

# Accepted Manuscript



Work limitations four years following mild traumatic brain injury: A Cohort Study

Alice Theadom, PhD, Suzanne Barker-Collo, PhD, Kelly Jones, PhD, Michael Kahan, MD, Braden Te Ao, PhD, Kathryn McPherson, PhD, Nicola Starkey, PhD, Valery Feigin, PhD

PII: S0003-9993(17)30067-9

DOI: [10.1016/j.apmr.2017.01.010](https://doi.org/10.1016/j.apmr.2017.01.010)

Reference: YAPMR 56790

To appear in: *ARCHIVES OF PHYSICAL MEDICINE AND REHABILITATION*

Received Date: 27 October 2016

Revised Date: 9 December 2016

Accepted Date: 6 January 2017

Please cite this article as: Theadom A, Barker-Collo S, Jones K, Kahan M, Te Ao B, McPherson K, Starkey N, Feigin V, on behalf of the BIONIC4you Research Group, Work limitations four years following mild traumatic brain injury: A Cohort Study, *ARCHIVES OF PHYSICAL MEDICINE AND REHABILITATION* (2017), doi: 10.1016/j.apmr.2017.01.010.

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

**Title: Work limitations four years following mild traumatic brain injury: A Cohort Study**

**Running head: Mild TBI and long-term work**

**Alice Theadom PhD,<sup>1</sup> Suzanne Barker-Collo PhD<sup>2</sup>, Kelly Jones PhD<sup>1</sup>, Michael Kahan MD<sup>3</sup>, Braden Te Ao PhD<sup>1</sup>, Kathryn McPherson PhD<sup>4</sup>, Nicola Starkey PhD<sup>5</sup>, Valery Feigin PhD<sup>1</sup> on behalf of the BIONIC4you Research Group\***

<sup>1</sup> National Institute for Stroke and Applied Neuroscience, School of Public health & Psychosocial Studies, Auckland University of Technology, New Zealand

<sup>2</sup> School of Psychology, University of Auckland, Auckland, New Zealand

<sup>3</sup> Waikato Occupational Services; Hamilton, New Zealand.

<sup>4</sup> Health Research Council of New Zealand, Auckland, New Zealand

<sup>5</sup> School of Psychology, University of Waikato, Hamilton, New Zealand

**Corresponding Author:**

Dr Alice Theadom  
National Institute for Stroke and Applied Neuroscience  
School of Rehabilitation and Occupation Studies  
Faculty of Health and Environmental Sciences  
AUT North Shore Campus, AA254C  
90 Akoranga Drive, Private Bag 92006  
Auckland 1142, New Zealand  
Telephone: +64 9 921 9999 x7805  
Fax: +64 9 921 9620  
Email: [alice.theadom@aut.ac.nz](mailto:alice.theadom@aut.ac.nz)

**DECLARATION OF COMPETING INTERESTS**

Alice Theadom was co-funded by ABI Management Rehabilitation. Kathryn McPherson was appointed as the Chief Executive of the Health Research Council in 2015 but was neither in post nor involved in the funding decisions for this study. The authors declare no conflicts of interest.

## **FUNDING**

This work was funded by the Health Research Council of New Zealand (09/063A, 11/192, 13/408)

## **ETHICAL APPROVAL**

Ethical approval was obtained from the Northern Y Health and Disability Ethics Committee of New Zealand (NTY/09/09/095) and the Auckland University of Technology Ethics Committee (09/265).

# 1      **Work limitations four years following mild traumatic brain injury: A cohort study**

2

## 3      **Abstract**

4

5

6      *Objective:* To explore employment status, work limitations and productivity loss following  
7      mild traumatic brain injury (TBI).

8      *Study Design:* Inception cohort study over four years.

9      *Setting:* General community.

10     *Participants:* 245 Adults (>16 years at the time of injury), who experienced a mild TBI and  
11     who were employed prior to their injury.

12     *Interventions:* Not applicable

13     *Main Outcome Measures:* Details of the injury, demographic information and pre-injury  
14     employment status were collected from medical records and self-report. Symptoms and  
15     mood were assessed one-month post-injury using the Rivermead Post-Concussion Symptom  
16     Questionnaire and the Hospital Anxiety and Depression Scale. Post-injury employment  
17     status and work productivity were assessed four-years post-injury using the Work  
18     Limitations Questionnaire.

19     *Results:* Four-years following mild TBI, 17.3% of participants had exited the workforce  
20     (other than for reasons of retirement or to study) or had reduced their working hours  
21     compared to pre-injury. A further 15.5% reported experiencing limitations at work as a  
22     result of their injury. Average work productivity loss was 3.6% The symptom of ‘taking

23 longer to think' one month post-injury significantly predicted work productivity loss four  
24 years later ( $\beta = 0.47$ ,  $t = 3.79$ ,  $p = <0.001$ ).

25

26 *Conclusions:* Whilst changes in employment status and difficulties at work are likely over  
27 time, the results indicate increased unemployment rates, work limitations and productivity  
28 loss in the longer-term following a mild TBI. Identification of cognitive difficulties one  
29 month following TBI in working aged adults and subsequent interventions to address these  
30 difficulties are required to facilitate work productivity.

