

1 The psychometric validation of the Dutch version 2 of the Rivermead Post-Concussion Symptoms 3 Questionnaire (RPQ) after Traumatic Brain Injury 4 (TBI)

5 *Short Title:* Psychometric Validation of the Dutch RPQ

6 Anne Marie C Plass^{1*}, Dominique Van Praag², Amra Covic¹, Anastasia Gorbunova¹, Ruben
7 Real¹, Nicole von Steinbuechel¹, and Dutch and Flemish CENTER TBI investigators.

8 ¹Institute of Medical Psychology and Medical Sociology, University Medical Center Göttingen (UMG)/
9 Georg-August-University, Göttingen, Germany

10 ²Department of Neurosurgery, Antwerp University Hospital and University of Antwerp, Edegem,
11 Belgium

12

13 *corresponding Author:

14 dr Anne Marie Plass, Ph.D,

15 Institute of Medical Psychology and Medical Sociology,

16 University Medical Centre Göttingen (UMG)/ Georg-August-University

17 Waldweg 37, Eingang A| 37073 Göttingen| Germany

18 (P)+49 551 39-8014/ -8192 (secretary)

19 (F) +49 551 39-8194|

20 (E) annemarie.plass@med.uni-goettingen.de

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22

23 **Abstract**

24 *Background*

25 Traumatic brain injury (TBI) is one of the most common neurological conditions. It can have wide-
26 ranging physical, cognitive and psychosocial effects. Most people recover within weeks to months after
27 the injury, but a substantial proportion are at risk of developing lasting post-concussion symptoms.
28 The Rivermead Post-Concussion Syndrome Questionnaire (RPQ) is a short validated 16-items self-
29 report instrument to evaluate post-concussive symptoms. The aim of this study was to test
30 psychometrics characteristics of the current Dutch translation of the RPQ.

31 *Methods*

32 To determine the psychometric characteristics of the Dutch RPQ, 472 consecutive patients six months
33 after they presented with a traumatic brain injury in seven medical centers in the Netherlands ($N=397$),
34 and in two in Belgium (Flanders) ($N=75$) took part in the study which is part of the large prospective
35 longitudinal observational CENTER-TBI-EU-study. Psychometric properties at six months post TBI, were
36 assessed using exploratory and confirmatory factor analyses. Sensitivity was analyzed by comparing
37 RPQ scores of patients after mild vs. moderate and severe TBI.

38 *Findings*

39 The Dutch version of RPQ proved good, showing excellent psychometric characteristics: high internal
40 consistency (Cronbach's α .93), and good construct validity, being sensitive to self-reported recovery
41 status at six months post TBI. Moreover, data showed a good fit to the three dimensions structure of
42 separate cognitive, emotional and somatic factors ($Chi^2=119$; $df=117$; $p=.4$; CFI=.99; RMSEA=.006),
43 reported earlier in the literature.

44 *Discussion*

45 Psychometric characteristics of the Dutch version of RPQ proved excellent to good, and can the
46 instrument therefore be applied for research purposes and in daily clinical practice.

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49 Validation; Psychometric Properties, Post-Concussion (PCS)

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52 **Introduction**

53 Traumatic brain injury (TBI) is one of the most common neurological conditions, and occurs
54 when an external force causes brain trauma (1). TBI can be classified as mild, moderate or
55 severe, and may have wide-ranging physical and psychological effects (2–7). Some signs or
56 symptoms appear immediately after the traumatic event, while others days or weeks later. In
57 the Netherlands, about 85,000 people are confronted with a traumatic brain injury, on a
58 yearly basis ([https://www.hersenstichting.nl/alles-over-hersenen/hersenaandoeningen/cijfers-](https://www.hersenstichting.nl/alles-over-hersenen/hersenaandoeningen/cijfers-over-patienten)
59 [over-patienten](https://www.hersenstichting.nl/alles-over-hersenen/hersenaandoeningen/cijfers-over-patienten)). On average 30,000 of these seek help at the Emergency Room (ER) of the
60 hospital, and about 21,000 require hospital stays (8). Yearly, about 1,000 die because of TBI
61 (9). Most people suffer from mild TBI (mTBI), e.g. concussion, for which they often do not
62 seek professional help, or seek advice from their General Practitioner (GP): In the Netherlands,
63 the GP functions as gatekeeper to the rest of the medical system. (8). Virtually all non-
64 institutionalized Dutch citizens are registered with a GP controlling access to specialized
65 medical care. (10)

66 In about one third of the cases mTBI leads to long-term consequences (5,11). A
67 substantial proportion (about 15-30%) of individuals after mTBI are at risk to developing
68 post-concussion symptoms (2). These can be classified into four categories: cognitive
69 difficulties (e.g. concentration and memory loss), behavioral maladaptation (e.g. impulsivity,

70 and aggressive behavior), psychiatric conditions (e.g. posttraumatic stress), and physical
71 disorders (e.g. chronic pain) (11). Whether patients develop post-concussion symptoms, is
72 associated with cognitive, emotional, behavioral and social risk factors, and does not
73 necessarily depend on the severity of the traumatic brain injury, see fig 1 (2). Years of
74 education, pre-injury psychiatric disorders, neck pain and prior TBI were found strong
75 predictors of 6-month post-concussive symptoms (3,5), as were patient's perceptions of their
76 brain injury, their behavioral responses, passive and avoidant coping styles and emotional
77 distress in response to this (2,5).

