, and
3 8		page 3-7 in Manuscript)
Setting	5	Describe the setting, locations, and relevant dates, including periods of
		recruitment, exposure, follow-up, and data collection (page 3 in
		Manuscript)
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of
1		participants. Describe methods of follow-up (pages 3 & 4 in Manuscript)
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed (not applicable)
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders,
variables	,	and effect modifiers. Give diagnostic criteria, if applicable ( <b>Appendix A</b>
		and pages 3 & 4 in Manuscript)
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of
Data sources/ measurement	O	assessment (measurement). Describe comparability of assessment methods
		if there is more than one group (pages 3-5 in Manuscript)
Bias	9	Describe any efforts to address potential sources of bias (pages 4-7 in
Dias	9	Manuscript)
Study size	10	Explain how the study size was arrived at (page 3 & 4 in Manuscript)
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If
Quantitudi ( variation )		applicable, describe which groupings were chosen and why (pages 5 & 6 in
		Manuscript)
Statistical methods	12	(a) Describe all statistical methods, including those used to control for
Statistical inclinate	12	confounding (pages 3-7 in Manuscript)
		(b) Describe any methods used to examine subgroups and interactions (page
		6 in Manuscript)
		(c) Explain how missing data were addressed (page 5 in Manuscript)
		(d) If applicable, explain how loss to follow-up was addressed ( <b>not</b>
		applicable)
		(e) Describe any sensitivity analyses (not applicable)
Doculto		( <u></u>
Results Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers
		potentially eligible, examined for eligibility, confirmed eligible, included in
		the study, completing follow-up, and analysed ( <b>Table 1 and page 8 in</b>
		Manuscript)
		(b) Give reasons for non-participation at each stage ( <b>not applicable</b> )
		(1) 21.1 reasons for non-participation at each stage (not applicable)

Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders ( <b>Table 1</b> )
		(b) Indicate number of participants with missing data for each variable of interest ( <b>Table 1</b> )
		(c) Summarise follow-up time (eg, average and total amount) (Figure 1)
Outcome data	15*	Report numbers of outcome events or summary measures over time (page 8 in Manuscript)
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included ( <b>Table 2</b> )
		(b) Report category boundaries when continuous variables were categorized (Table 2)
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period ( <b>not relevant</b> )
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses (page 8 in Manuscript)
Discussion		, , , , , , , , , , , , , , , , , , , ,
Key results	18	Summarise key results with reference to study objectives (page 10 in Manuscript)
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias (pages 10 & 11 in Manuscript)
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence (page 13 in Manuscript)
Generalisability	21	Discuss the generalisability (external validity) of the study results (page 11 in Manuscript)
Other information		
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based (page 14 in Manuscript)

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at http://www.strobe-statement.org.

# **BMJ Open**

The association of worker characteristics and early reimbursement for physical therapy, chiropractic and opioid prescriptions with Workers' Compensation claim duration, for cases of acute low back pain: An observational cohort study

Journal:	BMJ Open
Manuscript ID:	bmjopen-2015-007836.R2
Article Type:	Research
Date Submitted by the Author:	13-Jul-2015
Complete List of Authors:	Busse, Jason; McMaster University, Anesthesia; McMaster University, Clinical Epidemiology & Biostatistics Ebrahim, Shanil; Mcmaster University, Clinical Epidemiology and Biostatistics Heels-Ansdell, Diane; Mcmaster University, Clinical Epidemiology and Biostatistics Wang, Li; McMaster University, Anesthesia Couban, Rachel; McMaster University, Anesthesia Walter, Stephen; Mcmaster University, Clinical Epidemiology and Biostatistics
<b>Primary Subject Heading</b> :	Rehabilitation medicine
Secondary Subject Heading:	Complementary medicine, Evidence based practice
Keywords:	COMPLEMENTARY MEDICINE, Pain management < ANAESTHETICS, OCCUPATIONAL & INDUSTRIAL MEDICINE

SCHOLARONE™ Manuscripts

Predicting Workers' Compensation claim duration

The association of worker characteristics and early reimbursement for physical therapy, chiropractic and opioid prescription with Workers' Compensation claim duration, for cases of acute low back pain: An observational cohort study

Jason W. Busse,<sup>1,2,3\*</sup> Shanil Ebrahim,<sup>2,3,4,5</sup> Diane Heels-Ansdell,<sup>3</sup> Li Wang,<sup>2</sup> Rachel Couban,<sup>2</sup> Stephen D. Walter<sup>3</sup>

\*Corresponding Author: email: bussejw@mcmaster.ca

### **Abstract**

**Objective:** To assess the association between early reimbursement for physiotherapy, chiropractic, and opioid prescription for acute low back pain (LBP) with disability claim duration.

**Design:** Observational cohort study.

**Setting and participants:** From a random sample of 6,665 claims for acute, uncomplicated LBP approved by the Ontario Workplace Safety and Insurance Board (WSIB) in 2005, we analyzed 1,442 that remained on full benefits at four weeks after claim approval.

**Primary outcome measure:** Our primary outcome was WSIB claim duration.

Results: We had complete data for all but 3 variables, which had <15% missing data, and we included missing data as a category for these factors. Our time-to-event analysis was adjusted for demographic, workplace, and treatment factors, but not injury severity although we attempted to include a sample with very similar, less severe injuries. Regarding significant factors and treatment variables in our adjusted analysis, older age (e.g. hazard ratio [HR] for age ≥55 versus <25 = 0.52; 99% confidence interval [CI] = 0.36, 0.74) and WSIB-reimbursement for opioid prescription in the first 4-weeks of a claim (HR = 0.68; 99% CI = 0.53, 0.88) were associated with longer claim duration. Higher pre-disability income was associated with longer claim duration, but only among persistent claims (e.g. HR for active claims at 1-year with a pre-disability income >\$920/week versus ≤\$480 = 0.34; 99% CI = 0.17, 0.68). Missing data for union membership (HR = 1.27; 99% CI = 1.01, 1.59), and working for an employer with a return-to-work program was associated with fewer days on claim (HR = 1.78; 99% CI = 1.45, 2.18). Neither reimbursement for physiotherapy (HR = 1.01; 99% CI = 0.86, 1.19) or chiropractic care (HR for

Predicting Workers' Compensation claim duration

active claims at 60 days = 1.15; 99% CI = 0.94, 1.41) within the first 4-weeks were associated with claim duration. Our meta-analysis of 3 studies (n=51,069 workers) confirmed a strong association between early opioid use and prolonged claim duration (HR = 0.57, 95%CI = 0.48 to 0.69; moderate certainty evidence).

Conclusions: Our analysis found that early WSIB reimbursement for physiotherapy or chiropractic care, in claimants fully off work greater than 4 weeks, was not associated with claim duration, and that early reimbursement for opioids predicted prolonged claim duration. Well-designed randomized controlled trials are needed to verify our findings and establish causality between these variables and claim duration.

**Key Words:** Low back pain; disability; compensation; Workers Compensation Board; physical therapy; chiropractic; opioid; survival analysis

Predicting Workers' Compensation claim duration

#### **Strengths and Limitations**

- A priori creation of our regression model and the anticipated direction of included independent variables, and assessment of the proportional hazards assumption for all independent variables, provide greater confidence in our findings.
- The reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are
  uncertain, and despite our adjustments for potential confounders (but not injury
  severity) it remains possible that LBP claimants who chose to receive these healthcare
  interventions were prognostically different than those claimants who did not.
- A number of variables that may be important to consider were unavailable (e.g., patient
  expectations regarding recovery), and chiropractic and physiotherapy are professions,
  not modalities, and details of the treatment provided could not be obtained for our
  analysis.
- Our primary outcome, time to claim closure, is a surrogate for patient-important outcomes such as functional restoration or return to work.

Predicting Workers' Compensation claim duration

# Introduction

lived with disability.3

Back pain is a common problem among working adults in North America, with a lifetime prevalence of 63% and a point prevalence of 21%. After the common cold, low back pain (LBP) is the most frequent cause of lost time from work.<sup>2</sup> Globally, LBP is the primary cause of years

In Canada, annual medical expenditures for LBP are estimated to be between \$6 and \$12 billion, with additional costs associated with loss in worker productivity from time off work and associated disability payments. 4 Canadian workers who are disabled secondarily to a workrelated LBP injury are typically eligible for wage replacement benefits through their provincial Workers' Compensation Board.

In 2013, the Ontario Workplace Safety and Insurance Board (WSIB) approved approximately 232,000 claims that were associated with \$2,761 million in payments, and 18% of all allowed lost-time claims were for low back injuries. 5, 6 The WSIB's liability for disability claims greatly exceeds their assets, and as of March 31, 2013 the WSIB's unfunded liabilities were \$12.4 billion<sup>7</sup> – more than double their unfunded liability of \$5.9 billion in 2006.<sup>8</sup> Unfunded liability is the amount by which future payment obligations exceed the present value of funds available to pay them. To reduce their unfunded liability, the WSIB has become more aggressive about denying claims, decreasing disability benefits, and increasing employee premiums; 9, 10 however, these measures do not address optimal management of disability claims.

Interventions that are commonly reimbursed by WSIB for LBP claims include physiotherapy, chiropractic care and opioids; however, there is limited evidence about their effectiveness. Our systematic review of the Cochrane Back Review Group trial registry found no randomized controlled trials of these interventions focused on workers fully disabled by acute LBP and receiving benefits for lost-time claims (Tables S1 & S2, and Figure S1, in File S1). We also found that many trials of LBP patients use receipt of disability benefits as an exclusion criterion, likely because of concerns that secondary gain (e.g., receipt of financial compensation conditional on disability) will reduce the impact of study interventions. Henshke and colleagues followed a cohort of 973 consecutive primary care patients with non-specific, acute LBP recruited from the clinics of 170 general practitioners, physiotherapists, and chiropractors for 1year. They found that, in an analysis adjusted for age, gender, injury severity and psychological factors, receipt of disability benefits was strongly associated with delayed recovery (hazard ratio [HR]=0.59; 95%CI = 0.47 to 0.74). 11 As compensated acute LBP has a worse prognosis than uncompensated acute LBP, the results of trials that do not enroll patients receiving disability benefits cannot be confidently generalized to patients who are receiving compensation.

Using administrative data from the Ontario WSIB, we evaluated the association between receiving early reimbursement for physiotherapy, chiropractic care, or prescription for opioids for uncomplicated, acute LBP and disability claim duration. On the basis of prior observational studies, <sup>12-16</sup> we hypothesized that early reimbursement for opioids would be associated with delayed recovery, and early reimbursement for physiotherapy or chiropractic care would be associated with faster recovery. We reported our findings in concordance with the STROBE <sup>17</sup> and TRIPOD <sup>18</sup> statements.

# **Methods**

## Patient characteristics and eligibility criteria

Using WSIB administrative data, we identified an inception cohort of workers with uncomplicated, acute low back injuries (i.e., strain or sprain) who were fully disabled from working and receiving wage replacement benefits from the Ontario WSIB. We excluded workers if they were approved for no-lost-time claims. The prognosis for chronic LBP (duration >12 weeks) is different than for acute LBP, <sup>19</sup> and we excluded all claims in which the number of days between accident date and registration date of the claim was greater than 30 days. Most LBP claims resolve within the 1<sup>st</sup> month, <sup>20</sup> and Workers' Compensation Boards are primarily interested in factors that predict claim resolution among claimants who remain disabled after this time. We therefore excluded claims that ended before 4 weeks.

Between January 1 and June 30, 2005, the Ontario WSIB approved 18,974 lost-time claims for an uncomplicated, acute low back injury. Using the WSIB's administrative database, we acquired a random sample of 6,665 injured workers from this population; 1,442 unique workers remained on full benefits at four weeks and provided data for our analysis. If a worker had more than 1 claim for acute LBP, their first claim was used. The WSIB database recorded benefit status for two years after the first day of injury. Patient information was anonymized and de-identified prior to analysis. The Health Sciences Research Ethics Board of the University of Toronto approved our study protocol.

### **Administrative Variables**

Our primary outcome was time to claim closure, defined as the duration in days from disability claim approval until the claim was closed. The WSIB database also contained demographic, administrative, and clinical information, which were acquired from forms completed by the worker, their employer, and their primary health-care provider. The employer form (Employer's Report of Injury/Disease Form; Form 7), which is used to indicate whether there is doubt regarding the work-relatedness of an employee's back injury, is mandatory and must be submitted within three days of a work-related injury. The form asks employer's "Do you have any reason to doubt the injury/disease is work-related?" and they can indicate either "no" or "yes".