31

32 Keywords: Mild Traumatic Brain Injury, Work, Employment, Longitudinal studies,  
33 Productivity

34

35 Abbreviations:

36

37 TBI = Traumatic brain injury

38 WLQ-25 = Work limitations Questionnaire

39 HADS = Hospital Anxiety and Depression Scale

40 RPQ = Rivermead Post Concussion Symptom Questionnaire

41

42

43

44 The majority of traumatic brain injuries (TBIs) are classified as being in the mild severity  
45 range (70-95%),<sup>1,2</sup> with 63% of mild TBIs occurring in adults of working age (between 16-  
46 64years).<sup>2</sup> Following a mild TBI the most common symptoms include headaches, fatigue,  
47 forgetfulness and poor concentration and taking longer to think<sup>3</sup>, all of which can affect an  
48 individual's ability to function in everyday life, including productivity at work. Whilst  
49 symptoms may resolve quickly in many cases, estimates suggest between 22% and 48% of  
50 those sustaining a mild TBI experience persistent symptoms (including headaches, fatigue  
51 and difficulties processing information) and functional losses for at least one year following  
52 injury.<sup>3,4</sup> However, further work is needed to understand the longer-term consequences of  
53 mild TBI on occupational functioning.<sup>5</sup>

54

55 Return to work is increasingly identified as a key outcome in TBI, being linked to improved  
56 psychological well-being and quality of life.<sup>6,7</sup> Most adults (90%) return to pre-injury  
57 employment within two months of sustaining a mild TBI.<sup>8</sup> Delayed return to work after  
58 injury results in considerable lost earnings and long-term treatment costs.<sup>9</sup> Thus, there is a  
59 substantial economic incentive to facilitate return to work given the disproportionate risk of  
60 TBI in younger adults.<sup>2</sup> Studies have revealed that poorer mood, reduced global functioning,  
61 lower level of education, acute symptoms of nausea, vomiting and dizziness, extracranial  
62 injuries, cognitive impairment, occupational factors such as limited independence, increased  
63 age, multiple injuries and fatigue all predict delayed return to work following mild TBI.<sup>8,10</sup>  
64 <sup>11</sup> However, studies have been limited by small sample sizes and short follow up periods.<sup>10</sup>  
65 Consequently, it remains unknown if mild TBI has an impact on employment status in the  
66 longer term. Additionally, the current focus of research on return to work rates following

67 mild TBI has failed to identify the impact of mild TBI on work productivity or changes an  
68 individual may have needed to make to remain in their employment, and/or whether mild  
69 TBI increases absence from work. These are all key factors in a person's ability to sustain  
70 employment in the longer term.<sup>10, 12</sup> Work productivity is defined as how efficiently people  
71 perceived that they are performing in their employment including managing their time,  
72 coping with physical demands, relationships with colleagues and clients as well as the  
73 quality and quantity of their work).<sup>13</sup> It remains unclear if factors that are predictive of  
74 return to work also predict work productivity.

75

76 The potential impact of prior and recurrent brain injuries on work productivity has also not  
77 been explored. This is particularly pertinent given 32% of people have experienced at least  
78 one prior TBI before the age of 25 years<sup>14</sup> and 10% of people experience a recurrent injury  
79 within a year of an initial injury, which can exacerbate symptoms and negatively impact  
80 recovery.<sup>15</sup> This study aimed to explore employment status, work productivity and  
81 productivity loss four years after mild TBI and to identify acute predictors of these  
82 outcomes. It was hypothesised that lower age, acute symptoms and prior-recurrent injuries  
83 would be predictive of longer-term work productivity.

84

## 85 **METHODS**

86

87

### 88 **Participants**

89

90 According to government statistics, in New Zealand (NZ), only one in ten people go on to  
91 further study (including both degree and non-degree qualifications) and so many people

92 enter the workforce at the age of 16 years.<sup>16</sup> Adults (aged >16 years) who experienced a  
93 mild TBI during a 12 month period were identified through a longitudinal incidence and  
94 outcome study (*Brain Injury Incidence and Outcomes New Zealand in the Community*,  
95 BIONIC).<sup>17</sup> The BIONIC study identified 1369 cases (including all ages and severities) of  
96 TBI between 1 March 2010 through 28 February 2011 in a mixed urban and rural region  
97 (Hamilton and Waikato Districts) of the North Island of New Zealand (NZ). Multiple  
98 overlapping sources of case ascertainment were employed including searches of sports club  
99 accident records, general practitioners and allied health professional referrals and self-  
100 referrals, in addition to searches of hospital admission and discharge records and national  
101 health care databases. TBI was defined using the World Health Organisation criteria as: an  
102 acute brain injury resulting from mechanical energy to the head from external physical  
103 forces.<sup>18</sup> Information on all potential TBI cases based on self-report and information  
104 obtained from medical records was reviewed weekly by a diagnostic adjudication group to  
105 determine if they met the inclusion criteria for TBI. Mild TBI severity was defined using  
106 Glasgow Coma Score (GCS; 13-15) and/or Post-traumatic amnesia (< 24 hours). All cases  
107 meeting the TBI inclusion criteria that did not have a recorded GCS score were classified as  
108 mild in severity.