78 As most people recover from their TBI within weeks to months after the injury, post-
79 concussion symptoms might easily be overlooked, since these residual complaints may be
80 deferred. Furthermore, imaging techniques often do not show any structural brain damage in
81 this population (12,13). Still, one in three mTBI patients will not be able to resume work and
82 activities six months after the event at a level similar to that before the accident(5). As a
83 consequence, (m)TBI is associated with substantial ongoing disability and distress for
84 patients, and high healthcare costs (2,8). A possible instrument for (early) identification in
85 order to timely guide clinical management of post-concussion symptoms after TBI, is the
86 Rivermead Post Concussion Syndrome Questionnaire (RPQ). The RPQ is a validated
87 measurement-instrument to survey post-concussion symptoms, relying on self-report as to
88 the presence and severity of 16 symptoms (14–17). The items form one scale, but were not
89 always found to tap into the same underlying construct (3,4,14)). Eyres et al (2005) found no
90 evidence for a single factor structure, and proposed to split the RPQ into two subscales
91 consisting of the first three items 'RPQ3', representing immediate symptoms (headaches,
92 dizziness, and nausea) and the remaining 13 items 'RPQ13', representing symptoms that
93 might occur at a later stage. On the other hand, Lannsjö et al (2011) found strong support for

94 both a single and two factor structure in their RPQ validation study, but failed to reproduce
95 the RPQ3/13 two-category model as suggested by Eyres et al (2005). Furthermore, a
96 'rationally-based' three categories model was proposed by Smith-Seemiller and colleagues
97 (2003), comprising of the following domains: 1. cognitive deficits (impaired memory, poor
98 concentration, slow thinking), 2. somatic complaints (headaches, dizziness, nausea, blurred or
99 double vision, noise or light sensitivity, sleep disturbance, fatigue), and 3. emotional
100 complaints (irritably, depression, frustration, restlessness), serving as framework in various
101 studies on post-concussion symptoms (3,18). The results of Potter and colleagues (2006)
102 supported the existence of separate cognitive, emotional and somatic factors (17). So far, the
103 RPQ has been validated in various languages (4,6), but until now this has not been the case
104 for the Dutch version of this questionnaire. Therefore, this study aims to investigate the
105 psychometric characteristics of the current Dutch translation of the RPQ.

106 **Methods**

107 *Study sample*

108 This study is part of the Collaborative European NeuroTrauma Effectiveness Research in TBI
109 (CENTER-TBI) study, which is a prospective longitudinal observational study conducted in 72
110 centers from 21 countries (8). In the Netherlands, patients were recruited from seven medical
111 centers spread over the country: Leiden University Medical Center (LUMC), University Medical
112 Center Groningen (UMCG), Erasmus MC Rotterdam, Radboud University medical center
113 Nijmegen, Medical center Haaglanden The Hague, Elisabeth Hospital Tilburg, HAGA hospitals
114 The Hague. Furthermore, two centers in the Dutch-speaking-part of Belgium (Flanders) were
115 included in the study: the Antwerp University Hospital, and the University Hospital in Leuven.
116 Patients that presented within 24 hours after brain injury at the hospital, that were diagnosed
117 with TBI, and had a clinical indication for CT scan, were eligible for the study, and were all

118 invited to participate in this convenience sample. Those willing to participate provided written
119 informed consent prior to inclusion. Patients with severe pre-existing neurological disorder
120 that could confound the outcome assessment were excluded. A written informed consent to
121 participate in the study was obtained at the time of inclusion. At six months post TBI, the
122 nurse at the center administered the RPQ during a visit, or was sent by postal mail to those
123 who did not need to attend the hospital, for completion at home. A pre-franked envelope
124 was included to send it back.

125 *Translation of the Dutch RPQ*

126 Two native Dutch speakers who are proficient in English translated the RPQ into Dutch, after
127 which a native English speaker, who is fluent in Dutch, backward translated the harmonized
128 version. This version was then compared to the original English RPQ version possible
129 differences were identified and resolved by the two parties. In addition, a team of researchers
130 and CENTER-TBI collaborators refined and reshaped the measurement-instrument until
131 consensus was reached, following an iterative process. This multiple-step procedure resulted
132 in a final version of the Dutch RPQ.

133 *Ethical Approval*

134 The CENTER-TBI study (EC grant 602150) has been conducted in accordance with all relevant
135 laws of the EU if directly applicable or of direct effect, and all relevant laws of the country
136 where the Recruiting sites were located, including, but not limited to, the relevant privacy and
137 data protection laws and regulations (the "Privacy Law"), the relevant laws and regulations on
138 the use of human materials, and all relevant guidance relating to clinical studies from time to
139 time in force including, but not limited to, the ICH Harmonised Tripartite Guideline for Good
140 Clinical Practice (CPMP/ICH/135/95) ("ICH GCP") and the World Medical Association

141 Declaration of Helsinki entitled “Ethical Principles for Medical Research Involving Human
142 Subjects”. Ethical approval was obtained for each recruiting site. Informed Consent was
143 obtained for all patients recruited in the Core Dataset of CENTER-TBI and documented in the
144 e-CRF. The list of sites, Ethical Committees, approval numbers and approval dates can be
145 found on the official Center TBI website (www.center-tbi.eu/project/ethical-approval).