The worker may elect to fill out a form (Worker's Report of Injury/Disease; Form 6) if they have expenses related to their injury. The healthcare provider can elect to complete a form (Health Professional's Report; Form 8) to support their patient's claim that their injury is work-related, which is a pre-requisite for wage replacement benefits through the WSIB. Healthcare providers are asked to complete and submit a Functional Abilities Form for Planning Early and Safe Return to Work for each claim, and the WSIB provides compensation as incentive.

In order to increase confidence in our findings, we defined our regression model before conducting any analyses. Guided by the results from our ongoing systematic review of observational studies evaluating predictors of recovery in patients receiving disability benefits, <sup>21</sup> feedback from administrators at WSIB, and content experts within our research team, we selected, *a priori*, 11 variables from the WSIB database that we judged may be associated with claim closure; we also specified the direction of anticipated effects on claim

duration (Table 1): age, gender, native language, pre-disability income, prior disability claim, union membership, working for an employer with a return to work (RTW) program, employer's doubt that the injury was work-related, and early (1<sup>st</sup> month) receipt of reimbursement for opioids, physiotherapy, or chiropractic care (Table 1). In 2005, the Ontario WSIB did not capture any measure of LBP injury severity on Forms 6, 7 or 8, and as such we were unable to adjust for this variable. Injured workers may attend a healthcare provider for assessment purposes and we required ≥3 reimbursed visits for physiotherapy or chiropractic, within the first 28-days of claim, in order to qualify as reimbursement for treatment.

We hypothesized that workers represented by a union would resolve their claim faster, as we felt they would have more support for re-engagement with competitive employment (e.g. graduated work hours) versus workers who were not members of a union. We also hypothesized that claims due to injuries that employers reported were work-related would resolve faster than injuries in which the employer doubted that the employee was injured at work, as we felt this may be a surrogate for the influence of non-medical factors (e.g. secondary gain). Based on the findings of a recent systematic review that found RTW coordination was associated with faster RTW for disabled employees, <sup>22</sup> we hypothesized that claimants employed by companies with formal RTW programs would resolve their claim faster.

### **Data Extraction**

The Ontario WSIB's database consists of scanned paper documents, and data must therefore be extracted manually for analysis. Two reviewers extracted data, independently and in duplicate, from the first 100 eligible claims into an Access database (Microsoft Access, Filemaker). In order

to minimize data entry mistakes, we developed data entry forms that included range checks and missing value alerts. The PROC COMPARE procedure in SAS version 9.2 (SAS Institute Inc., Cary, NC) revealed 98% agreement for the initial 100 claims; therefore, to increase feasibility, only a single abstractor completed data entry for the remaining claims used in our analyses.

We screened all data to identify outliers, inconsistencies and missing data by calculating summary statistics, and explored distributions graphically. We worked with WSIB representatives to correct identified outliers and inconsistencies. If inconsistencies could not be corrected, we treated them as missing data. Some WSIB forms are voluntary, and so we included "missing data" as a discrete category for independent variables when applicable.

## Statistical analysis

We generated frequencies for all collected data. We reported the mean and standard deviation (SD) of continuous variables, and the number of occurrences represented as proportions for categorical variables. Age was negatively skewed and pre-disability gross income was positively skewed and were therefore entered as categorical variables into our regression model; by decade for age, starting at age 15 and ending at age 65, and by quartiles for pre-disability income.

We performed a time-to-event analysis using a Cox proportional hazards regression model to assess the association between time to claim closure and all 11 independent variables described in Table 1. To avoid overfitting our models, we required at least 10 observations per variable term for our Cox regression model, for a total of 190 disabled workers. We set a threshold of at least 50 observations per category for each independent factor in our regression

model to provide some reassurance that each variable had sufficient discriminant power to detect an association with claim duration, if such an association existed.

For claims that were unresolved when the data was extracted, we used 2-years after claim approval as a censoring point. In order to be more stringent and minimize the likelihood of spurious findings, an independent variable was considered statistically significant if it had a p-value of ≤0.01 in our adjusted model. We calculated adjusted hazard ratios (HRs) for our time-to-event analyses, their associated 99% confidence intervals (Cls), and the associated p-values. We assessed each independent variable in our model to ensure that the proportional hazards assumption was met by entering each variable in the model separately and calculating its interaction with time. We considered a p-value of ≤0.05 for the interaction term as significant. We reported the HRs for independent variables that violated the proportional hazards assumption at 60-days, 6-months, and 1-year. We conducted a sensitivity analysis to investigate the impact of entering receipt of WSIB-reimbursed chiropractic care or physiotherapy as time-dependent covariates in a Cox proportional hazards regression model. This approach accounts for when treatments were initiated during the course of the disability claim.

# Pooling data from similar studies

When possible, we pooled the association between early opioid, physiotherapy, or chiropractic care and claim duration in our sample with similar data from observational studies identified through a systematic review (search strategy, Table S4 in File S1). We considered studies to be similar if they enrolled Workers' Compensation patients who were completely disabled from

working secondary to acute, uncomplicated LBP and, explored the association of early treatment with opioids, physiotherapy or chiropractic care with claim duration. Using standardized, pilot-tested forms, 2 reviewers screened, independently and in duplicate, titles and abstracts of identified citations and then full texts of potentially eligible studies. The same reviewers extracted patient characteristics, methodology, and measures of association between early use of opioids, physiotherapy or chiropractic care and disability claim duration from eligible articles.

We used the following criteria to gauge risk of bias: (1) representativeness of the study population (low risk of bias when using random sampling or consecutive sampling, high risk of bias when the source of study population was not reported or acquired through convenience sampling); (2) validity of outcome assessment (low risk of bias when claim duration was acquired directly from the benefits administrator); (3) proportion of lost to follow-up (high risk of bias if >20%); and (4) whether or not predictive models were appropriately adjusted (low risk of bias if adjusted for age, gender, and injury severity).

When possible, we pooled measures of association between early opioid, physiotherapy, or chiropractic care and claim duration, and presented the pooled estimate as a HR and the associated 95%CI. When necessary, we converted odds ratios (ORs) to a relative risk (RR), then to a HR, using the following formula: <sup>24</sup>

$$RR = OR/(1-P_0 + P_0 \times OR)$$

$$HR = (In(1-RR \times P_0))/(In(1-P_0))$$

where  $P_0$  is the proportion of patients in the control group who had an event by the follow-up time.

Predicting Workers' Compensation claim duration

We used random-effects meta-analyses, which are usually conservative in that they take both within-study and between-study variability into account. We examined heterogeneity using both a chi-squared test and the I<sup>2</sup> statistic, the latter being the percentage of the total variation in outcomes that is associated with between-study variability (i.e., true differences between studies rather than with sampling error (chance)). Heterogeneity of 0% to 40% was considered 'might not be important', 30% to 60% to be 'moderate heterogeneity', 50% to 90% to be 'substantial heterogeneity', and 75% to 100% to be 'considerable heterogeneity'. The Cochrane Collaboration has proposed overlapping categories, to convey that there are no strict cut-offs for interpreting heterogeneity, and that this decision will depend on the magnitude and direction of effects, as well as the strength of evidence for heterogeneity.

We used the GRADE approach to summarize the certainty of evidence for the effect of early opioid use on claim duration as high, moderate, low, or very low. <sup>28</sup> Using GRADE, observational studies begin as moderate certainty but can be rated down due to: (1) risk of bias; (2) inconsistency; (3) indirectness; (4) imprecision; or (5) publication bias. GRADE suggests considering rating up certainty of evidence one level when methodologically rigorous observational studies show at least a two-fold reduction or increase in risk, and rating up two levels for at least a five-fold reduction or increase in risk. <sup>29</sup> We assessed publication bias by visually observing asymmetry of funnel plots, but only if there were ≥10 studies eligible for meta-analysis. We performed all statistical analyses using SAS version 9.2 (SAS Institute Inc., Cary, NC, USA). All hypothesis tests were 2-tailed and p≤ 0.05 was considered statistically significant.

We estimated the cumulative proportion of claims closed in our WSIB dataset at 90-days for disabled workers that did, and did not, receive early opioids by using the following formula:

$$P_1 = 1 - (1 - P_0)^{HR}$$

where  $P_1$  is the cumulative proportion of claims closed by 90-days in the early opioid group,  $P_0$  is the cumulative proportion of claims closed by 90-days in the group that did not receive early opioids, and HR is the pooled estimate of the HR from our meta-analysis.

Predicting Workers' Compensation claim duration

# Results

Table 2 presents the baseline characteristics for the 1,442 disabled workers included in our analysis. The Ontario WSIB reimbursed 786 (55%) claimants for physiotherapy and 391 (27%) for chiropractic care. In the first 4 weeks of their claim, 27% (n=388) were reimbursed for ≥3 physiotherapy treatments, 17% (n=247) were reimbursed for ≥3 chiropractic treatments, and 9% of claimants (n=136) were reimbursed for an opioid prescription. Figure 1 presents the Kaplan-Meier curve for time to claim closure for LBP claimants. Most workers (67%, n=966) had resolved their claim by 90 days, 84% (n=1211) by 180 days, and 91% by 1 year (n=1312); 1348 (93.5%) claims were closed prior to 2 years and 94 (6.5%) were censored.

#### **Time-to-Event Analysis**

Our adjusted regression analysis showed that older age (e.g., HR for age ≥55 versus <25 = 0.52; 99% CI = 0.36, 0.74) and opioid prescription reimbursed by the Ontario WSIB in the first 4weeks of claim (HR = 0.68; 99% CI = 0.53, 0.88) were associated with longer claim duration. The HRs for pre-disability income and receiving reimbursement for early chiropractic care with claim duration were not proportional over time (p=0.001 and 0.031, respectively), and older claims showed a significant association of greater pre-disability income with longer claim duration (e.g. HR for active claims at 1-year with a pre-disability income >\$920/week versus ≤\$480/week = 0.34; 99% CI = 0.17, 0.68). Working for an employer that had a RTW program (HR = 1.78; 99% CI = 1.45, 2.18), and missing data regarding union membership (HR = 1.27, 99%CI = 1.01, 1.59) were associated with shorter claim duration. Contrary to our predictions, neither early receipt

of WSIB-reimbursement for physiotherapy (HR = 1.01; 99% CI = 0.86, 1.19) nor chiropractic care (e.g. HR for active claims at 60-days = 1.15; 99% CI = 0.94, 1.41) were associated with claim duration (Table 3). We found no important differences using alternative analytic methods (Table S3 in File S1). Figures 2-4 present the Kaplan-Meier curves for time to claim closure for acute LBP claimants who received reimbursement for early opioid prescription, physiotherapy and chiropractic care.

#### **Systematic Review**

Our systematic review of observational studies identified 2998 unique records, of which we retrieved 99 in full text; three were eligible for our review and explored early opioid use (Figure S2 in File S1). All 3 observational studies that explored early use of opioids for Workers' Compensation claims due to uncomplicated, acute LBP reported a significant association with prolonged claim duration, and two studies adjusted for injury severity in their regression models (Table 4). When our results were pooled with the 2 studies that reported measures of association in relative units,  $^{14, 15}$  resulting in a total of 51,069 participants, the association between early opioid use and prolonged claim duration was consistent with our findings (adjusted HR = 0.57, 95% CI = 0.48 to 0.69), heterogeneity test p=0.02,  $I^2$ =75%; moderate certainty evidence) (Figure 5). Applying this effect to our WSIB dataset means that, at 90-days, 69% of workers without reimbursement for early opioids had resolved their disability claim versus 49% of workers who received reimbursement for early opioids.

The pooled effect was associated with substantial heterogeneity; however, statistical tests of heterogeneity can be misleading when sample sizes are very large and confidence

intervals for measures of association are therefore very narrow. 30 These results provide an excellent example of the phenomenon. The three studies all show consistent, large effect estimates (0.52 to 0.68), which increases confidence in our findings.