109

## 110 **Procedure**

111

112 Participant demographic and clinical information about the person's TBI history (including  
113 injuries sustained prior to the registering TBI identified as part of the incidence study and  
114 recurrent injuries sustained over the subsequent four year period of follow up) were  
115 obtained from medical records and self-report (for consenting participants) at baseline.

116

117 All confirmed TBI cases were invited to participate in follow-up assessments that were  
118 completed in person at the participant's place of residence or at another mutually convenient  
119 location (e.g., private room at medical practice). Participants were also offered the option to  
120 complete the assessment via the telephone if they preferred. Participants provided written  
121 consent to participate all time-points. Assessments used in the current analysis were  
122 completed at one month and four years post-injury. Participants were eligible for inclusion  
123 in this analysis if they were: 1) aged >16 years at the time of injury; 2) had experienced a  
124 mild TBI within the recruitment period; 3) were in full or part-time paid employment pre-  
125 injury. The participant's occupation was classified into the 8 main job categories outlined by  
126 the Australian and New Zealand Standard Classification of Occupations (2013 Version  
127 1.2).<sup>19</sup> As the aim of this study was to identify the acute factors that may predict longer term  
128 work status and productivity (to indicate targets for early intervention), potential predictors  
129 (mood and symptoms) were assessed one month post-injury, with work productivity  
130 assessed at four years post-injury.

131

### 132 **Assessments at one month post-injury**

133

134 The *Rivermead Post Concussion Symptoms Questionnaire*<sup>20</sup> (RPQ) was specifically  
135 developed to assess the experience of symptoms after a brain injury in comparison to before  
136 the injury. Participants are presented with a list of 16 cognitive, emotional and physical  
137 symptoms commonly experienced after a brain injury including headaches, noise sensitivity  
138 and balance difficulties. Participants were asked to rate on a 5-point scale, ranging from the  
139 symptom is not experienced (0) to it being a severe problem (4). To explore the role of

140 specific acute symptoms on work productivity, individual item (i.e., symptom) level scores  
141 were used for this analysis.

142

143 As mood has been found to influence return to work following injury<sup>11</sup>, the *Hospital Anxiety*  
144 *and Depression Scale (HADS)*<sup>21</sup> was used to assess levels of depression and anxiety. The scale  
145 was developed specifically to assess mood in people with physical health conditions and has  
146 been found to be sensitive to changes in anxiety and depression following TBI.<sup>22</sup> High  
147 scores (range 0-21) indicate poorer mood status.

148

#### 149 **Outcome measures 4 years post-injury**

150

151 In addition to the measures collected up to one year as part of the BIONIC study,  
152 participants employed four years post-injury were asked to complete the *Work Limitations*  
153 *Questionnaire (WLQ-25)*.<sup>23, 24</sup> This questionnaire was developed to assess the extent to  
154 which a person experiences limitations at work and lost productivity due to their health.  
155 Participants are asked to respond on a 5-point scale about the level of difficulty that they  
156 have had in completing specific job tasks over the previous two weeks. Responses are  
157 summed to yield four subscale scores for time management; physical demands; mental-  
158 interpersonal demands; and an outputs demands scale assessing diminished work quality  
159 and quantity. Higher scores indicate poorer work productivity. In accordance with the  
160 scoring instructions of the measure, an algorithm was used to convert the four sub-scale  
161 scores into an estimate of total percentage of productivity loss. Two additional time loss  
162 questions were included which asked how many half and full days the person had taken off  
163 work in the past two weeks. The questionnaire has well-established criterion and construct

164 validity and reliability.<sup>13, 24</sup> Additionally, an open text question asked participants if they had  
165 needed to make any changes within their employment as a result of their brain injury.

166

### 167 **Statistical analysis**

168

169 Descriptive statistics were used to examine the participant sample, changes in employment  
170 status between pre-injury and four year follow up, changes required to remain in  
171 employment and numbers of days taken off work in the past two weeks. Chi square or t-tests  
172 were used to determine any differences between participants consenting to take part in the  
173 four year follow up and those who did not. Spearman's correlation co-efficients were used  
174 to explore demographic, injury related and symptom variables associated with work  
175 productivity four years post-injury. The significance level was set at the more stringent level  
176 of  $p < 0.01$  to account for multiple comparisons. Acute factors that were significantly  
177 correlated with work productivity and which were not multi-collinear ( $r = > 0.8$ ), were  
178 entered into a multiple linear regression to determine their unique contribution. Forward  
179 conditional selection was used to develop the regression model. Predictors were retained in  
180 each model if the p-value was  $\leq 0.05$ .

181

## 182 **RESULTS**

183

184

185 There were 245 adults (>16 years) who met the inclusion criteria for this analysis. At four  
186 year follow up two (0.8%) participants had died and 50 (20.4%) were unable to be  
187 contacted. Of the remaining 193 participants, 110 (57.0%) consented to the four year follow  
188 up assessment, 31 (12.7%) declined, 19 (7.8%) initially agreed but were then unable to be

189 contacted and 33 (13.5%) had declined follow up at a previous assessment. Characteristics  
190 of the sample are presented in Table 1 alongside a comparison with those who were eligible  
191 for this analysis but who did not consent to follow up to explore risk of selection and  
192 attrition bias. As can be seen in Table 1, there were no significant differences on most  
193 clinical and demographic variables with the exception that participants in this analysis were  
194 older than those who did not take part. The majority of assessments were completed face-to-  
195 face with the person in their own home (96.6%), with only a few people preferring to  
196 complete the assessment via telephone interview (3.4%). As there were minimal missing  
197 data due to the majority of assessments being conducted by the researcher in person,  
198 participants with missing data were excluded from the analysis. Prior to their injury, 11.8%  
199 of the sample were in managerial roles, 15.5% professionals, 10.9% Technician/Trade  
200 workers, 12.7% community workers, 10.9% clerical or administrative workers, 8.2% sales  
201 workers, 10.0% machinery operators, 13.6% labourers and 6.4% who did not specify their  
202 pre-injury occupation.