146 *Measurement-Instrument*

147 The Rivermead Post-Concussion Questionnaire (RPQ) consists of 16 common symptoms
148 related to post concussion. Patients are asked to rate how problematic symptoms were
149 compared to the situation before the head injury on a 5-point Likert scale (0-4). A score of 0
150 indicating ‘not experienced at all; 1 indicating ‘no more of a problem (than before)’, 2
151 indicating ‘a mild problem’; 3 indicating ‘a moderate problem; 4 indicating ‘a severe problem’
152 (14). Scores are taken as sum of all symptom scores, excluding scores of 1, as these indicate
153 symptoms are unchanged since the brain injury. This gives a potential total score range of 0
154 (representing no change in symptoms since the head injury) to 64 (most severe symptoms)
155 (4). If more than 5 of the items were missing from the 16, a score was not calculated and
156 treated as missing. The RPQ total score is calculated using prorating as imputation method, if
157 up to one third of the items were missing. In addition, the RPQ scoring method of Stulemeijer
158 et. al. (2008) was applied where a score of highest 2 (a mild problem) to at least 13 of the 16
159 items is defined a favorable outcome. Stulemeijer et al (2008) showed that 94% of non-brain-
160 injured patients (wrist-, or ankle distortion) would meet this criterion (19).

161 Further, TBI severity was rated using the Glasgow Coma Scale (GCS), with scores
162 between 3-8 indicating severe, 9-12 moderate, and 13-15 mild TBI (20–24). The GCS was
163 administered within the first 24 hours after the brain injury occurred. Current disability was
164 assessed by administering the extended Glasgow Outcome Scale (GOSE) (25). GOSE scores

165 were used to differentiate between patients with remaining severe disability (3-4), moderate
166 disability (5-6), and good recovery (21,22). In addition, socio- demographic data were
167 collected, including gender, age, working status, education level, etc.

168 *Analyses*

169 The psychometric characteristics of the Dutch version of the RPQ were determined at six
170 months post TBI, using SPSS version 24, AMOS version 24, and R version 3.3.3 to performing
171 classical and modern test-theory analyses.

172 Internal consistency was determined by calculating Cronbach's alpha, including the
173 scale if any item were deleted. To testing construct validity, Principal Axis Factoring (PAF) was
174 done by unweighted least squares and oblimin rotation on the 16 RPQ items, exploring the
175 underlying constructs in the Dutch version of the RPQ, as no consistent underlying factor
176 structure has been established so far. Items were included if the factor loading was 0.5 or
177 higher and if factor loadings on the other factors were 0.1 or lower. Confirmatory factor
178 analysis (CFA) was used to examine the fit of the Dutch RPQ data to the various factor
179 structures that had been described earlier in literature: For this, we used the single model
180 factor, reflecting post-concussion symptoms as unitary entity (15); the RPQ3 and RPQ13 two
181 factor model (4); and the three factor model (17,18).

182 Concurrent criterion validity was assessed by analyzing the influence of important
183 covariates on RPQ scores (GCS, GOSE) using t-tests and one-way ANOVA. Descriptive
184 analyses were performed for sociodemographic variables (gender, age, education level, etc.).
185 Although people in The Netherlands and Flanders (Belgium) both speak Dutch, the language
186 used differs, and words might have a different meaning. Therefore, all tests were performed
187 both for the entire research sample and for each country separately where possible.

188 **Results**

189 *Sample*

190 In total 472 patients filled in the Dutch version of the RPQ at six months post TBI. Of these,
191 397 were administered in the Netherlands and 75 in Belgium. Twenty-five participants were
192 aged under 18 (18 in the Netherlands, and seven in Belgium) and were excluded from this
193 study. Country of residence was registered for 437 patients, who were either living in Belgium
194 (N=67), or the Netherlands (N= 368), apart from two in Nepal. Not all respondents were born
195 in the Netherlands or Belgium (see table 1), but since their understanding of the Dutch
196 language was sufficient to fill in the RPQ, and since they currently were living in the
197 Netherlands or Flanders, they were not excluded from the study. There were 277/ 447 (62%)
198 male respondents (resp. 231/ 379 (60.9%) in the Netherlands, and 46/ 68 (67.6%) in Belgium).
199 The vast majority of the study population (68.2%) belonged to the middle aged and older
200 age groups (38% was aged 45-65, 30.2% 65 and up). More than half were higher educated
201 (64.2%), and were either married or living together (56.6%). Nearly half were not, or no longer
202 employed (48.4%), see table 2 for further details. There were no significant differences in total
203 RPQ scores at six months post TBI for study participants in the Netherlands ($M=12.63$;
204 $SD=13.77$) and participants in Belgium ($M=12.64$; $SD=11.94$; $t(445)=-110$; $p=.9$). The
205 magnitude of the differences in the means was very small (eta squared < .0001).