# **Discussion**

### **Statement of Principle Findings**

Our analysis of the Ontario WSIB's administrative data revealed that older claimants who were fully disabled at 4 weeks due to uncomplicated, acute LBP and who are reimbursed for opioid prescription in the first 4 weeks of their claim were more likely to experience prolonged claim duration. Higher pre-injury income was also associated with prolonged claim duration, but only among persistent claims. Injured workers employed by organizations with a RTW program and/or missing information on union affiliation were likely to resolve their claim faster. Neither early receipt of reimbursement for physiotherapy nor chiropractic care for uncomplicated LBP was associated with claim duration.

#### **Strengths & Weaknesses**

A priori specification of our regression model and stating the anticipated direction of included independent variables, as well as the assessment of the proportional hazards assumption for all independent variables, provide greater confidence in our findings. In 2005 the Ontario WSIB did not capture any measure of LBP injury severity on Forms 6-8 and we were therefore unable to adjust for this factor. The Ontario WSIB Health Professional's Report (Form 8) was updated in August 2011, and now includes capture of baseline pain. We did attempt to include patients with similar injuries by restricting our cohort to acute, uncomplicated LBP, and while there may still be important differences in injury severity in our cohort, our findings regarding the association with early opioid use and delayed claim recovery are consistent with

other studies that have adjusted for low back injury severity.<sup>13, 14</sup> Other strengths include limited missing data, correction of identifiable data errors and inconsistencies, and validation checks to ensure the accuracy of the data used to inform our regression model.

Our study has several limitations. First, it was a retrospective cohort study in which the reasons for reimbursement of physiotherapy, chiropractic or opioid prescription are uncertain. Thus, despite our adjustments for potential confounders, it remains possible that acute LBP claimants who chose to receive these healthcare interventions were prognostically different than those claimants who did not. Second, the WSIB database captures only those physiotherapy or chiropractic treatments that are reimbursed by the WSIB, and it is possible that some patients paid out-of-pocket to receive these services. It is highly unlikely that patients would have received opioids outside of WSIB reimbursement (personal communication, Dr. Norman Buckley, Chair of Anesthesiology, McMaster University). Third, a number of variables that may be important to consider were unavailable (e.g., patient expectations regarding recovery<sup>32</sup>), and some variables were not optimally collected. For example, chiropractic and physiotherapy are professions and not modalities, and there were no details of treatment provided. Fourth, our study focused on workers with acute LBP who were receiving disability benefits from the Ontario WSIB for at least 4-weeks in 2005, and we cannot say whether our findings are generalizable to other disabled workers. We are, however, unaware of any major changes in practices among Ontario chiropractors or physiotherapists since 2005, and there is evidence that both rates of opioids prescriptions and average morphine equivalent dose for non-malignant pain have increased since 2005, which would suggest that our findings regarding early reimbursement for opioids apply to a greater proportion of current

BMJ Open: first published as 10.1136/bmjopen-2015-007836 on 26 August 2015. Downloaded from http://bmjopen.bmj.com/ on April 27, 2024 by guest. Protected by copyright.

Predicting Workers' Compensation claim duration

WSIB LBP claimants.<sup>33</sup> Finally, our primary outcome, time to claim closure, is a surrogate for patient-important outcomes such as functional restoration or return to work; however, claim closure and faster claim resolution is associated with functional recovery among adults disabled by non-severe whiplash injuries,<sup>34</sup> which provides some assurances that patients who resolve their disability claim are also likely to experience clinical improvement.

## Our Findings in the Context of Other Relevant Literature

Our finding that older age is associated with prolonged claim duration is consistent with the literature. <sup>35</sup> We predicted that injured workers employed by companies that had formal RTW programs would resolve their claims faster, and this was supported by our findings. It is unclear why missing data for union membership was associated with shorter claim duration. Similarly, reasons why higher pre-disability income was associated with prolonged claim duration, but only among persistent claims, are uncertain. Possibilities include that injured workers with higher salaries who do not resolve their claims in the initial 6-months may find it more difficult to identify suitable employment at similar earnings levels, or that compared to other workers (i.e. with lower pre-disability income) workers with higher salaries can accommodate limited compensatory income for longer periods of time.

Although there are no randomized controlled trials exploring the effect of physiotherapy, chiropractic care, or early opioid use for workers with acute, uncomplicated LBP receiving lost-time compensation benefits (Table S2 in File S1), our systematic review identified 6 observational studies that are relevant to our findings. Turner and colleagues followed 1,885 workers from Washington for 1 year after they had been awarded Workers'

Compensation benefits for acute LBP. In a comprehensive regression model adjusted for multiple sociodemographic, employment, clinical, healthcare, and administrative factors – including injury severity – they found that workers who attended a chiropractor first, versus a primary care provider, were significantly less likely to remain on disability benefits at 1 year (adjusted OR = 0.41, 95% CI = 0.24, 0.70). <sup>16</sup>

Wasiak and colleagues found that workers in Florida, USA, with acute low back injuries who were reimbursed for limited chiropractic care (<30 days) experienced an 8.6% shorter duration of work disability versus Workers' Compensation claimants who were reimbursed for prolonged chiropractic care (>30 days). The authors did exclude severe injuries from their population, but were unable to adjust for injury severity within their sample. Further, chiropractic care was collected after baseline and not treated as time-dependent, and so it is not surprising that claims with longer duration also received more chiropractic care.

Lemstra and Olszynski explored the effect of standard care (which included long waiting lists for physiotherapy) to provision of rapid rehabilitation services on Workers' Compensation claim duration from a company in Saskatchewan, Canada. After adjusting for a number of factors, including age and injury severity, longer claim duration was associated with both chiropractic care (adjusted HR = 2.88, 95% CI = 1.45, 5.73) and physical therapist involvement (adjusted HR = 19.88, 95% CI = 7.95, 39.77). The authors collected healthcare provider utilization data after baseline and did not treat these variables as time-dependent, and so an alternative explanation is that claims with longer duration are simply more likely to involve either chiropractors or physical therapists.

Predicting Workers' Compensation claim duration

Canada is currently the largest per-capita consumer of opioids in the world; <sup>38</sup> however, prescribing patterns in Ontario show considerable variation between primary care providers. <sup>39</sup> Workers' Compensation data from the US has shown an almost 10-fold range (5.7% to 52.9%) in the early prescription of opioid medications between various states, suggesting that local prescribing patterns have significant influence on the use of these analgesics. <sup>40</sup> Findings from a study of Workers' Compensation nonspecific LBP claims (that did not adjust for injury severity) revealed that, compared with a no opioid reference group, the odds of chronic work loss were six times greater for claimants who used strong opioids and 11-14 times greater for claimants with opioid prescriptions which exceeded 90 days. <sup>41</sup> We found that reimbursement for early opioid use by the Ontario WSIB was associated with prolonged claim duration for uncomplicated back pain, and pooling of our data with similar studies <sup>14, 15</sup> shows a consistent effect, which increases confidence in our findings.

#### **Implications & Future Research**

To manage their growing unfunded liability, the Ontario WSIB has focused on increasing their claim denial rate, decreasing benefits to injured workers, reducing WSIB staff, and raising employer premiums. <sup>9, 10</sup> Most employers are obligated to pay WSIB premiums because they are legally bound to provide injury benefits to their employees, and the Ontario WSIB is protected by laws prohibiting competition in the marketplace. Another strategy is to optimize clinical management of injured workers. Our findings, which were not adjusted for illness severity, suggest that receiving reimbursement from the Ontario WSIB for early chiropractic care or physiotherapy for acute, uncomplicated LBP is not associated with shorter time to claim closure

Predicting Workers' Compensation claim duration

and receiving reimbursement for early opioids was linked with longer claim duration. We did find a non-significant association between early reimbursement for chiropractic care and shorter claim duration, which was lost by 6-months. In 2005 the Ontario WSIB typically limited reimbursement for chiropractic care to no more than 3-months after a low back injury and the change in association that we found may reflect discontinuation of reimbursement by the WSIB and consequent termination of chiropractic care.

The Ontario WSIB should continue to incorporate established prognostic factors for recovery into their baseline data collection forms, as well as outcomes of direct importance to patients, such as functional recovery, to facilitate more rigorous analyses of their administrative data. However, observational data cannot establish causality and high quality, randomized controlled trials are urgently needed to confirm or refute our findings.

#### **Author affiliations**

- <sup>1</sup> The Michael G. DeGroote Institute for Pain Research and Care, McMaster University,
  Hamilton, Canada
- <sup>2</sup> Department of Anesthesia, McMaster University, Hamilton, Canada
- <sup>3</sup> Department of Clinical Epidemiology and Biostatistics, McMaster University, Hamilton, Canada
- <sup>4</sup> Stanford Prevention Research Center, Department of Medicine, Stanford University, Stanford, California, USA
- <sup>5</sup> Department of Anaesthesia & Pain Medicine, the Hospital for Sick Children, Toronto, Canada

## Acknowledgements

The authors thank Dr. Gordon Guyatt for helpful discussion.

#### **Contributors**

JWB, SE, D-HA and SDW designed the study. DH-A and LW conducted data analysis and SDW provided statistical advice. JWB, SE, D-HA and SDW were involved in interpreting the data. RC designed and conducted all literature searches. JWB drafted the manuscript and wrote the final version. All authors critically revised the manuscript, provided comment and approved the final version for publication.

#### **Funding**

This study was funded by research grants from the Ontario Workers Safety and Insurance Board Research Advisory Council and the Ontario Chiropractic Association. Shanil Ebrahim was

supported by a MITACS Elevate Postdoctoral Fellowship Award. The funding sources had no role in design or conduct of the study; the collection, management, analysis, or interpretation

Predicting Workers' Compensation claim duration

### **Competing interests**

Jason Busse acts as a consultant to Prisma Health Canada, a private incorporated company funded by employers and insurers that consults on and manages long-term disability claims.

of the data; or the preparation, review, or approval of the manuscript.

### **Ethics approval**

The study was approved by the Health Sciences Research Ethics Board of the University of Toronto.

#### Provenance and peer review

Not commissioned; externally peer reviewed.

#### Data sharing statement

No additional data are available.

### **Open Access**

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work noncommercially, and license their derivative works on different terms,

provided the original work is properly cited and the use is non-commercial. See: http:// creativecommons.org/licenses/by-nc/4.0/



Predicting Workers' Compensation claim duration

#### References

- 1. Thiese MS, Hegmann KT, Wood EM, Garg A, Moore JS, Kapellusch JM, et al. Low-back pain ratings for lifetime, 1-month period, and point prevalences in a large occupational population. *Hum Factors* 2014; 56: 86-97.
- 2. Deyo RA, Phillips WR. Low back pain. A primary care challenge. *Spine (Phila Pa 1976)* 1996; 21: 2826-2832.
- 3. Hoy D, March L, Brooks P, Blyth F, Woolf A, Bain C, et al. The global burden of low back pain: estimates from the Global Burden of Disease 2010 study. *Ann Rheum Dis* 2014; 73: 968-974.
- Brown A, Angus D, Chen S, Tang Z, Milne S, Pfaff J, et al. Costs and outcomes of chiropractic treatment for low back pain [Technology report no 56]. Ottawa: Canadian Coordinating Office for Health Technology Assessment, 2005.
- 5. WSIB-CSPAAT Ontario. By the Numbers: 2012 WSIB Statistical Report. Schedule 1 Toronto, ON: WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2014. [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S1.pdf] (Accessed August 3, 2014).
- 6. WSIB-CSPAAT Ontario. By the Numbers: 2012 WSIB Statistical Report. Schedule 2 WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2014.
  [http://www.wsibstatistics.ca/WSIB-StatisticalReport\_S2.pdf] (Accessed August 3, 2014).

Predicting Workers' Compensation claim duration

- 7. WSIB-CSPAAT Ontario. First Quarter Sufficiency Report to Stakeholders WSIB-CSPAAT Ontario, Workplace Safety & Insurance Board, Commission de la sécurité professionnelle et de l'assurance contre les accidents du travail, 2013.
  [http://www.wsib.on.ca/files/Content/SufficiencySufficiencyReport2013/2013Q1WSIBSufficiencyReport.pdf] (Accessed August 3, 2014).
- 8. Ontario WSIB funding fiasco [Editorial] (November 18, 2008). National Post. 2008. [http://www.nationalpost.com/opinion/story.html?id=eb83e348-1660-4771-9804-e47a502aa756] (Accessed August 3, 2014).
- 9. UFCW Canada. WSIB President gets \$80K bonus on the backs of injured workers. Media & News Canada's Best Labour and Social Justice News. 2012.
  [http://www.ufcw.ca/index.php?option=com\_content&view=article&id=2760:wsib-president-gets-80k-bonus-on-the-backs-of-injured-workers&Itemid=6&lang=en]
  (Accessed August 3, 2014).
- 10. WSIB raises its rates to tackle \$12 billion in unfunded liabilities. Machinery and Equipment MRO Maintenance, Repair and Operations. 2010.
  [http://www.mromagazine.com/news/wsib-raises-its-rates-to-tackle-12-billion-in-unfunded-liabilities/1000391835/?&er=NA] (Accessed August 3, 2014).
- 11. Henschke N, Maher CG, Refshauge KM, Herbert RD, Cumming RG, Bleasel J, et al.