203

**204 INSERT TABLE 1**

205

206 As shown in Table 2, 19 (17.3%) of participants had exited their employment at four years  
207 post-injury for reasons other than retirement or study. Rates of unemployment were slightly  
208 higher (7.0%) in our mild TBI sample four years post-injury than reported in the New  
209 Zealand general population in the same year of follow up (5.8% in 2015).<sup>25</sup> Additionally,  
210 17 (15.5%) people reported needing to make changes to enable them to continue in their  
211 employment role because of the impact of their brain injury. The most common changes  
212 included becoming more reliant on note taking and allowing themselves more time to  
213 process information, increasing their sleep/rest/breaks to assist in managing fatigue and

214 reducing working hours. A greater proportion of labourers exited the workforce, than  
215 remained in their employment, which neared statistical significance  $X^2 = 3.31$ ,  $p = 0.07$ .  
216 There were no observable differences in employment status at four years within the other  
217 occupation categories. Of those who had left the workforce four years post-injury, 45.8%  
218 were under the age of 35 years, with 63.6% of these participants becoming unemployed.

219  
220 **INSERT TABLE 2**

221

222 There were  $N = 86$  participants who completed the WLQ-25 four years post-injury. The four  
223 sub-scales of the WLQ-25 demonstrated good internal consistency for this mild TBI sample,  
224 with Cronbach alpha scores ranging between 0.84 and 0.92. To provide context for the  
225 scores on the WLQ-25, NZ normative data matched for age, gender, education level and  
226 ethnicity were supplied by the developers of the WLQ-25. As can be seen in figure 1, the  
227 mild TBI sample experienced far greater difficulties at work four years post-injury than  
228 reported in the NZ general population.

229

230 **INSERT FIGURE 1**

231

232 **Figure 1. Comparison of mean scores between the mild TBI sample four years post-**  
233 **injury and NZ norms**

234

235

236 Participants who were employed lost on average half a day off work in the past two weeks  
237 due to health or medical care (although this was not directly specified as due to their TBI),  
238 with 15% of people having taken two days off work or more. Work productivity loss ranged  
239 between 0 – 18%, with a mean productivity loss score (3.6%) that was higher than the NZ

240 norm of 2.3%. (Figure 1). Adults who had experienced a mild TBI had particular difficulties  
241 with losing their train of thought, concentration and sticking to a routine. One in six reported  
242 having to make changes to their working style to assist them in managing persistent  
243 symptoms at work.

244

245 To explore the acute factors associated with longer-term work productivity, baseline socio-  
246 demographic factors and injury characteristics, self-reported symptoms (RPQ items) and  
247 mood assessed at one month, were correlated with the sub-scales of the work limitations  
248 questionnaire assessed at four years. As shown in Table 3, the symptom of taking longer to  
249 think at one month was significantly correlated with all work productivity subscale scores  
250 and work productivity loss. Symptoms of fatigue, irritability and poor concentration were  
251 also related to more than three of the five WLQ outcome domains. Demographic factors  
252 were not correlated with the subscales of the WLQ, with prior and recurrent TBI being the  
253 only injury characteristics associated with work limitations four years post-injury.

254

255 **INSERT TABLE 3**

256

257 Variables significantly correlated with each work productivity outcome were entered into a  
258 regression model to explore their unique contribution to aspects of work productivity.

259 Taking longer to think at one month was independently predictive of work productivity loss,  
260 physical demands and, output demands at four years. Poor concentration was significantly  
261 associated with time management difficulties and mental and interpersonal concerns at four  
262 years (Table 4).

263

264 **INSERT TABLE 4**

265

266 **DISCUSSION**

267

268

269 This study highlights that whilst previous research has shown high return to work rates  
270 following a mild TBI in the short term, nearly one in five people struggle to meet the  
271 demands of their employment in the longer term and may go on to exit employment or  
272 reduce their hours. Those working as labourers may be at particular risk of maintaining  
273 employment post-injury. The findings highlight that cognitive impairments, particularly  
274 taking longer to think and poor concentration observed one month following injury, may  
275 help to predict those who will go on to experience difficulties in their employment and who  
276 may benefit from early intervention to help reduce the likelihood and impact of mild TBI on  
277 longer term work limitations and productivity loss.