206 At the time of the injury, 13.4% (N=60) of the study population solely attended the ER
207 without further hospitalization. 52.6% (N=235) were hospitalized, and 34 % (N=152) needed
208 a stay in the ICU. Initially, within 24 hours after TBI, 80.5% (N=316) were diagnosed mTBI,
209 5.8% (N=21) were diagnosed moderate TBI, and 6.4% (N=23) with severe TBI. Of 87
210 respondents (19.5%) these data were missing. Six months after the brain injury, GOSE scores
211 reveal that 55.5% (N=248) of the respondents showed good recovery, 25% (N=112) reported

212 moderate disabilities, and 6.7% (N=30) suffered from severe disabilities at that point in time.
213 Of 57 respondents (12.8%) these data were missing. Those patients that solely attended the
214 ER without further hospitalization, all were initially diagnosed mTBI (see Table 3a). Of 313
215 respondents all three data types were available (hospitalization, initial diagnosis and six
216 months post recovery status), see table 3b for further descriptives.

217 *Factor analyses*

218 Prior to performing PAF the suitability of data for factor analyses was assessed. Inspection of
219 the correlation matrix revealed the presence of many coefficients of .3 and above. The Kaiser-
220 Meyer-Okin (KMO) value was .94, exceeding the recommended value of .6 (Kaiser 1970,
221 1974) and the Barlett's Test of sphericity (Barlett, 1954) reached statistical significance (p
222 $<.0001$), supporting the factorability of the correlation matrix. PAF, using Oblimin rotation,
223 revealed the presence of three components with eigenvalue exceeding 1, explaining 47.7%,
224 5.0%, and 4.0% of the variance respectively. The scree plot revealed a clear break after the
225 first component, see figure 2. Moreover, all items except for three (nausea (.42), blurred vision
226 (.42) and double vision (.34)) show factor loadings of .5 and up on the first factor, but high
227 factor loadings ($>.1$) on at least one of the other factors too. Confirmatory Factor analysis
228 (CFA) was run to testing model fit to possible underlying factor structures that had been
229 described in literature earlier (4,6,17,18). A central assumption is that the data are distributed
230 normally. However, substantial problems with univariate skew and kurtosis were identified,
231 see table 4. To correct for this data were dichotomized, computing 0 and 1 responses into 0,
232 and 2, 3, and 4 into 1. Following this, CFA indicated a lack of fit to unitary model, given the
233 significant Chi-squares (15) ($Chi^2=285.5$; $df=120$; $p<.001$; CFI=.99; RMSEA=.06), and a lack of
234 fit to the RPQ3/ RPQ13 two component model(4) ($Chi^2=271$; $df=119$; $p<.001$; CFI=.99;

235 RMSEA=.05), but showed a good fit to the three factor structure(18) ($Chi^2=119$; $df=117$; $p=.4$;
236 CFI=.99; RMSEA=.006). The Belgian sample was too small to performing separate CFAs.

237 *Quality Criteria*

238 The RPQ showed high consistency with Cronbach's alpha being .93. For the sample in the
239 Netherlands Cronbach's alpha was .94, and for the sample in Belgium Cronbach's was alpha
240 .91. The scale did not improve if any items were deleted. Spearman Brown Coefficient r_{sb1} was
241 .91). Further, item characteristics showed high item correlations (>.55, except for double
242 vision and nausea), and acceptable asymmetry for all items but double vision (skewness
243 being 2.6), see table 5

244 *Concurrent Criterion validity*

245 RPQ total scores at six months post TBI of patients (self-)reporting severe ($M=20.7$; $SD=18.3$;
246 $N=30$) or moderate ($M=20.2$; $SD=13.9$; $N=112$) disabilities according to their total scores on
247 the GOSE scale, differed significantly to those that showed good recovery at this point in time
248 ($M=8.3$; $SD=10.6$; $N=248$) ($F(2, 387)=42.7$; $p<.001$). RPQ total scores at six months post TBI
249 further were found to differentiate between initial mTBI ($M=11.6$; $SD=13.2$; $N=316$) and
250 moderate TBI ($M=20.2$; $SD=16.8$; $N=21$) diagnoses (GCS-scores) ($F(2, 357)=4.5$; $p=.012$) .
251 Remarkably, RPQ total scores of those initially diagnosed with severe TBI ($M=14.8$, $SD =13.9$;
252 $N=23$), resembled most those initially diagnosed with mTBI (*NS*). When recalculating RPQ
253 total score into favorable (a score of highest 2 to at least 13 of the 16 items (26)) vs
254 unfavorable, 74.7% ($N=334$) of the study population had a favorable outcome at six months
255 post TBI, indicating that the symptoms reported, do not differ from what can be found in a
256 non-TBI population (26,27). 25.3% ($N=113$) still had an unfavorable, strongly related to TBI,

257 outcome. The RPQ score was found to discriminate between recovery status (GOSE scores) at
258 six months post TBI ($Chi^2=45.2$; $df=2$; $p<.001$), although not strongly ($Cramer's V =.11$).
259 When solely taking the sample from the Netherlands into account, a stronger relationship
260 between favorable and unfavorable RPQ outcomes and recovery status (GOSE scores) at six
261 month post TBI was found ($Cramer's V =.35$, $Chi^2=43.8$; $df=2$; $p<.001$). Further, the RPQ total
262 scores at six months post TBI of the Dutch sample were found to discriminate between
263 recovery status (GOSE scores) at six months post TBI ($F(2, 358)=39.2$; $p<.001$), and initial
264 diagnoses within 24 hours after the brain injury occurred (GCS Scores) ($F(2, 299)=3.7$; $p=.3$).
265 The number of participants from Belgium that could be included in these analyses were too
266 low for further analyses.