  Prognosis in patients with recent onset low back pain in Australian primary care:
  inception cohort study. *BMJ* 2008; 337: a171.

Predicting Workers' Compensation claim duration

- and guideline adherent physical therapy for low back pain on utilization and costs. BMC Health Serv Res 2015; 15: 150.
- 13. 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: 2127-2132.
- Franklin GM, Stover BD, Turner JA, Fulton-Kehoe D, Wickizer TM. Early opioid prescription 14. and subsequent disability among workers with back injuries: the Disability Risk Identification Study Cohort. Spine (Phila Pa 1976) 2008; 33: 199-204.
- Gross DP, Stephens B, Bhambhani Y, Haykowsky M, Bostick GP, Rashiq S. Opioid prescriptions in Canadian workers' compensation claimants: prescription trends and associations between early prescription and future recovery. Spine (Phila Pa 1976) 2009; 34: 525-531.
- Turner JA, Franklin G, Fulton-Kehoe D, Sheppard L, Stover B, Wu R, et al. ISSLS prize 16. winner: early predictors of chronic work disability: a prospective, population-based study of workers with back injuries. Spine (Phila Pa 1976) 2008; 33: 2809-2818.
- von Elm E, Altman DG, Egger M, Pocock SJ, Gotzsche PC, Vandenbroucke JP. The 17. Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) Statement: guidelines for reporting observational studies. Int J Surg 2014; 12: 1495-1499.
- Collins GS, Reitsma JB, Altman DG, Moons KG. Transparent Reporting of a multivariable prediction model for Individual Prognosis Or Diagnosis (TRIPOD): the TRIPOD Statement. Br J Surg 2015; 102: 148-158.

- 19. da CMCL, Maher CG, Hancock MJ, McAuley JH, Herbert RD, Costa LO. The prognosis of acute and persistent low-back pain: a meta-analysis. *CMAJ* 2012; 184(11): E613-E624.
- 20. Frank JW, Brooker AS, DeMaio SE, Kerr MS, Maetzel A, Shannon HS, et al. Disability resulting from occupational low back pain. Part II: What do we know about secondary prevention? A review of the scientific evidence on prevention after disability begins. *Spine* (*Phila Pa 1976*) 1996; 21: 2918-2929.
- 21. Busse J, Steenstra I, Riva J, Ebrahim S, de Bruin L, Guyatt G. Predictors of prolonged recovery following acceptance for disability benefits: a systematic review of observational studies. *Occup Environ Med* 2011; 68(Suppl 1): A97.
- 22. Schandelmaier S, Ebrahim S, Burkhardt SC, de Boer WE, Zumbrunn T, Guyatt GH, et al.

  Return to work coordination programmes for work disability: a meta-analysis of randomised controlled trials. *PLoS ONE* 2012; 7: e49760.
- 23. Harrell FE. Multivariate modeling strategies. In: Harrell FE, editor. Regression Modeling Strategies With Applications to Linear Models, Logistic Regression and Survival Analysis. New York, NY: Springer; 2001. p. 53-85.
- 24. Zhang J, Yu KF. What's the relative risk? A method of correcting the odds ratio in cohort studies of common outcomes. *JAMA* 1998; 280: 1690-1691.
- 25. Montori V, Ioannidis J, Cook DJ. Fixed-effects and random-effects models. In: Guyatt GH, Rennie D, Meade MO, Cook DJ, editors. Users' guides to the medical literature: A manual for evidence-based clinical practice (2nd ed). USA: McGraw Hill; 2008. p. 555-62.
- 26. Higgins JP, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. BMJ 2003; 327: 557-560.

- 27. Higgins JPT, Green Se. Cochrane Handbook for Systematic Reviews of Interventions

  Version 5.1.0 [updated March 2011]: The Cochrane Collaboration; 2011.
- 28. Atkins D, Best D, Briss PA, Eccles M, Falck-Ytter Y, Flottorp S, et al. Grading quality of evidence and strength of recommendations. *BMJ* 2004; 328: 1490.
- 29. Guyatt GH, Oxman AD, Sultan S, Glasziou P, Akl EA, Alonso-Coello P, et al. GRADE guidelines: 9. Rating up the quality of evidence. *J Clin Epidemiol* 2011; 64: 1311-1316.
- 30. Rucker G, Schwarzer G, Carpenter JR, Schumacher M. Undue reliance on I(2) in assessing heterogeneity may mislead. *BMC Med Res Methodol* 2008; 8: 79.
- 31. The Ontario WSIB Health Professional's Report (Form 8)

  [http://www.wsib.on.ca/cs/groups/public/documents/staticfile/c2li/mdey/~edisp/wsib01
  2197.pdf] (Accessed July 2, 2015).
- 32. Cole DC, Mondloch MV, Hogg-Johnson S. Listening to injured workers: how recovery expectations predict outcomes--a prospective study. *CMAJ* 2002; 166: 749-754.
- 33. Gomes T, Mamdani MM, Paterson JM, Dhalla IA, Juurlink DN. Trends in high-dose opioid prescribing in Canada. *Can Fam Physician* 2014; 60: 826-832.
- 34. Cote P, Hogg-Johnson S, Cassidy JD, Carroll L, Frank JW. The association between neck pain intensity, physical functioning, depressive symptomatology and time-to-claim-closure after whiplash. *J Clin Epidemiol* 2001; 54: 275-286.
- 35. Hadler NM. The bane of the aging worker. *Spine (Phila Pa 1976)* 2001; 26: 1309-1312.
- 36. Wasiak R, Kim J, Pransky GS. The association between timing and duration of chiropractic care in work-related low back pain and work-disability outcomes. *J Occup Environ Med* 2007; 49: 1124-34.

Predicting Workers' Compensation claim duration

- Lemstra M, Olszynski WP. The effectiveness of standard care, early intervention, and occupational management in worker's compensation claims. *Spine (Phila Pa 1976)* 2003;
   28: 299-304.
- 38. Pain and Policy Studies Group. PPSG Opioid Consumption Motion Chart. University of Wisconsin Paul P Carbone Comprehensive Cancer Center Madison, Wisconsin. 2011 [https://ppsg.medicine.wisc.edu/chart] (Accessed May 15, 2015).
- 39. Dhalla IA, Mamdani MM, Gomes T, Juurlink DN. Clustering of opioid prescribing and opioid-related mortality among family physicians in Ontario. *Can Fam Physician* 2011; 57: e92-e96.
- 40. Webster BS, Cifuentes M, Verma S, Pransky G. Geographic variation in opioid prescribing for acute, work-related, low back pain and associated factors: a multilevel analysis. *Am J Ind Med* 2009; 52: 162-171.
- 41. Volinn E, Fargo JD, Fine PG. Opioid therapy for nonspecific low back pain and the outcome of chronic work loss. *Pain* 2009; 142: 194-201.

Predicting Workers' Compensation claim duration

# **Figure Legends**

- Figure 1: Kaplan-Meier curve for time to claim closure
- Figure 2: Kaplan-Meier curve for early reimbursement for opioid prescription
- Figure 3: Kaplan-Meier curve for early reimbursement for physiotherapy
- Figure 4: Kaplan-Meier curve for early reimbursement for chiropractic care
- Figure 5: The association between early opioid use/prescription and claim duration

# **Supporting Information Legends**

- Table S1: Search strategy for the Cochrane Back Group trials registry
- Table S2: Randomized controlled trials of acute low back pain assessing the effect of opioids, chiropractic care, or physical therapy, and their generalizability to injured workers receiving disability benefits
- Table S3: Sensitivity analysis of factors associated with time to claim closure, entering chiropractic care and physiotherapy as time-dependant co-variates (n=1,442)
- Table S4: Search strategy for observational studies exploring the association of early opioid, physiotherapy, or chiropractic care for workers with acute low back pain with claim duration
- Figure S1: A flow diagram of the literature search process for randomized controlled trials assessing the effect of opioids, physiotherapy, or chiropractic care for acute low back pain.
- Figure S2: A flow diagram of the literature search process for observational studies assessing the effect of early opioids, physiotherapy, or chiropractic care for acute low back pain.

### Table 1: Description of model variables

Variable	Anticipated Direction of Effect	
Age (in decades)	older age: -	
Gender	female: -	
First language	non-English: -	
Pre-disability income	higher income: -	
Reimbursement for opioid prescription	opioid reimbursement: -	
in the first 4-weeks of a claim		
Prior claim(s)	prior claim: -	
Union membership	Union member: +	
Employer RTW-program	RTW program: +	
Work-relatedness	work-related: +	
Reimbursement for ≥3 chiropractic	DC care: +	
treatments in the first 4-weeks of a		
claim		
Reimbursement for ≥3 physiotherapy	PT care: +	
treatments in the first 4-weeks of a		
claim		

- +: associated with faster claim closure
- -: associated with slower claim closure

Predicting Workers' Compensation claim duration

Age in years, mean (SD)	41.3 (10.5)
Gender, n (%)	
Female	552 (38.3)
Male	890 (61.7)
First language, n (%)	
English	1372 (95.1)
Other	70 (4.9)
Pre-disability income (dollars/week)	
mean (SD)	731.4 (332.5)
Opioid prescription reimbursed by WSIB in the first 4-weeks of	
claim, n (%)	
Yes	136 (9.4)
No	1306 (90.6)
Prior WSIB claim, n (%)	
Yes	1091 (75.7)
No	351 (24.3)
Union membership, n (%)	
Yes	610 (42.3)
No	656 (45.5)
Missing data	176 (12.2)
Employer RTW program, n (%)	
Yes	1042 (72.3)
No	278 (19.3)
Missing data	122 (8.5)
Employer doubts work-relatedness of injury, n (%)	
Yes	195 (13.5)
No	1051 (72.9)
Missing data	196 (13.6)
Chiropractic care reimbursed by WSIB during claim, n (%)	391 (27.1)
Early Chiropractic care (≥3 treatments received within the first 28	
days), n (%)	247 (17.1)
Dhe daile and a daile and by MCID de day a daile a (0/)	706 (54.5)
Physiotherapy reimbursed by WSIB during claim, n (%)	786 (54.5)
Early Physiotherapy (≥3 treatments received within the first 28 days), n (%)	388 (26.9)