278

279 Over the period of the study, unemployment in NZ remained relatively stable, declining by  
280 0.7% between 2010 when TBI cases were identified and recruited and 2015 when the four  
281 year follow up assessments were conducted. Given that the majority of people (91.7%)  
282 return to work within two months following a mild TBI, reductions in the employment rate  
283 were not expected.<sup>8</sup> Whilst nearly half of the sample did not experience any difficulties at  
284 work following their injury, this study has clearly highlighted that more than half of adults  
285 experiencing a mild TBI experience challenges in their workplace. These levels are higher  
286 compared with findings from other clinical populations, such as colorectal cancer patients,  
287 where only 22-39% of patients report experiencing work difficulties.<sup>26</sup>

288

289 A work productivity loss of 3.6% as assessed by the work limitations questionnaire was  
290 similar to the 3.0% long-term productivity loss costs based on health economic projections  
291 for TBI.<sup>27</sup> This level of productivity loss is higher than has been reported for colorectal  
292 cancer patients of 1.9%,<sup>26</sup> but lower than reported for psoriatic and rheumatoid arthritis<sup>28, 29</sup>  
293 and systematic vasculitis<sup>30</sup> patients with reported productivity losses of 5.9%, 7.1% and  
294 8.2% respectively. These findings provide more evidence of the possible persistent nature of  
295 symptoms following mild TBI, and specifically the implications of symptoms on  
296 employment and work productivity<sup>3, 4</sup>

297

298 One month symptoms of taking longer to think and poor concentration one month post-  
299 injury were significantly correlated with sub-scale work limitations and productivity scores  
300 at four years. As the reliability and validity of individual items of the post-concussion  
301 symptoms scale have not been demonstrated, these findings should be interpreted with  
302 caution, however the findings indicate an important trend in need of further investigation.  
303 This study indicates that screening, to see if these particular symptoms remain present one  
304 month after injury, may help to identify those at risk of developing longer term difficulties  
305 at work. It was interesting to note that prior and recurrent TBIs were the only injury  
306 characteristics that were significantly correlated with longer term work limitations and  
307 productivity loss. This suggests that symptoms may be more informative in identifying  
308 those who may be more at risk of longer term difficulties. Whilst a high proportion of those  
309 exiting employment following injury were aged less than 35 years, age was not an  
310 independent predictor of longer term work productivity. It may be the case that length of  
311 time in the job or whether the role is part of a planned career path may have more of  
312 influence on employment status and productivity. Further exploration of potential job and

313 contextual factors may be useful to understand the links between mild TBI and work  
314 productivity.

315

### 316 **Study Limitations**

317

318 This study was based on an incidence cohort of people who sustained an injury within a one  
319 year period and took advantage of the multiple sources of cases ascertainment used to  
320 identify those often missed in TBI research (e.g. those who do not go the hospital following  
321 their injury). However, it is acknowledged that participants in this sample were older in age  
322 than those with mild TBI in general, and only included those employed at the time of injury,  
323 limiting generalizability of the findings. Whilst one in six participants did report making  
324 direct changes to their working style as a result of their injury, it is not clear what changes to  
325 employment status and work productivity were directly the result of TBI. For example,  
326 some work difficulties may have been due to additional injuries sustained at the time of the  
327 accident, comorbid medical conditions or changing life circumstances as a result of external  
328 factors. Although data on employment status at time of injury were available, the impact of  
329 recall bias meant the person's level of pre-work functioning was not able to be reliably  
330 obtained. However the findings do suggest a link between symptoms following a mild TBI  
331 and longer term work productivity.

332

333

### 334 **CONCLUSIONS**

335

336

337 Whilst return to work rates following a mild TBI may be high in the short term, nearly one  
338 in five people struggle to meet the demands of their employment in the longer term and may  
339 go on to exit employment or reduce their hours. The findings highlight that cognitive  
340 impairments, particularly taking longer to think and poor concentration observed one month  
341 following injury, may help to predict those who will go on to experience difficulties in their  
342 employment and who may benefit from early intervention to help reduce the impact of mild  
343 TBI on longer term work limitations and productivity loss.

344

345

## 346 **ACKNOWLEDGEMENTS**

347

348

349 We thank the research team and the participants for their contribution to this study and the  
350 many doctors, nurses and rehabilitation professionals who assisted in the long-term follow  
351 up of this sample.

352

## 353 **REFERENCES**

354

- 355 1. Cassidy JD, Carroll LJ, Peloso PM, Borg J, von Holst H, Holm L, Kraus J, Coronado VG,  
356 W. H. O. Collaborating Centre Task Force on Mild Traumatic Brain Injury. Incidence, risk  
357 factors and prevention of mild traumatic brain injury: results of the WHO Collaborating  
358 Centre Task Force on Mild Traumatic Brain Injury. *J Rehab Med.* 2004(43 Suppl):28-60.
- 359 2. Feigin VF, Theadom A, Barker-Collo SL, Starkey N, McPherson K, Kahan M, Dowell A,  
360 Brown P, Parag V, Kydd R, Jones K, Jones A, Ameratunga S, Group obotBS. Incidence of  
361 traumatic brain injury in New Zealand: a population-based study. *Lancet Neurol.*  
362 2013;12(1):53-64.
- 363 3. Theadom A, Parag V, Dowell T, McPherson K, Starkey N, Barker-Collo S, Jones K,  
364 Ameratunga S, Feigin VL, Group BR. Persistent problems 1 year after mild traumatic brain  
365 injury: a longitudinal population study in New Zealand. *Br J Gen Pract.* 2016;66(642):e16-  
366 23.
- 367 4. McMahon P, Hricik A, Yue JK, Puccio AM, Inoue T, Lingsma HF, Beers SR, Gordon WA,  
368 Valadka AB, Manley GT, Okonkwo DO, Casey SS, Cooper SR, Dams-O'Connor K, Menon  
369 DK, Sorani MD, Yuh EL, Mukherjee P, Schnyer DM, Vassar MJ. Symptomatology and