267 **Discussion**

268 The current Dutch translation of the RPQ showed good psychometric characteristics, with
269 high internal consistency, and good construct validity. As for these aspects, it can be applied
270 for research purposes and in daily clinical practice, as an instrument to identify post-
271 concussion symptoms. Besides, it proved sensitive for recovery status at six months post TBI,
272 showing that those who (self-) reported moderate or severe disabilities six months after the
273 brain injury took place, had significant lower RPQ total scores compared to those reporting
274 good recovery at that time point. Further, RPQ total scores at six months post TBI were found
275 to distinguish between initial TBI diagnoses: Those initially diagnosed with moderate TBI had
276 higher RPQ total (sum) scores at six months post TBI compared to those initially diagnosed
277 with mild TBI (mTBI). However, the number of people in our research sample that were
278 initially diagnosed with moderate TBI was low. Moreover, RPQ total scores at six months post
279 TBI of those initially diagnosed with severe TBI resembled more the total score of those
280 diagnosed with mTBI, rather than those with moderate TBI. A possible explanation for this

281 might be that moderate TBI and the amount of care needed was underestimated. This type of
282 TBI might need more intensive care than what was provided. However, again the number of
283 people in this group was too low to base further conclusions upon. Another explanation for
284 this might be that the large number of mTBI-diagnosed patients who were admitted to the
285 ICU, needed intensive care because of other injuries, and thus were diagnosed mTBI correctly.

286 Consistent with the findings of others, we found multidimensionality as underlying
287 structure of the RPQ measurement-instrument (3,4,6,17,18,26). However, high factor loadings
288 of items on multiple factors, and the clear break after the first factor in the scree plot, would
289 suggest a one factor structure rather than multidimensionality. Confirmative factor analyses
290 on the other hand revealed that our data would fit best to the three-component model
291 dividing the RPQ post-concussion symptoms into the following three categories: 1. cognitive
292 deficits (impaired memory, poor concentration, slow thinking), 2. somatic complaints
293 (headaches, dizziness, nausea, blurred or double vision, noise or light sensitivity, sleep
294 disturbances), and 3. emotional complaints (irritably, depression, frustration, restlessness).
295 Small differences were found between the Belgium and Dutch sample, with only the Dutch
296 sample showing a good fit to this model. However the number of respondents in the Belgium
297 sample was below 250, due to which the criteria for model fit may not be valid (28).

298 The variation in underlying structure of the RPQ differs between studies and countries
299 and might be due to various reasons. One reason could be the convenience sample used for
300 this research. Another explanation might be the different analyses techniques used in the
301 various studies, as modern techniques often tend to disqualify measurement-instrument
302 validity that had been established before by classical analyses methods (29). A third possible
303 explanation underpinning this may be the way in which measurement-instruments are being
304 translated in accordance to the WHO guidelines of forward and back translation (30). In order

305 not to lose the potential to comparing data, researchers prefer to stay as close as possible to
306 the original version. However, through this, the principles of cultural interpretation and
307 translating the correct meaning of the items, might be missed out, due to which country
308 differences might occur, even though the instrument used is very similar. (31,32).

309 In addition, we argue that despite the underlying multidimensionality found, the
310 Dutch version of the RPQ needs not necessarily be divided into subscales when applied for
311 research purposes and in daily clinical practice. The underlying multidimensionality might
312 indicate that post-concussion symptoms represent more than one dimension, but factors
313 highly correlated, and items were not unique for just one of the factors. Moreover, there is a
314 large body of evidence that supports the use of total scores of scales to which
315 multidimensionality is a precondition, e.g. attitude scales that usually exist of a cognitive and
316 affective component(33). Furthermore, as the psychometric properties of the Dutch version of
317 the RPQ proved good, it would be of interest to implement this measurement-instrument
318 into primary care settings, in order to timely recognize the possible long-term consequences
319 of TBI. This would especially be effective in countries as the Netherlands where one has to
320 see the GP first, before entering the rest of the medical system, the so-called gatekeeper
321 system, and in ER settings, where people usually are only checked medically and then send
322 home, in order to timely identify potential patients at risk. However, more clarity is needed on
323 how to best interpret RPQ scores(17), since similar symptoms can too be reported by those
324 suffering from different injuries and, disorders, or by members of the general population as
325 fatigue, headaches, nausea etc., are very common. As such, Stulemeijer and colleague (2016)
326 found that 94% of non-TBI patients with wrist or ankle distortion too score positive on a
327 maximum of three RPQ items.

328 *Limitations:* At six months post TBI, three quarters of the research sample no longer showed
329 post-concussion symptoms, due to which data were skewed and not normally distributed.
330 Validation of the RPQ at this point in time might therefore be difficult. Further, items were
331 strongly correlated, due to which items strongly loaded on one and the same factor. Another
332 limitation of this study was the limited Belgian sample, which was often too small for sound
333 complex analyses, such as CFA. Further, the lacking of a construct validation phase making
334 use of cognitive interviewing limits the overall conclusion on validity of the Dutch version of
335 the RPQ, especially since there were between-country differences. Knowing what our
336 respondents think we are asking, and knowing how they interpret the questions we are
337 asking might help to explain the variety in underlying symptom structure found too (29).
338 Moreover, the response scale used (0-4) might be confusing since patients might find the
339 following order of the score of zero indicating 'no problem', and the score of one indicating
340 'no more of a problem' difficult to understand, as a more natural following order would be:
341 zero indicating no problem, and one indicating a small problem. Last, the convenience
342 hospital sample used in this study might be limited representative to the entire mTBI
343 population, as most people in the Netherlands tend not to seek specialized medical help for
344 their head injury.