Table 3: Factors associated with time to claim closure (n=1,442)\*

	Univariate		Multivariable		
	Hazard Ratio (99% CI) p-value		Adjusted Hazard Ratio (99% CI)	p-value	
Baseline predictors					
Age categories in years		<0.001		<0.001	
15 to <25	1.00		1.00		
25 to <35	0.88 (0.65, 1.19)		0.79 (0.58, 1.09)		
35 to <45	0.78 (0.59, 1.04)		0.70 (0.52, 0.95)		
45 to <55	0.76 (0.56, 1.02)		0.67 (0.49, 0.91)		
55 to 65	0.56 (0.40, 0.80)		0.52 (0.36, 0.74)		
Gender		0.114		0.446	
Females	1.09 (0.95, 1.26)		0.96 (0.82, 1.12)		
Males	1.00		1.00		
First language		0.137		0.312	
English	1.00		1.00		
Other	0.83 (0.59, 1.15)		0.88 (0.63, 1.23)		
Pre-disability income in	, -,				
dollars per week					
At 60 days:					
≤480	1.00	_	1.00	_	
481-694	1.09 (0.89, 1.34)	0.273	1.03 (0.83, 1.27)	0.749	
695-920	1.13 (0.92, 1.39)	0.137	0.96 (0.77, 1.21)	0.678	
>920	1.08 (0.88, 1.33)	0.326	0.93 (0.74, 1.18)	0.443	
At 180 days:	2.00 (0.00) 2.00)	0.020	0.00 (0.1. 1, 2.20)	01110	
≤480	1.00		1.00	_	
481-694	0.89 (0.68, 1.17)	0.267	0.89 (0.67, 1.17)	0.266	
695-920	0.84 (0.63, 1.12)	0.115	0.73 (0.54, 0.98)	0.006	
>920	0.72 (0.54, 0.97)	0.004	0.63 (0.46, 0.86)	<0.001	
At 365 days:	0.72 (0.5 1, 0.57)	0.001	0.05 (0.10) 0.00)	10.001	
≤480	1.00	_	1.00	_	
481-694	0.65 (0.36, 1.18)	0.064	0.71 (0.38, 1.31)	0.147	
695-920	0.53 (0.28, 1.02)	0.012	0.47 (0.24, 0.91)	0.003	
>920	0.39 (0.20, 0.75)	<0.001	0.34 (0.17, 0.68)	<0.001	
Early Reimbursement for	0.33 (0.20, 0.73)	<0.001	0.34 (0.17, 0.00)	<0.001	
Opioid prescription (within		\0.001		10.001	
the first 28 days)					
Yes	0.62 (0.48, 0.80)		0.68 (0.53, 0.88)		
No	1.00		1.00		
Prior claims	1.00	0.306	1.00	0.709	
Yes	1.07 (0.91, 1.26)	0.300	1.03 (0.86, 1.22)	0.703	
No	1.00		1.00		
Union membership	1.00	<0.001	1.00	0.016	
Yes	1.29 (1.11, 1.50)	\0.001	1.14 (0.96, 1.35)	0.010	
No	1.00		1.00		
Missing	1.34 (1.07, 1.68)		1.27 (1.01, 1.59)		
	1.34 (1.07, 1.00)	<0.001	1.27 (1.01, 1.33)	<0.001	
Employer RTW program Yes	1.73 (1.43, 2.09)	<0.001	1 79 /1 /5 2 19\	<0.001	
	1.73 (1.43, 2.09)		1.78 (1.45, 2.18) 1.00		
No Missing					
Missing  Doubt Work relatedness	1.17 (0.87, 1.58)	0.174	1.17 (0.86, 1.59)	0.130	
Doubt Work-relatedness	0.94 (0.76, 1.16)	0.174	0.88 (0.71, 1.08)	0.138	

No	1.00		1.00	
Missing	1.13 (0.92, 1.39)		1.08 (0.87, 1.33)	
Early Reimbursement for				
Chiropractic care				
At 60 days:	1.19 (0.99, 1.45)	0.017	1.15 (0.94, 1.41)	0.067
At 180 days:	0.91 (0.65, 1.24)	0.437	0.90 (0.65, 1.24)	0.392
At 365 days:	0.60 (0.29, 1.23)	0.067	0.61 (0.29, 1.29)	0.089
Early Reimbursement for	0.98 (0.84, 1.15)	0.726	1.01 (0.86, 1.19)	0.848
Physiotherapy				

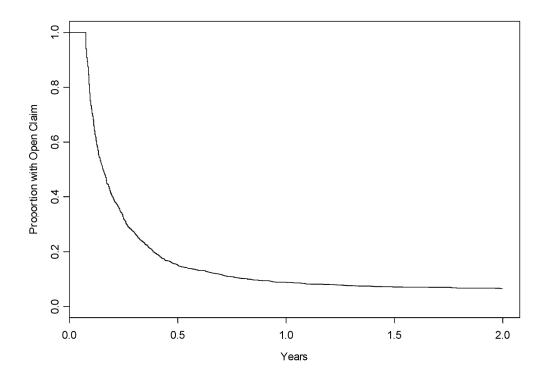
<sup>\*</sup> HR>1 indicates faster claim closure; RTW = return to work; Early reimbursement for chiropractic or physiotherapy = 3 or more treatments received within the first 28 days 3 01 me.

Table 4: Observational studies exploring the association between early opioid use and Workers' Compensation claim duration

Study	Population	Opioid variable tested	Adjustments	Dependent variable	Results*
Webster et al.,	8,443 American	Receipt of	Injury severity,	Change in	1-140mg MEA
2007	Workers'	opioids within	age, gender,	mean	5.4 days, 95%CI = -14.6 to 25.0
	Compensation	the first 15 days	length of job	disability	141-225mg MEA
	claimants with	of claim	tenure	duration	21.9 days, 95%CI = 3.2 to 40.6
	new-onset,				226-450mg MEA
	disabling LBP	100			43.8 days, 95%CI = 23.7 to 63.9
	_				>450mg MEA
					69.1 days, 95%CI = 49.3 to 89.0
Franklin et al.,	1,843 Washington,	Reimbursement	Age, gender,	Receipt of	1-150mg MED
2008	US, Workers'	for opioids	race,	wage	OR = 1.9, 95%Cl = 1.2 to 3.1
	Compensation	within 6 weeks	education,	replacement	151-300mg MED
	claimants with	of 1 <sup>st</sup> medical	injury severity,	benefits at 1-	OR = 2.0, 95%Cl = 1.2 to 3.3
	new-onset,	visit for LBP	pain intensity,	year	301-650mg MED
	disabling LBP		Roland		OR = 1.6, 95%CI = 0.9 to 2.6
			disability		>650mg MED
			questionnaire		OR = 1.9, 95%CI = 1.2 to 2.9
Gross et al.,	47,784 Alberta,	Reimbursement	Age, gender,	Receipt of	No early opioids
2009	Canada, Workers'	for opioids	annual salary,	wage	HR = 1.94, 95%CI = 1.86 to 2.02
	Compensation	within the first 2	year of claim,	replacement	
	claimants with	weeks of claim	number of	benefits at 1-	
	new-onset,		previous	year	
	disabling LBP		claims		

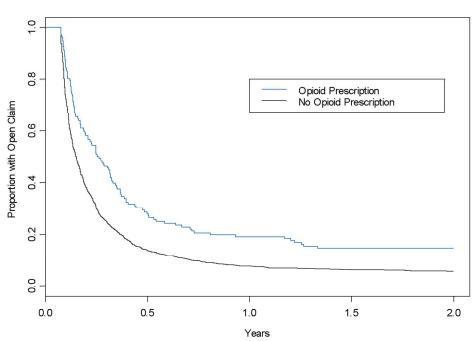
<sup>\*</sup> The reference group is no early receipt of opioids for Franklin et al., and Webster et al. 95% CI = 95% confidence interval; LBP = low back pain; MEA = morphine equivalent amount; MED = morphine equivalent dose; OR = odds ratio; HR = hazard ratio

Predicting Workers' Compensation claim duration



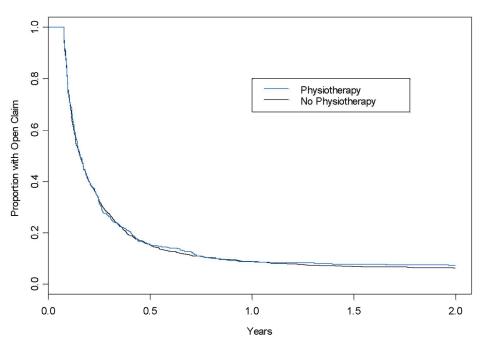
Kaplan-Meier curve for time to claim closure 181x130mm (300 x 300 DPI)

# Claim Closure by Opioid Prescription



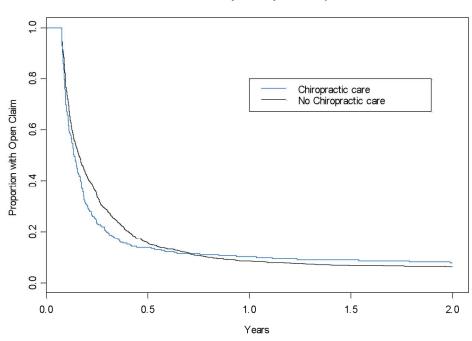
Kaplan-Meier curve for early reimbursement for opioid prescription  $145 \times 110 \text{mm}$  (300 x 300 DPI)

## Claim Closure by Early Physiotherapy

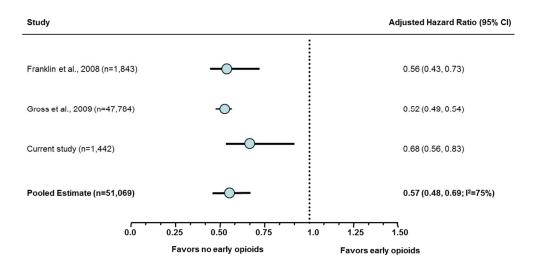


Kaplan-Meier curve for early reimbursement for physiotherapy 146x107mm (300 x 300 DPI)

## Claim Closure by Early Chiropractic Care



Kaplan-Meier curve for early reimbursement for chiropractic care 149x109mm~(300~x~300~DPI)



The association between early opioid use/prescription and claim duration 177x88mm (300 x 300 DPI)

### Table S1: Search strategy for the Cochrane Back Group trials registry\*

Search Strategy in Cochrane Library:

\*:ti,ab,kw in Trials, with Back Group in Review Groups 6476

Endnote search strategy:

Any field contains: acute AND Any field contains chiropractic (43) Any field contains: acute AND Any field contains physiotherapy\* (87)

Any field contains: acute AND Any field contains opioid (30)

Any field contains: opioid NOT Any field contains: surgery NOT Any field contains:

chronic (28)

Any field contains: emergency (57) NOT Any field contains: whiplash (18)

\*On May 1, 2015 we used the Cochrane Library to search the Cochrane Back Review Group (CBRG) Trials Register, which we exported to Endnote. We also screened the included studies lists of recent CBRG Reviews of chiropractic and physiotherapy interventions for acute low back pain (Franke et al., 2015, Rubinstein et al., 2012).

Table S2: Randomized controlled trials of acute low back pain assessing the effect of opioids, chiropractic care, or physical therapy, and their generalizability to injured workers receiving disability benefits

Study	Participants and Interventions	Representation of injured workers receiving lost-time disability benefits for acute low back pain
Glover 1974	84 patients with unilateral low back pain (LBP), randomized to manipulation or control (sham diathermy)	No mention regarding receipt of lost-time disability benefits
Bergquist- Ullman 1977	217 patients consulting a workplace health centre with acute or subacute LBP randomized to back school, physiotherapy or placebo	At least 88% of patients were enrolled with acute LBP, and of the 217, 184 were "sick-listed" for a median of 21 days during the study, but there was no mention regarding receipt of lost-time disability benefits
Rasmussen 1979	26 patients with LBP duration <3weeks, randomized to manipulation or diathermy	No mention regarding receipt of lost-time disability benefits
Hoehler 1981	95 patients with palpatory cues indicating hyperalgesia or a restricted or painful range of vertebral motion, randomized to rotational manipulation of the trunk or massage	No mention regarding receipt of lost-time disability benefits
Farrell 1982	48 subjects with acute LBP duration <3 weeks, randomized to passive mobilization and manipulation or combination of diathermy, exercises and ergonomic advice	No mention regarding receipt of lost-time disability benefits
Gilbert 1985	270 patients presenting with LBP, randomized to bed rest or physiotherapy	The authors reported that "people who were receiving workman's compensation were also slower to recover." p.794, but they did not report any associated data (e.g. how many patients were receiving disability benefits, or the quantitative results for this subgroup).
Waterworth 1985	112 patients with acute mechanical LBP, randomized to ergonomic advice/ Diflunisal or ergonomic advice/ conservative physiotherapy (ultrasound and exercise) or ergonomic advice/manipulation	The authors enrolled a mixed group of patients, that may include up to 54% receiving lost-time claim benefits, but that the proportion is unequal between study group (ranging from 47% to 63%) and the results are provided for the total population which precludes confident generalizability to only those who were receiving lost-time claim benefits.
Hadler 1987	54 subjects with acute LBP, randomized to mobilization or manipulation	Patients who were receiving disability benefits were explicitly excluded from this trial: "neither workers' compensation nor disability insurance should be at issued [sic] and the acute low-back pain must not be considered work-related."pg 703
MacDonald 1990	95 subjects with LBP duration <4 weeks, randomized to osteopathic manipulation or control (advice to rest and resume activities gradually)	"Less than 30% [of the study group] suffered loss of income because of disability" pg. 366