- 370 Functional Outcome in Mild Traumatic Brain Injury: Results from the Prospective TRACK-  
 371 TBI Study. *J Neurotrauma*. 2014;31(1):26-33
- 372 5. Carroll LJ, Cassidy JD, Cancelliere C, Côté P, Hincapié CA, Kristman VL, Holm LW, Borg  
 373 J, Nygren-de Boussard C, Hartvigsen J. Systematic Review of the Prognosis After Mild  
 374 Traumatic Brain Injury in Adults: Cognitive, Psychiatric, and Mortality Outcomes: Results  
 375 of the International Collaboration on Mild Traumatic Brain Injury Prognosis. *Arch Phys  
 376 Med Rehabil*. 2014;95(3):S152-S173.
- 377 6. O'Neill JH, Hibbard MR, Brown M, Jaffe M, Sliwinski M, Vandergoot D, Weiss MJ. The  
 378 effect of employment on quality of life and community integration after traumatic brain  
 379 injury. *J Head Trauma Rehabil*. 1998;13(4):68-79.
- 380 7. Steadman-Pare D, Colantonio A, Ratcliff G, Chase S, Vernich L. Factors associated with  
 381 perceived quality of life many years after traumatic brain injury. *J Head Trauma Rehabil*.  
 382 2001;16(4):330-342.
- 383 8. Waljas M, Iverson GL, Lange RT, Liimatainen S, Hartikainen KM. Return to Work  
 384 Following Mild Traumatic Brain Injury. *J Head Trauma Rehabil*. 2014;29(5):443-450.
- 385 9. Sengupta I, Reno V, Burton JH. Workers' compensation benefits, coverage, and costs, 2008.  
 386 Washington: DC: National Academy of Social Insurance.; 2010.
- 387 10. Cancelliere C, Kristman VL, Cassidy JD, Hincapié CA, Cote P, Boyle E, Carroll LJ,  
 388 Stalnacke BM, Nygren-de Boussard C, Borg J. Systematic review of return to work after  
 389 mild traumatic brain injury: results of the international collaboration on mild traumatic brain  
 390 injury prognosis. *Arch Phys Med Rehabil Nurs*. 2014;95(3):S201-S209.
- 391 11. Vikane E, Hellstrom T, Roe C, Bautz-Holter E, Assmus J, Skouen JS. Predictors for Return  
 392 to Work in Subjects with Mild Traumatic Brain Injury. *Behav Neurol*. 2016;2016:8026414.
- 393 12. Sherer M, Novack TA, Sander AM, Struchen MA, Alderson A, Thompson RN.  
 394 Neuropsychological assessment and employment outcome after traumatic brain injury : a  
 395 review. *Clin Neuropsychol*. 2002;16(2):157-178.
- 396 13. Lerner D, Amick BC, Rogers WH, Malspeis S, Bungay K, Cynn D. The Work Limitations  
 397 Questionnaire. *Med Care*. 2001;39(1):72-85.
- 398 14. McKinlay A, Grace RC, Horwood LJ, Fergusson DM, Ridder EM, MacFarlane MR.  
 399 Prevalence of traumatic brain injury among children, adolescents and young adults:  
 400 prospective evidence from a birth cohort. *Brain Inj*. 2008;22(2):175-181.
- 401 15. Theadom A, Parmar P, Jones K, Barker-Collo S, Starkey NJ, McPherson KM, Ameratunga  
 402 S, Feigin VL, Group BR. Frequency and impact of recurrent traumatic brain injury in a  
 403 population-based sample. *J Neurotrauma*. 2015;32(10):674-681.
- 404 16. New Zealand Government, Ministry of Education.  
 405 [https://www.educationcounts.govt.nz/statistics/indicators/main/student-engagement-  
 406 participation/participation\\_rates\\_in\\_tertiary\\_education](https://www.educationcounts.govt.nz/statistics/indicators/main/student-engagement-participation/participation_rates_in_tertiary_education).
- 407 17. Theadom A, Barker-Collo S, Feigin V, Starkey N, Jones K, Jones A, Ameratunga S, Barber  
 408 P, on behalf of the BIONIC Research Group. The spectrum captured: a methodological  
 409 approach to studying incidence and outcomes of traumatic brain injury on a population  
 410 level. *Neuroepidemiol*. 2012(38(1):18-29.
- 411 18. Carroll LJ, Cassidy JD, Holm L, Kraus J, Coronado VG, WHO Collaborating Centre Task  
 412 Force on Mild Traumatic Brain Injury. Methodological issues and research  
 413 recommendations for mild traumatic brain injury: the WHO Collaborating Centre Task  
 414 Force on Mild Traumatic Brain Injury. *J Rehabil Med*. 2004;43 Suppl:113-125.
- 415 19. Trewin D, Pink B. Australian and New Zealand Standard Classification of Occupations.  
 416 Canberra, Australia: Australian Bureau of Statistics 2006.
- 417 20. King NS, Crawford S, Wenden FJ, Moss NE, Wade DT. The Rivermead Post Concussion  
 418 Symptoms Questionnaire: a measure of symptoms commonly experienced after head injury  
 419 and its reliability. *J Neurol*. 1995;242(9):587-592.
- 420 21. Zigmond AS, Snaith RP. The Hospital Anxiety and Depression Scale. *Acta Psychiatr Scand*.  
 421 1983;67:361-370.
- 422 22. Whelan-Goodinson R, Ponsford J, Schonberger M. Validity of the Hospital Anxiety &  
 423 Depression Scale to assess depression & anxiety following traumatic brain injury as