345 **Conclusion**

346 The psychometric characteristics of the Dutch version of RPQ proved good, showing high
347 consistency, and good construct validity, being sensitive to self-reported recovery status at
348 six months post TBI and initial TBI-diagnosis sensitive. The Dutch version of the RPQ can
349 therefore be applied for research purposes and in daily clinical practice. Further discussion is
350 needed with regard to the scoring of the RPQ, as underlying multidimensionality may not
351 necessarily stand in the way of using a total score.

352

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360 Principal Investigators and contact information: Professor A.I. Maas: Andrew.Maas@azu.be; Professor D. Menon:
361 dkm31@wbic.cam.ac.uk.

362 Ardon Hilko³, Bartels Ronald⁴, Carpenter K⁵, Covic Amra¹, Cnossen Maryse⁶, De Keyser Véronique², De Ruiter
363 Godard C.W²³, Depreitere Bart⁷, Dippel Diederik⁸, Engemann Heiko¹, Foks Kelly⁸, Geleijns Karin⁵, Haagsma Juanita
364 A.⁶, Haitisma Iain¹⁰, Hoedemaekers Astrid¹¹, Jacobs Bram¹², Janssens Koen², Kalala Jean-Pierre¹³, Ketharanathan
365 Naomi⁵, Kompanje Erwin¹⁴, Lecky Fiona¹⁵, Lingsma Hester⁶, Loeckx Dirk¹⁶, Luijten-Arts Chantal¹¹, Maas Andrew
366 I.R.², Menon David⁹, Menovsky Tomas², Schoonman Guus²⁰, Oldenbeuving Annemarie¹⁷, Parizel Paul M.¹⁸, Peul
367 Wilco¹⁹, Polinder Suzanne⁶, Pullens Pim¹⁸, Roks Gerwin²⁰, Ruiz de Arcaute Felix¹⁶, Schipper Inger²¹, Sir Özcan²²,
368 Smakman Lidwien²³, Smeets Dirk¹⁶, Steyerberg Ewout W.⁶, Tibboel Dick⁵, Vande Vyvere Thijs¹⁶, Van Der Jagt
369 Mathieu²⁵, Van Der Naalt Joukje¹², Van Dijck Jeroen²⁴, Van Hecke Wim¹⁶, Van Vlierberghe Eline¹⁶, Verheyden Jan¹⁶,
370 Wildschut Eno⁵, Van Essen Thomas A.¹⁹, Van Praag Dominique², Van Roost Dirk¹³, Vleggeert-Lankamp Carmen²³,
371 Volovici Victor¹⁰, Von Steinbüchel Nicole¹.

372 ¹ Institute of Medical Psychology and Medical Sociology, Universitätsmedizin Göttingen, Göttingen, Germany

373 ² Department of Neurosurgery, Antwerp University Hospital and University of Antwerp, Edegem, Belgium

374 ³ Department of Neurosurgery, Elisabeth-Tweesteden Ziekenhuis, Tilburg, the Netherlands

375 ⁴ Department of Neurosurgery, Radboud University Medical Center, The Netherlands

376 ⁵ Intensive Care and Department of Pediatric Surgery, Erasmus Medical Center, Sophia Children's Hospital,
377 Rotterdam, The Netherlands

378 ⁶ Department of Public Health, Erasmus Medical Center-University Medical Center, Rotterdam, The Netherlands

379 ⁷ Department of Neurosurgery, University Hospitals Leuven, Leuven, Belgium

380 ⁸ Department of Neurology, Erasmus MC, Rotterdam, the Netherlands

381 ⁹ Division of Anaesthesia, University of Cambridge, Addenbrooke's Hospital, Cambridge, UK

382 ¹⁰ Department of Neurosurgery, Erasmus MC, Rotterdam, the Netherlands

383 ¹¹ Department of Intensive Care Medicine, Radboud University Medical Center, The Netherlands

384 ¹² Department of Neurology, University Medical Center Groningen, Groningen, Netherlands

385 ¹³ Department of Neurosurgery, UZ Gent, Gent, Belgium

386 ¹⁴ Department of Intensive Care and Department of Ethics and Philosophy of Medicine, Erasmus Medical Center,
387 Rotterdam, The Netherlands

388 ¹⁵ Emergency Medicine Research in Sheffield, Health Services Research Section, School of Health and Related
389 Research (SchARR), University of Sheffield, Sheffield, UK

390 ¹⁶ icoMetrix NV, Leuven, Belgium

391 ¹⁷ Department of Intensive Care, Elisabeth-Tweesteden Ziekenhuis, Tilburg, the Netherlands

392 ¹⁸ Department of Radiology, Antwerp University Hospital and University of Antwerp, Edegem, Belgium

393 ¹⁹ Dept. of Neurosurgery, Leiden University Medical Center, Leiden, The Netherlands and Dept. of Neurosurgery,
394 Medical Center Haaglanden, The Hague, The Netherlands

395 ²⁰ Department of Neurology, Elisabeth-TweeSteden Ziekenhuis, Tilburg, the Netherlands

396 ²¹ Department of Traumasurgery, Leiden University Medical Center, Leiden, The Netherlands

397 ²² Department of Emergency Care Medicine, Radboud University Medical Center, The Netherlands

398 ²³ Neurosurgical Cooperative Holland, Department of Neurosurgery, Leiden University Medical Center and Medical
399 Center Haaglanden, Leiden and The Hague, The Netherlands

400 ²⁴ Department of Neurosurgery, The HAGA Hospital, The Hague, The Netherlands

401 ²⁵ Department of Intensive Care, Erasmus MC, Rotterdam, the Netherlands

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407 decision to publish, or preparation of the manuscript.