Cramer 1993	36 subjects with mechanical LBP less than two weeks duration randomized to side-lying manipulation, electrical stimulation and cold packs or control (detuned ultrasound, cold packs and 15-30 sec. gentle massage)	Patients who were receiving disability benefits were explicitly excluded from this trial. Inclusion criteria stipulate "no litigation or workers' compensation" (as per the review by Rubinstein 2012, pg. 52)	
Skargren 1997	323 patients with back and neck pain of mixed duration, randomized to chiropractic or physiotherapy.	51% of patients (166 of323) were on "sick-leave" when enrolled (Table 3, pg. 2170), but there was no mention regarding receipt of lost-time disability benefits.	
Innes 1998	123 patients with acute LBP, randomized to ketorolac or acetaminophen/codeine	Patients who were receiving disability benefits were limited during enrollment: "Because Workers Compensation status might influence response to therapy, we limited each site to 10 work-related back injuries, hoping to limit such cases to no more than half the total study enrollment." pg. 550	
Cherkin 1998	321 adults with acute LBP randomized to the McKenzie method of physical therapy, chiropractic manipulation, or provision of an educational booklet	patients who were receiving disability benefits were explicitly excluded from this trial: "Subjects who were involved in claims for compensation or litigation because of the back injurywere also excluded"; pg. 1022	
Seferlis 1998	180 patients sick-listed for < 2 weeks for LBP randomized to General Practitioner Program (rest, sick-leave, analgesics etc.) or Manual Therapy Program (autotraction, manipulation, mobilization etc.) or Intensive Training Program (information, muscle training and general condition training 3x/week for 8 weeks)	All enrolled patients were "sick listed for acute low-back pain for up to 2 weeks", but there is no mention regarding receipt of lost-time disability benefits.	
Morton 1999	29 patients with acute mechanical LBP, randomized to manipulation/ exercise or exercise alone	Patients who were receiving disability benefits were explicitly excluded from this trial: "Exclusion criteria were third-party, public liability or workers' compensation claimants"; pg. 185	
Veenema 2000	155 patients with musculoskeletal LBP, randomized to meperidine or ketorolac	No mention regarding receipt of lost-time disability benefits	
Metscher 2001	192 Patients with acute LBP randomized to dexketoprofen-trometamol or tramadolhydrochloride	Abstract in English, paper in German. No mention in the abstract about receipt of lost-time disability benefits	
Palangio 2002	147 patients with acute LBP, first episode or exacerbation of chronic condition with onset <48 hours before enrolment, randomized to combination hydrocodone 7.5 mg and ibuprofen 200 mg (HC/IB) or combination oxycodone 5 mg and acetaminophen 325 mg (OX/AC)	No mention regarding receipt of lost-time disability benefits	
Hofstee 2002	250 patients with sciatica of less than 1 months duration randomized to bed rest, physiotherapy or continuation of activities of daily living	No mention regarding receipt of lost-time disability benefits	
Johnstone 2002	12 patients with acute LBP with signs of psychological distress (DRAM score Modified Zung score >17) randomized to cognitive behavioral therapy and conventional physiotherapy or conventional physiotherapy alone	Patients with "ongoing medico legal issues" were excluded (pg.183). No mention regarding receipt of lost-time disability benefits	

Childs 2004	131 patients with LBP of median duration of 27 days, randomized to manipulation and exercise or exercise alone	39.8% of patients had missed work due to LBP, Table 2 pg. 925, but there was no mention regarding receipt of lost-time disability benefits
Hoiriis 2004	192 patients with LBP of 3 to 6 weeks duration randomized to chiropractic adjustments with placebo medicine, muscle relaxants with sham adjustments, or placebo medicine with sham adjustments	No mention regarding receipt of lost-time disability benefits
Salvador 2005	28 subjects randomly allocated to a muscle energy technique or transcutaneous electrical nerve stimulation (TENS)	Abstract in English, paper in Portuguese. No mention in the abstract about receipt of lost-time disability benefits
Brennan 2006	123 patients referred to physiotherapy for LBP less than 90 days duration, randomized to manipulation or specific exercise or stabilization	No mention regarding receipt of lost-time disability benefits
Santilli 2006	102 patients with acute moderate to severe radiating LBP of duration <10 days with MRI evidence of disc protrusion, randomized to manipulation or simulated manipulation	No mention regarding receipt of lost-time disability benefits
Hancock 2007	240 subjects with acute LBP duration < 6 weeks, randomized to four groups: control (placebo drug and placebo manipulation) or NSAIDs (diclofenac and placebo manipulation) or manipulation (placebo drug and active manipulation) or manipulation and NSAIDs (diclofenac and active manipulation)	No mention regarding receipt of lost-time disability benefits
Lee 2008	Study of 78 musculoskeletal pain patients, 67% with LBP, randomized to tramadol/paracetemol (n=28 with LBP) or ketorolac/paracetemol (n=24 with LBP)	No mention regarding receipt of lost-time disability benefits
Lau 2008	110 patients with acute LBP, randomized to immediate intervention (advice to stay active, Back Care booklet, reassurance, advice, interferential current therapy) or control (walking training and prescription of walking aids as indicated) followed by outpatient physiotherapy (for both groups)	12% of patients (13 of 110) had work-related injuries (see Table 1), but no mention regarding receipt of lost-time disability benefits
Selkow 2009	20 subjects with acute LBP, randomly allocated to muscle energy technique or sham manual treatment	No mention regarding receipt of lost-time disability benefits
Cleland 2009	112 subjects with LBP, that met 4 out of 5 criteria for a clinical prediction rule for LBP likely to respond to manipulation, randomized to supine thrust manipulation, side-lying thrust manipulation or non-thrust manipulation	Only 6% of patients were unable to work due to LBP. Table 2, pg.2724.
Hallegraef 2009	64 patients with acute nonspecific LBP duration <16 days, randomized to manipulative therapy plus physical therapy or physical therapy alone	No mention regarding receipt of lost-time disability benefits

Sutlive 2009	60 subjects with LBP meeting 3 out of 5 criteria for a clinical prediction rule for LBP likely to respond to manipulation, randomized to lumbopelvic manipulation or neutral gap manipulation	Patients "with litigation pending for their LBP" were excluded.  No mention regarding receipt of lost-time disability benefits
Juni 2009	104 patients with acute LBP duration < 4 weeks, randomized to standard care with manipulation or standard care alone	No mention regarding receipt of lost-time disability benefits
Machado 2010	148 adults with acute LBP duration <6 weeks, randomized to the McKenzie method and first-line care (advice, reassurance and time-contingent acetaminophen) or first-line care alone	Only 3% of participants (4 of 146) were receiving disability benefits for their injury (see Table 1)
Lewis 2011	89 patients with acute LBP duration < 3 months, randomized to strain-counterstrain manual therapy/exercise or exercise alone	No mention regarding receipt of lost-time disability benefits
Biondi 2013	1664 patients with acute LBP, randomized to tapentadol or oxycodone	No mention regarding receipt of lost-time disability benefits
Goertz 2013	91 patients with acute LBP, duration < 4 weeks, randomized to standard medical care and chiropractic manipulation or standard care alone	No mention regarding receipt of lost-time disability benefits
Behrbalk 2014	65 adults with acute LBP, randomized to morphine or morphine/promethazine	No mention regarding receipt of lost-time disability benefits
Eken 2014	137 patients with moderate or severe acute LBP, randomized to paracetemol, morphine or dexketoprophen	No mention regarding receipt of lost-time disability benefits
Tanen 2014	44 patients with acute radicular LBP, randomized to lidocaine or ketorolac	No mention regarding receipt of lost-time disability benefits

#### References for Tables S1 and S2

- 1. Franke H, Fryer G, Ostelo RW, Kamper SJ. Muscle energy technique for non-specific low-back pain. *Cochrane Database Syst Rev* 2015; 2: CD009852.
- 2. Rubinstein SM, Terwee CB, Assendelft WJ, de Boer MR, van Tulder MW. Spinal manipulative therapy for acute low-back pain. *Cochrane Database Syst Rev* 2012; 9: CD008880.
- 3. Glover JR, Morris JG, Khosla T. Back pain: a randomized clinical trial of rotational manipulation of the trunk. *British journal of industrial medicine* 1974; 31: 59-64.
- 4. Bergquist-Ullman M, Larsson U. Acute low back pain in industry. A controlled prospective study with special reference to therapy and confounding factors. *Acta orthopaedica Scandinavica* 1977; 48(Suppl. 170): 1-117.
- 5. Rasmussen GG. Manipulation in treatment of low back pain (a randomized clinical trial. *Man-Med* 1979; 17: 8-10.
- 6. Hoehler FK, Tobis JS, Buerger AA. Spinal manipulation for low back pain. *JAMA* 1981; 245: 1835-1838.
- 7. Farrell JP, Twomey LT. Acute low back pain. Comparison of two conservative treatment approaches. *Medical journal of Australia* 1982; 1: 160-164.
- 8. Gilbert JR, Taylor DW, Hildebrand A, Evans C. Clinical practice of common treatments for low back pain in family practice. *British Medical Journal Clinical Research Ed* 1985; 291: 791-794.
- 9. Waterworth RF, Hunter IA. An open study of diflunisal, conservative and manipulative therapy in the management of acute mechanical low back pain. *New Zealand medical journal* 1985; 98: 372-375.
- 10. Hadler NM, Curtis P, Gillings DB, Stinnett S. A benefit of spinal manipulation as adjunctive therapy for acute low-back pain: a stratified controlled trial. *Spine* 1987; 12: 702-706.
- 11. MacDonald RS, Bell CMJ. An open controlled assessment of osteopathic manipulation in nonspecific low-back pain. *Spine* 1990; 15: 364-370.
- 12. Cramer GD, Humphreys CR, Hondras MA, McGregor M, Triano JJ. The Hmax/Mmax ratio as an outcome measure for acute low back pain. *Journal of manipulative and physiological therapeutics* 1993; 16: 7-13.
- 13. Skargren EI, Oberg BE, Carlsson PG, Gade M. Cost and effectiveness analysis of chiropractic and physiotherapy treatment for low back and neck pain. Six-month follow-up. *Spine* 1997; 22: 2167-2177.
- 14. Innes GD, Croskerry P, Worthington J, Beveridge R, Jones D. Ketorolac versus acetaminophen-codeine in the emergency department treatment of acute low back pain. *Journal of emergency medicine* 1998; 16: 549-556.
- 15. Cherkin DC, Deyo RA, Battie M, Street J, Barlow W. A comparison of physical therapy, chiropractic manipulation, and provision of an educational booklet for the treatment of patients with low back pain. *N Engl J Med* 1998; 339: 1021-1029.
- 16. Seferlis T, Németh G, Carlsson AM, Gillström P. Conservative treatment in patients sicklisted for acute low-back pain: a prospective randomised study with 12 months' follow-up. *European spine journal* 1998; 7: 461-470.
- 17. Morton JE. Manipulation in the treatment of acute low back pain. *Journal of manual & manipulative therapy* 1999; 7: 182-189.
- 18. Veenema KR, Leahey N, Schneider S. Ketorolac versus meperidine: ED treatment of severe musculoskeletal low back pain. *American journal of emergency medicine* 2000; 18: 404-407.

19. Metscher B, Kübler U, Jahnel-Kracht H. Dexketoprofen-trometamol and tramadol in acute lumbago [German]. *Fortschritte der Medizin Originalien* 2001; 118: 147-151.