- 424 compared with the Structured Clinical Interview for DSM-IV. *J Affect Dis.* 2008;114:94-  
425 102.
- 426 **23.** Lerner DJ, Amick BI, Wellcome G. *Work Limitations Questionnaire*. Boston, MA: The  
427 Health Institute, Tufts-New England Medical Center; 1998.
- 428 **24.** Lerner D, Rogers WH, Chang H. The work limitations questionnaire. *Quality of Life.*  
429 2002;28:9-10.
- 430 **25.** Statistics NZ. Unemployment rate table.  
431 [http://www.stats.govt.nz/browse\\_for\\_stats/snapshots-of-nz/nz-progress-](http://www.stats.govt.nz/browse_for_stats/snapshots-of-nz/nz-progress-indicators/home/economic/unemployment-rate.aspx)  
432 [indicators/home/economic/unemployment-rate.aspx](http://www.stats.govt.nz/browse_for_stats/snapshots-of-nz/nz-progress-indicators/home/economic/unemployment-rate.aspx).
- 433 **26.** McGrath C, Mihala G, Beesley VL, Lynch BM, Graves N, Gordon LG. "Cancer Put My  
434 Life on Hold": Work-Related Challenges Among Middle-aged Adults 12 Months After a  
435 Diagnosis of Colorectal Cancer. *Cancer Nursing.* 2016;26(Epub ahead of print).
- 436 **27.** Te Ao B, Brown P, Tobias M, Ameratunga S, Barker-Collo S, Theadom A, McPherson K,  
437 Starkey N, Dowell A, Jones K, Feigin VL, Group. obotBS. Cost of traumatic brain injury in  
438 New Zealand: Evidence from a population-based study. *Neurol.* 2014;83(18):1645-1652.
- 439 **28.** Walsh JA, McFadden ML, Morgan MD, Sawitzke AD, Duffin KC, Krueger GG, Clegg DO.  
440 Work productivity loss and fatigue in psoriatic arthritis. *J Rheumatol.* 2014;41(8):1670-  
441 1674.
- 442 **29.** van Vilsteren M, Boot CR, Knol DL, van Schaardenburg D, Voskuyl AE, Steenbeek R,  
443 Anema JR. Productivity at work and quality of life in patients with rheumatoid arthritis.  
444 *BMC Musculoskel Diss.* 2015;16(107).
- 445 **30.** Barra LJ, Bateman EA, Rohekar S, Pagnoux C, Moradizadeh M. Assessment of work  
446 limitations and disability in systemic vasculitis. *Clin Exp Rheumatol.* 2016;34(3):111-114.  
447  
448

**Table 1. Participant Characteristics of those employed pre-injury consenting to follow up and those who declined or could not be contacted**

	<b>Participant sample at 4 years (N=110)</b>	<b>Non-consenters to 4 year follow up (n = 135 )</b>	<b>Test of Difference</b>
<b>Age, mean (SD)</b>	39.63 (13.47)	34.50 (13.93)	T = 2.92 p = 0.04
<b>Gender</b>			
Male	63 (57.3)	92 (68.1)	X <sup>2</sup> = 2.6 P = 0.10
Female	47 (42.7)	43 (31.9)	
<b>Ethnicity</b>			
European	79 (71.8)	82 (60.7)	X <sup>2</sup> = 4.47 p = 0.11
Maori	24 (21.8)	46 (34.1)	
Other	7 (6.4)	7 (5.2)	
<b>Mechanism of injury</b>			
Falls	40 (36.4)	38 (28.1)	X <sup>2</sup> = 5.75 P = 0.22
Motor vehicle accident	29 (26.4)	35 (25.9)	
Exposure to mechanical force	20 (18.2)	27 (20.0)	
Assault	21 (19.1)	30 (22.2)	
Other/unknown	0 (0.0)	5 (3.7)	
<b>Additional injuries</b>			
Yes	80 (72.7)	103 (76.3)	X <sup>2</sup> = 4.67 p = 0.09
No	21 (19.1)	29 (21.5)	
Not recorded	9 (8.2)	3 (2.2)	
<b>Prior TBI</b>			
None	48 (43.6)	66 (48.9)	X <sup>2</sup> = 2.15 p = 0.54
One	24 (21.8)	23 (17.0)	
Two or more	33 (30.1)	43 (31.9)	
Unknown	5 (4.5)	3 (2.2)	
<b>Recurrent TBI (over 4 year study duration)</b>			
Yes	25 (22.7)	22 (16.3)	X <sup>2</sup> = 1.23 p = 0.27
No	85 (77.3)	113 (83.7)	
<b>Marital Status</b>			
Married/Civil Union/De facto	60 (54.5)	69 (51.1)	X <sup>2</sup> = 2.49 P = 0.29
Never Married/Single	35 (31.8)	54 (40.0)	
Divorced/Separated/Widowed	15 (13.6)	12 (8.9)	
<b>Area of residence</b>			
Urban	85 (77.3)	91 (67.4)	X <sup>2</sup> = 2.45 p = 0.12
Rural	25 (22.7)	44 (32.6)	