408 Data Availability

409 There are however legal constraints that prohibit us from making the data available. Since
410 there are only a limited number of centers per country included in this study (for two

411 countries only one center), data will be identifiable. Readers may contact Dr Hester Lingsma
412 (h.lingsma@erasmusmc.nl) for requests for the data.

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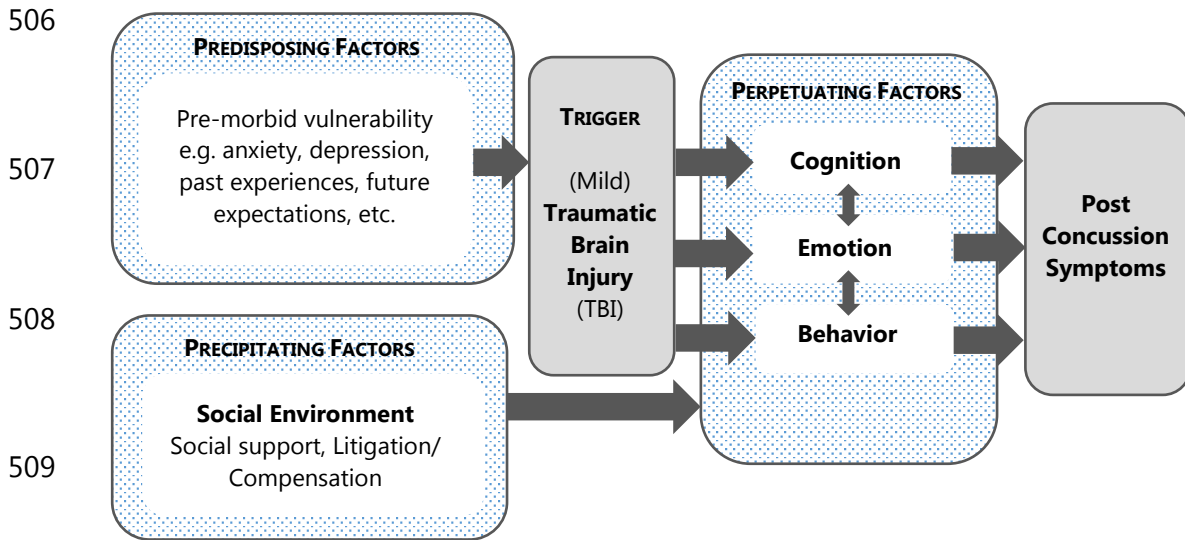
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- 504

505 **Figure 1** Factors influencing the development of Post-Concussion Symptoms after TBI (2)