- 20. Palangio M, Morris E, Doyle RT, Dornseif BE, Valente TJ. Combination hydrocodone and ibuprofen versus combination oxycodone and acetaminophen in the treatment of moderate or severe acute low back pain. *Clinical therapeutics* 2002; 24: 87-99.
- 21. Hofstee DJ, Gijtenbeek JM, Hoogland PH, Houwelingen HC, Kloet A, Lötters F, et al. Westeinde sciatica trial: randomized controlled study of bed rest and physiotherapy for acute sciatica. *Journal of neurosurgery* 2002; 96(1 Suppl): 45-49.
- 22. Johnstone R, Donaghy M, Martin D. A pilot study of a cognitive-behavioural therapy approach to physiotherapy, for acute low back pain patients, who show signs of developing chronic pain. *Advances in physiotherapy* 2002; 4: 182-188.
- 23. Childs JD, Fritz JM, Flynn TW, Irrgang JJ, Johnson KK, Majkowski GR, et al. A clinical prediction rule to identify patients with low back pain most likely to benefit from spinal manipulation: a validation study. *Annals of internal medicine* 2004; 141: 920-928.
- 24. Hoiriis KT, Pfleger B, McDuffie FC, Cotsonis G, Elsangak O, Hinson R, et al. A Randomized Clinical Trial Compring Chiropractic Adjustments to Muscle Relaxants for Subacute Low Back Pain. *Journal of manipulative and physiological therapeutics* 2004; 27: 388-398.
- 25. Salvador D, Neto PD, Ferrari FP. Application of muscle energy technique in garbage collectors with acute mechanical lumbar pain [Portuguese]. *Fisioterapia e Pesquisa* 2005; 12: 20-27.
- 26. Brennan GP, Fritz JM, Hunter SJ, Thackeray A, Delitto A, Erhard RE. Identifying subgroups of patients with acute/subacute "nonspecific" low back pain: results of a randomized clinical trial. *Spine* 2006; 31: 623-631.
- 27. Santilli V, Beghi E, Finucci S. Chiropractic manipulation in the treatment of acute back pain and sciatica with disc protrusion: a randomized double-blind clinical trial of active and simulated spinal manipulations. *Spine journal* 2006; 6: 131-137.
- 28. Hancock MJ, Maher CG, Latimer J, McLachlan AJ, Cooper CW, Day RO, et al. Assessment of diclofenac or spinal manipulative therapy, or both, in addition to recommended first-line treatment for acute low back pain: a randomised controlled trial. *Lancet* 2007; 370: 1638-1643.
- 29. Lee HKH, Ting SM, Lau FL. A randomised control trial comparing the efficacy of tramadol and paracetamol against ketorolac and paracetamol in the management of musculoskeletal pain in the emergency department. Hong Kong Journal of Emergency Medicine [Internet]. 2008; 15(1):[5-11 pp.]
- 30. Lau PM, Chow DH, Pope MH. Early physiotherapy intervention in an Accident and Emergency Department reduces pain and improves satisfaction for patients with acute low back pain: a randomised trial. *Australian journal of physiotherapy* 2008; 54: 243-249.
- 31. Selkow N, Grindstaff T, Cross K, Pugh K, Hertel J, Saliba S. Short-Term Effect of Muscle Energy Technique on Pain in Individuals with Non-Specific Lumbopelvic Pain: A Pilot Study. *Journal of Manual & Manipulative Therapy* 2009; 17: 14-18.
- 32. Cleland JA, Fritz JM, Kulig K, Davenport TE, Eberhart S, Magel J, et al. Comparison of the effectiveness of three manual physical therapy techniques in a subgroup of patients with low back pain who satisfy a clinical prediction rule: a randomized clinical trial. *Spine (Phila Pa 1976)* 2009; 34: 2720-2729.
- 33. Hallegraeff JM, Hallegraeff HJ, Greef M, Winters JC, Lucas C. Manipulative therapy and clinical prediction criteria in treatment of acute nonspecific low back pain. *Perceptual and motor skills* 2009; 108: 196-208.

- 34. Sutlive TG, Mabry LM, Easterling EJ, Durbin JD, Hanson SL, Wainner RS, et al. Comparison of short-term response to two spinal manipulation techniques for patients with low back pain in a military beneficiary population. *Military medicine* 2009; 174: 750-756.
- 35. Jüni P, Battaglia M, Nüesch E, Hämmerle G, Eser P, Beers R, et al. A randomised controlled trial of spinal manipulative therapy in acute low back pain. *Annals of the rheumatic diseases* 2009; 68: 1420-1427.
- 36. Machado LA, Maher CG, Herbert RD, Clare H, McAuley JH. The effectiveness of the McKenzie method in addition to first-line care for acute low back pain: a randomized controlled trial. *BMC Med* 2010; 8: 10.
- 37. Lewis C, Souvlis T, Sterling M. Strain-Counterstrain therapy combined with exercise is not more effective than exercise alone on pain and disability in people with acute low back pain: a randomised trial. *Journal of physiotherapy* 2011; 57: 91-98.
- 38. Biondi D, Xiang J, Benson C, Etropolski M, Moskovitz B, Rauschkolb C. Tapentadol immediate release versus oxycodone immediate release for treatment of acute low back pain. *Pain physician* 2013; 16: E237-E246.
- 39. Goertz CM, Long CR, Hondras MA, Petri R, Delgado R, Lawrence DJ, et al. Adding chiropractic manipulative therapy to standard medical care for patients with acute low back pain: results of a pragmatic randomized comparative effectiveness study. *Spine* 2013; 38: 627-634.
- 40. Behrbalk E, Halpern P, Boszczyk BM, Parks RM, Chechik O, Rosen N, et al. Anxiolytic medication as an adjunct to morphine analgesia for acute low back pain management in the emergency department: A prospective randomized trial. *Spine* 2014; 39: 17-22.
- 41. Eken C, Serinken M, Elicabuk H, Uyanik E, Erdal M. Intravenous paracetamol versus dexketoprofen versus morphine in acute mechanical low back pain in the emergency department: A randomised double-blind controlled trial. *Emergency medicine journal* 2014; 31: 177-181.
- 42. Tanen DA, Shimada M, Danish DC, Santos FD, Makela M, Riffenburgh RH. Intravenous lidocaine for the Emergency Department treatment of acute radicular low back pain, a randomized controlled trial. *Journal of emergency medicine* 2014; 47: 119-124.

Table S3: Sensitivity analysis of factors associated with time to claim closure, entering chiropractic care and physiotherapy as time-dependant co-variates (n=1,442)

	Univariat	:e	Multivariable	
	Hazard Ratio (99%	p-value	Adjusted Hazard Ratio (99%	p-value
	CI)		CI)	
Baseline predictors				
Age categories in years		<0.001		<0.001
15 to <25	1.00		1.00	
25 to <35	0.88 (0.65, 1.19)		0.79 (0.58, 1.08)	
35 to <45	0.78 (0.59, 1.04)		0.70 (0.52, 0.95)	
45 to <55	0.76 (0.56, 1.02)		0.67 (0.49, 0.91)	
55 to 65	0.56 (0.40, 0.80)		0.52 (0.36, 0.74)	
Gender		0.114		0.451
Females	1.09 (0.95, 1.26)		0.96 (0.82, 1.12)	
Males	1.00		1.00	
First language		0.137		0.327
English	1.00		1.00	
Other	0.83 (0.59, 1.15)		0.88 (0.63, 1.23)	
Pre-disability income in				
dollars per week				
At 30 days:				
≤480	1.00	_	1.00	_
481-694	1.15 (0.91, 1.45)	0.125	1.08 (0.85, 1.37)	0.396
695-920	1.21 (0.96, 1.54)	0.034	1.05 (0.81, 1.36)	0.614
>920	1.20 (0.94, 1.52)	0.050	1.04 (0.80, 1.35)	0.689
At 60 days:	1.20 (0.54, 1.52)	0.030	1.04 (0.80, 1.33)	0.003
480 ≤480	1.00		1.00	
481-694	1.09 (0.89, 1.34)	0.273	1.03 (0.83, 1.27)	0.710
695-920		0.273	0.97, (0.77, 1.22)	0.716
	1.13 (0.92, 1.39)			
>920	1.08 (0.88, 1.33)	0.326	0.94 (0.74, 1.19)	0.477
At 180 days:	4.00		1.00	
≤480	1.00	- 267	1.00	-
481-694	0.89 (0.68, 1.17)	0.267	0.85 (0.65, 1.12)	0.130
695-920	0.84 (0.63, 1.12)	0.115	0.71 (0.52, 0.96)	0.003
>920	0.72 (0.54, 0.97)	0.004	0.61 (0.45, 0.84)	<0.001
At 365 days:				
≤480	1.00	-	1.00	-
481-694	0.65 (0.36, 1.18)	0.064	0.63 (0.35, 1.16)	0.051
695-920	0.53 (0.28, 1.02)	0.012	0.44 (0.23, 0.84)	0.001
>920	0.39 (0.20, 0.75)	<0.001	0.32 (0.16, 0.63)	<0.001
Opioid prescription		<0.001		<0.001
Yes	0.62 (0.48, 0.80)		0.69 (0.53, 0.89)	
No	1.00		1.00	
Prior claims		0.306		0.661
Yes	1.07 (0.91, 1.26)		1.03 (0.87, 1.22)	
No	1.00		1.00	
Union membership		<0.001		0.014
Yes	1.29 (1.11, 1.50)		1.14 (0.96, 1.36)	
No	1.00		1.00	
Missing	1.34 (1.07, 1.68)		1.27 (1.01, 1.60)	
Employer RTW program	, , ,,	<0.001		<0.001
Yes	1.73 (1.43, 2.09)	2.001	1.77 (1.45, 2.18)	0.001

No	1.00		1.00	
Missing	1.17 (0.87, 1.58)		1.18 (0.86, 1.60)	
Doubt Work-relatedness		0.174		0.119
Yes	0.94 (0.76, 1.16)		0.87 (0.70, 1.08)	
No	1.00		1.00	
Missing	1.13 (0.92, 1.39)		1.08 (0.87, 1.33)	
Time-dependent predictors*				
Chiropractic care received	1.11 (0.95, 1.30)	0.096	1.08 (0.91, 1.29)	0.268
after the accident prior to				
claim closure				
Physiotherapy after the	0.96 (0.83, 1.10)	0.420	0.99 (0.85, 1.16)	0.913
accident prior to claim closure				

HR>1 indicates faster claim closure; RTW = return to work

<sup>\*</sup> The time-dependent predictors are "turned on" once the claimant has received their first service after their accident.

Table S4: Search strategy for observational studies exploring the association of early opioid, physiotherapy, or chiropractic care with Workers' Compensation claim duration, for cases of acute low back pain

#### MEDLINE (OvidSP)

1

2

4

5

6 7 8

9

10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26

27 28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52 53

54

55

56

57

58

- 1 exp Whiplash Injuries/
- 2 exp Soft Tissue Injuries/
- 3 repetitive strain injur\$.mp.
- 4 carpal tunnel syndrome.mp.
- 5 exp Cumulative Trauma Disorders/
- 6 exp Back pain/ or exp pain/ or chronic pain.tw.
- 7 exp Anxiety/
- 8 exp Depression/
- 9 exp Neck Pain/
- 10 exp Depressive Disorder/
- 11 exp Back Injuries/
- 12 injured worker\$.mp.
- 13 musculoskeletal injur\$.mp.
- 14 or/1-13
- 15 exp "Wounds and Injuries"/
- 16 Musculoskeletal System/ or Musculoskeletal Diseases/
- 17 15 and 16
- 18 14 or 17
- 19 exp insurance claim reporting/ or exp "insurance claim review"/ or exp insurance, disability/ or insurance, liability/
- 20 Insurance, Accident/
- 21 ((worker\$ or workman\$ or workmen&) adj compensation).mp. or exp Workers' compensation/
- 22 claim.mp.
- 23 claimant.mp.
- 24 or/19-23
- 25 prognosis.mp. or exp Prognosis/
- 26 Time/ or exp Time Factors/
- 27 exp "Recovery of Function"/
- 28 "Severity of Illness Index"/
- 29 exp Trauma Severity Indices/
- 30 (recovery or prognostic).mp.
- 31 or/25-30
- 32 18 and 24 and 31
- 33 exp Disability Evaluation/
- 34 24 and 31 and 33
- 35 exp Occupational Diseases/ or exp Accidents, Occupational/ or (occupational injur: or occupational accident:).mp.
- 36 24 and 31 and 35
- 37 exp Accidents, Traffic/
- 38 24 and 31 and 37
- 39 "Compensation and Redress"/
- 40 18 and 31 and 39

2

4

5

6 7

8

9

10

11 12 13

14 15

16

17

18

19

20 21

22

23

24

25

26 27

28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52

53 54

55

56

57

58

59 60

```
41 exp Work Capacity Evaluation/ or exp workload/ or workload.mp.
42 (18 or 35) and 24 and 41
43 32 or 34 or 36 or 38 or 40 or 42
44 18 or 33
45 31 or 41
46 35 and 44 and 45
47 46 or 43
```