**Table 2. Change in Employment Status from Pre-injury to Four Years post-injury**

	<b>N = 110</b>
<b>Exited employment</b>	<b>24 (21.8)</b>
Employed – student	4 (16.2)
Employed – retired	7 (29.2)
Employed – unemployed/beneficiary	12 (50.0)
Employed - homemaker	1 (4.2)
<b>Remained Employed</b>	<b>86 (78.2)</b>
Reduced hours	6 (7.0)
Increased Hours	6 (7.0)
No Change	74 (86.0)

**Table 3. Correlations between demographic/injury factors, symptoms and mood at one month post-injury and work productivity at four years, N = 86**

	WLQ Time Management	WLQ Physical Demands	WLQ Mental /Interpersonal	WLQ Output Demands	WLQ Productivity Loss
Age	-.04	.19	-.02	.09	.10
Gender	-.03	-.16	-.02	-.17	.13
Ethnicity	.04	-.09	-.12	-.07	-.09
Mechanism	.21	.11	.11	.08	.15
Education Level	-.16	-.22	-.12	-.02	-.06
Marital status	.07	.03	.02	.10	.08
Residence area	.03	.20	-.05	-.02	.01
Additional Injury	-.15	-.09	-.04	.04	-.07
Prior TBI	.26	.19	<b>.36</b>	.11	<b>.29</b>
Recurrent TBI	.17	.04	<b>.37</b>	.27	<b>-.37</b>
Symptoms (RPQ)					
Headaches	.15	.06	.27	.24	.24
Dizziness	.21	<b>.36</b>	<b>.38</b>	.25	<b>.37</b>
Nausea	-.22	.02	.11	.04	-.04
Noise sensitivity	.17	.11	.29	.35	<b>.37</b>
Sleep disturbance	.12	.09	-.02	-.19	.04
Fatigue	.25	.25	<b>.41</b>	<b>.36</b>	<b>.39</b>
Irritable	.19	.22	<b>.52</b>	<b>.36</b>	<b>.43</b>
Depressed	.11	.15	.27	.19	.22
Frustration	.14	.15	<b>.41</b>	.27	.30
Forgetful	.26	.31	<b>.41</b>	.30	<b>.49</b>
Poor concentration	.33	<b>.36</b>	<b>.53</b>	<b>.39</b>	<b>.55</b>
Longer to think	<b>.39</b>	<b>.38</b>	<b>.62</b>	<b>.49</b>	<b>.52</b>
Blurred vision	-.02	.09	-.10	-.06	.05
Light sensitivity	.06	.24	.20	.24	.29
Double vision	.02	.07	.13	.00	.10
Restlessness	.28	.20	.31	.20	.31
Mood (HADS)					
Anxiety	.34	.32	<b>.51</b>	.30	<b>0.44</b>
Depression	.18	.04	<b>.35</b>	<b>.40</b>	0.30

Significant correlations  $p < 0.01$  in bold

Table 4. Predictors of work productivity four years post mild TBI (N = 86)

Outcome variable	Parameter	$\beta$	t	P Value	Lower CI	Upper CI	R sq	F	P value
<b>Time Management</b>	Poor concentration	0.32	2.32	0.025	0.53	7.64			
						<b>Overall Model</b>	0.10	5.36	0.025
<b>Physical Demands<sup>≠</sup></b>	Taking longer to think	0.39	2.86	0.006	1.28	7.34			
						<b>Overall Model</b>	0.15	8.18	0.006
<b>Mental/ Interpersonal<sup>∞</sup></b>	Poor concentration	0.52	4.12	<0.001	3.44	10.01			
						<b>Overall Model</b>	0.27	16.94	<0.001
<b>Output Demands<sup>§</sup></b>	Taking longer to think	0.32	2.30	0.026	-0.62	9.21			
						<b>Overall Model</b>	0.10	5.30	0.026
<b>Productivity Loss<sup>†</sup></b>	Taking longer to think	0.47	3.79	<0.001	<0.001	0.02			
	Recurrent TBI	0.32	2.59	0.013	-0.06	-0.01			
						<b>Overall Model</b>	0.32	10.24	<0.001

<sup>≠</sup> Excluded variables = dizziness and poor concentration

<sup>∞</sup> Excluded variables = prior TBI, recurrent TBI, dizziness, fatigue, irritability, frustration, forgetfulness, taking longer to think, anxiety and depression

<sup>§</sup> Excluded variables = fatigue, irritability, poor concentration and depression

<sup>†</sup> Excluded variables = prior TBI, dizziness, noise sensitivity, fatigue, irritability, forgetfulness, poor concentration and anxiety.