#### EMBASE (OvidSP)

- 1 whiplash injur\$.mp. or exp whiplash injury/
- 2 exp soft tissue injury/
- 3 soft tissue injur\$.mp.
- 4 repetitive strain injur\$.mp.
- 5 carpal tunnel syndrome.mp. or exp carpal tunnel syndrome/
- 6 exp cumulative trauma disorder/
- 7 back pain.mp. or exp backache/
- 8 backpain.mp.
- 9 chronic pain.mp. or exp chronic pain/
- 10 exp pain/
- 11 anxiety/
- 12 exp depression/
- 13 neck pain.mp. or exp neck pain/
- 14 back injur\$.mp.
- 15 low back injury/
- 16 injured worker\$.mp.
- 17 musculoskeletal injury/
- 18 occupational injuries.mp. or exp occupational accident/
- 19 occupational accidents.mp.
- 20 occupational diseases.mp. or exp occupational disease/
- 21 or/1-20
- 22 insurance/ or exp compensation/ or exp workman compensation/ or exp health insurance/ or exp "health plan employer data and information set"/
- 23 accident insurance.mp.
- 24 exp workman compensation/
- 25 ((worker\$ or workman\$ or workmen&) adj compensation).mp.
- 26 (claim or claimant).mp.
- 27 or/22-26
- 28 prognosis.mp. or prognosis/
- 29 exp time/
- 30 recovery of function.mp. or convalescence/
- 31 disease severity/
- 32 exp injury scale/
- 33 (recovery or prognostic).mp.
- 34 workload.mp. or exp workload/
- 35 exp work capacity/
- 36 exp work resumption/ or return to work.mp.
- 37 or/28-36

```
38 21 and 27 and 37
```

39 or/1-17

1

2

4

5

6 7

8

9 10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26 27

28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46 47

48

49

50

51

52 53

54

55

56

57

58

59 60

- 40 or/18-20
- 41 37 and 39 and 40
- 42 38 or 41

#### PsycInfo (OvidSP)

- 1 exp Whiplash/
- 2 whiplash injur:.mp.
- 3 soft tissue injur\$.mp.
- 4 cumulative trauma disorder\$.mp.
- 5 repetitive strain injur\$.mp.
- 6 carpal tunnel syndrome.mp.
- 7 back pain.mp. or exp Back Pain/
- 8 (backpain or backache).mp.
- 9 chronic pain.mp. or exp Chronic Pain/
- 10 exp Musculoskeletal Disorders/ or exp Fibromyalgia/ or fibromyalgia.mp.
- 11 exp Anxiety/
- 12 exp "Depression (Emotion)"/ or exp Major Depression/
- 13 neck pain.mp.
- 14 back injur:.mp.
- 15 musculoskeletal injur\$.mp.
- 16 exp Industrial Accidents/
- 17 exp Occupational Safety/
- 18 (occupational injur: or occupational accident:).mp.
- 19 exp Work Related Illnesses/
- 20 or/1-19
- 21 exp Workers' Compensation Insurance/
- 22 exp Employee Health Insurance/
- 23 exp Insurance/
- 24 disability insurance.mp.
- 25 (claim or claimant).mp.
- 26 ((worker: or workman: or workmen:) adj compensation).mp.
- 27 accident insurance.mp.
- 28 or/21-27
- 29 prognosis.mp. or exp Prognosis/
- 30 exp Time/
- 31 time factors.mp.
- 32 exp "Recovery (Disorders)"/
- 33 recovery of function.mp.
- 34 exp "Severity (Disorders)"/ or severity of illness.mp.
- 35 (recovery or prognostic).mp.
- 36 exp Work Load/ or workload.mp. or exp Job Performance/
- 37 exp Vocational Evaluation/ or exp Disability Evaluation/ or work capacity evaluation.mp. or exp Reemployment/
- 38 work resumption.mp.
- 39 or/29-38
- 40 20 and 28 and 39

4

5

6

7 8

9 10

11

12

13

14 15

16

17

18

19

20 21

22

23

24

25

26

27 28

29

30

31

32

33 34

35

36

37

38

39 40

41

42

43

44

45

46

47 48

49

50

51

52

53 54

55

56

57

58

```
41 16 or 17 or 18 or 19
42 or/1-15
43 39 and 41 and 42
44 28 and 41
45 40 or 43 or 44
CINAHL (Ebsco)
49
    S46 or S48
S48 S22 and S45 and S47
S47 S23 or S24 or S25 or S26
S46 S27 and S33 and S45
S45 S34 or S35 or S36 or S37 or S38 or S39 or S40 or S41 or S42 or S43 or S44
S44 (MH "Disability Evaluation+")
S43 "work resumption"
S42 "return to work" OR (MH "Job Re-Entry")
S41 (MH "Work Capacity Evaluation")
S40 (MH "Workload Measurement") OR (MH "Workload") OR "workload"
S39 (recovery or prognostic)
S38 (MH "Severity of Illness") OR (MH "Severity of Illness Indices+")
S37 (MH "Recovery")
S36 "recovery of function"
S35 (MH "Time+") OR (MH "Time Factors")
S34 (MH "Prognosis+") OR "prognosis"
S33 S28 or S29 or S30 or S31 or S32
S32 "accident insurance"
S31 (worker* N2 compensation) OR (workman* N2 compensation) OR (workmen* N2
compensation)
S30 (Claim or claimant)
S29 (MH "Insurance") OR (MH "Insurance, Disability+")
S28 (MH "Worker's Compensation")
S27 S22 or S23 or S24 or S25 or S26
S26 "occupational accident" OR (MH "Accidents, Occupational+")
S25 "occupational inju*"
S24 (MH "Occupational-Related Injuries")
S23 (MH "Occupational Diseases+")
S22 S18 or S21
S21 S19 and S20
S20 (MH "Wounds and Injuries+") OR (MH "Occupational-Related Injuries")
S19 (MH "Musculoskeletal Diseases+") or (MH "Musculoskeletal System+")
S18 S1 or S2 or S3 or S4 or S5 or S6 or S7 or S8 or S9 or S10 or S11 or S12 or S13 or S14 or
S15 or S16 or S17
S17 "musculoskeletal injur*"
S16 "back injur*"
S15 (MH "Back Injuries+")
S14 (MH "Neck Pain") OR "neck pain"
S13 (MH "Depression+")
S12 (MH "Anxiety+")
S11 (MH "Pain+")
```

- S10 (MH "Fibromyalgia") OR "fibromyalgia"
- S9 (MH "Chronic Pain") OR "chronic pain"
- S8 (backpain or backache)

- S7 (MH "Back Pain") OR "back pain"
- S6 (MH "Carpal Tunnel Syndrome") OR "carpal tunnel syndrome"
- S5 "repetitive strain injur\*"
- S4 (MH "Cumulative Trauma Disorders+")
- S3 "soft tissue injur\*"
- S2 (MH "Soft Tissue Injuries")
- S1 "whiplash injur\*" OR (MH "Whiplash Injuries")

Figure S1: Flow diagram of the literature search process for randomized controlled trials assessing the effect of opioids, physiotherapy, or chiropractic care for acute low back pain.

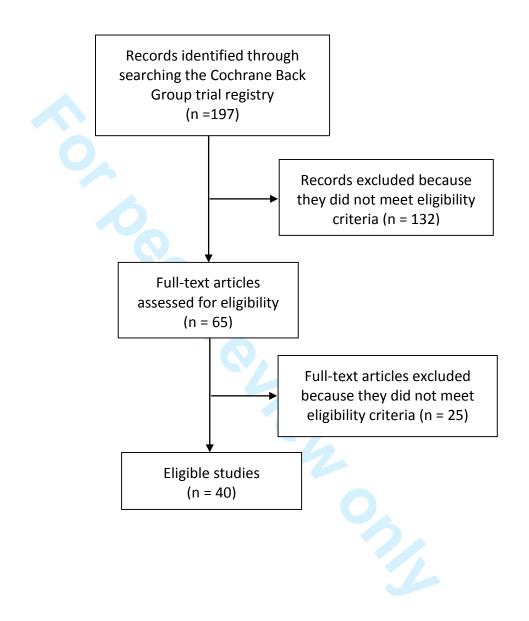
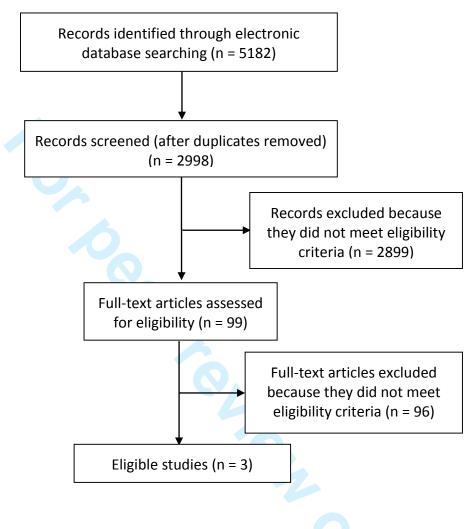


Figure S2: Flow diagram of the literature search process for observational studies assessing the effect of early opioids, physiotherapy, or chiropractic care for acute low back pain.



STROBE Statement—Checklist of items that should be included in reports of *cohort studies* 

	Item No	Recommendation
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the
		abstract (in title on title page)
		(b) Provide in the abstract an informative and balanced summary of what
		was done and what was found (in Abstract)
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being
		reported (page 1 & 2 in Manuscript)
Objectives	3	State specific objectives, including any prespecified hypotheses (Appendix
J		A)
Methods		,
Study design	4	Present key elements of study design early in the paper (In Abstract, and
area y area y		page 3-7 in Manuscript)
Setting	5	Describe the setting, locations, and relevant dates, including periods of
		recruitment, exposure, follow-up, and data collection (page 3 in
		Manuscript)
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection of
1		participants. Describe methods of follow-up (pages 3 & 4 in Manuscript)
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed (not applicable)
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders,
variables	,	and effect modifiers. Give diagnostic criteria, if applicable ( <b>Appendix A</b>
		and pages 3 & 4 in Manuscript)
Data sources/ measurement	8*	For each variable of interest, give sources of data and details of methods of
Data sources/ measurement	O	assessment (measurement). Describe comparability of assessment methods
		if there is more than one group (pages 3-5 in Manuscript)
Bias	9	Describe any efforts to address potential sources of bias (pages 4-7 in
Dias	9	Manuscript)
Study size	10	Explain how the study size was arrived at (page 3 & 4 in Manuscript)
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If
Quantitudi ( variation )		applicable, describe which groupings were chosen and why (pages 5 & 6 in
		Manuscript)
Statistical methods	12	(a) Describe all statistical methods, including those used to control for
Statistical inclinate	12	confounding (pages 3-7 in Manuscript)
		(b) Describe any methods used to examine subgroups and interactions (page
		6 in Manuscript)
		(c) Explain how missing data were addressed (page 5 in Manuscript)
		(d) If applicable, explain how loss to follow-up was addressed ( <b>not</b>
		applicable)
		(e) Describe any sensitivity analyses (not applicable)
Doculto		( <u></u>
Results Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers
		potentially eligible, examined for eligibility, confirmed eligible, included in
		the study, completing follow-up, and analysed ( <b>Table 1 and page 8 in</b>
		Manuscript)
		(b) Give reasons for non-participation at each stage ( <b>not applicable</b> )
		(1) 21.1 reasons for non-participation at each stage (not applicable)

Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders ( <b>Table 1</b> )
		(b) Indicate number of participants with missing data for each variable of interest ( <b>Table 1</b> )
		(c) Summarise follow-up time (eg, average and total amount) (Figure 1)
Outcome data	15*	Report numbers of outcome events or summary measures over time (page 8 in Manuscript)
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted estimates and their precision (eg, 95% confidence interval). Make clear which confounders were adjusted for and why they were included ( <b>Table 2</b> )
		(b) Report category boundaries when continuous variables were categorized (Table 2)
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period ( <b>not relevant</b> )
Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses (page 8 in Manuscript)
Discussion		, , , , , , , , , , , , , , , , , , , ,
Key results	18	Summarise key results with reference to study objectives (page 10 in Manuscript)
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias (pages 10 & 11 in Manuscript)
Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence (page 13 in Manuscript)
Generalisability	21	Discuss the generalisability (external validity) of the study results (page 11 in Manuscript)
Other information		
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based (page 14 in Manuscript)

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at http://www.strobe-statement.org.