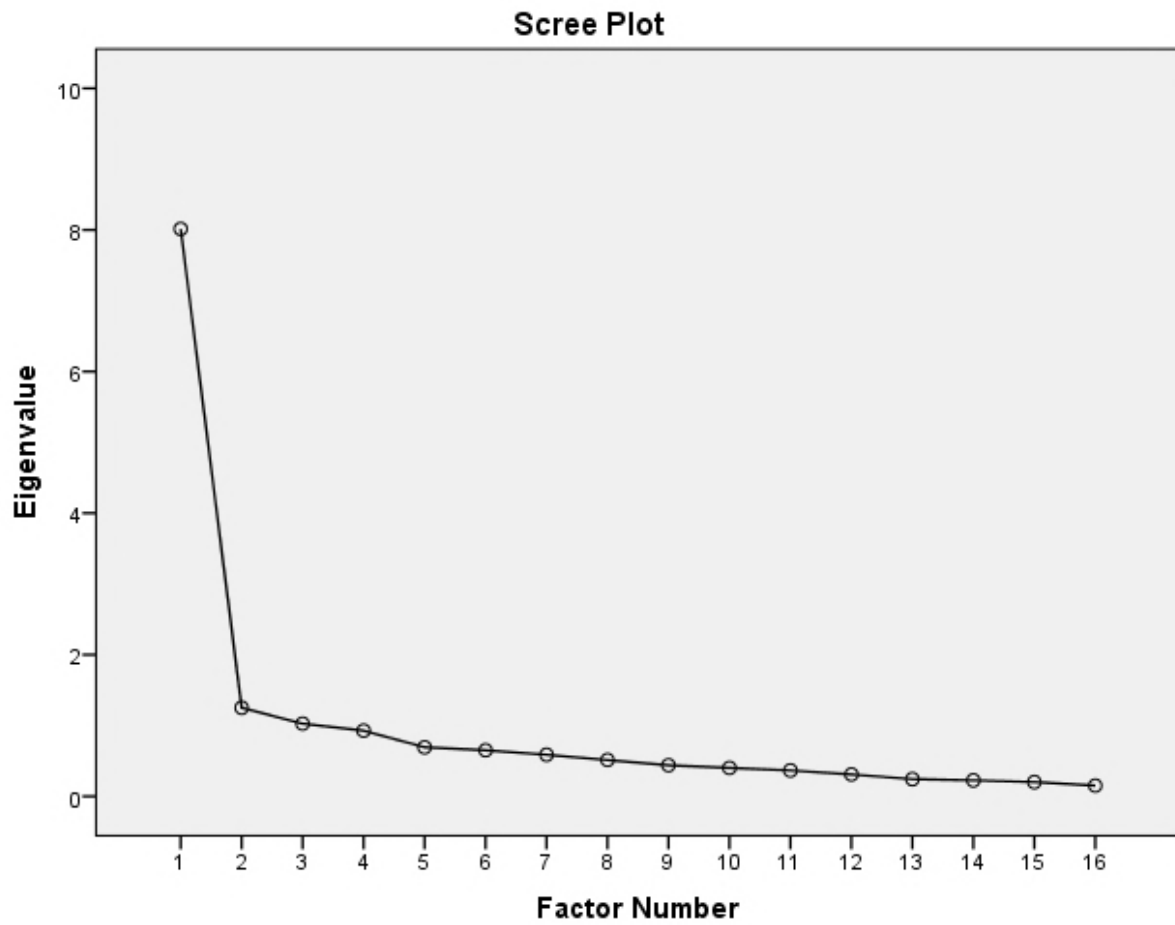


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512 **Figure 2** Visual representation of factor loadings

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Table 1 Overview of countries of birth of the respondents (N=447)

Aruba (N=1)	Brasil (N=1)	Germany (N=2)	Indonesia (N=4)	Morocco (N=3)	Surinam (N=6)
Bosnia and Herzegovina (N=1)	China (N=3)	Spain (N=1)	Ireland (N=1)	Netherlands (N=326)	Saint-Martin (N=1)
Belgium (N=59)	Colombia (N=1)	UK (N=3)	Iran (N=1)	Slovenia (N=1)	Turkey (N=3)
Bermuda (N=1)	Cape Verde (N=1)	Greece (N=1)	Italy (N=1)	Somalia (N=1)	Vietnam (N=1)

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523 **Table 2 Patient Demographics, presenting percentages and numbers**

Agegroup	% NL (N)	% BE (N)	% Total (N)
18 to 24	12.9 (49)	5.9 (4)	11.9 (53)
25 to 34	10.3 (39)	11.8 (8)	10.5 (47)
35 to 44	8.2 (31)	16.2 (11)	9.4 (42)
45 to 54	13.7 (52)	19.1 (13)	14.5 (65)
55 to 64	23.2 (88)	25.0 (17)	23.5 (105)
65 to 74	18.7 (71)	13.2 (9)	17.9 (80)
75 and up	12.9 (49)	8.8 (6)	12.3 (55)
Total (N)	379	68	447

524

Employment Status	% NL (N)	% BE (N)	% Total (N)
Working >34hpw	28.7 (118)	7.3 (30)	36.0 (148)
Working 20-34hpw	9.2 (38)	1.2 (5)	10.5 (43)
Working < 20hpw	3.6 (15)	.2 (1)	3.9 (16)
Currently sick leave	1.0 (4)	-	1.0 (4)
Special Employment	.2 (1)	-	.2 (1)
Unemployed	3.2 (13)	.7 (3)	3.9 (16)
Unable to work	2.2 (9)	.7 (3)	2.9 (12)
Retired	26.8 (110)	4.9 (20)	31.6 (130)
Student	7.1 (29)	.7 (3)	7.8 (32)
Homemaker			2.2 (9)
Total (N)	345	66	411
Missing (N)	34	2	36

525

Education Level	% NL (N)	% BE (N)	% Total (N)
None, not in school	.3 (1)	.3 (1)	.5 (2)
At school currently	2.8 (11)	-	2.8 (11)
Primary education	5.3 (21)	1.5 (6)	6.9 (27)
Secondary education	17.3 (68)	8.4 (33)	25.7 (101)
Post Highschool	33.1 (130)	2.3 (9)	35.4 (139)
University/ College	24.7 (97)	4.1 (16)	28.8 (113)
Total (N)	328	65	393
Missing (N)	51	3	54

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Marital staus	% NL (N)	% BE (N)	% Total (N)
Never Married	23.6 (99)	3.6 (15)	27.2 (114)
Married	41.3 (173)	7.9 (33)	49.2 (206)
Living Together	5.5 (23)	1.9 (8)	7.4 (31)
Divorced/ Seperated	8.6 (36)	1.7 (7)	10.3 (43)
Widowed	5.0 (21)	.7 (3)	5.7 (24)
Total (N)	352	66	418
Missing (N)	27	2	49

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530 **Table 3a TBI-related patient demographics, presenting percentages and numbers**
531

Type of hospital stay	% NL (N)	% BE (N)	% Total (N)
<i>ER</i>	13 (58)	.4 (2)	13.4 (60)
<i>Admission</i>	47.0 (210)	5.6 (25)	52.6 (235)
<i>ICU</i>	24.8 (111)	9.2 (41)	34.0 (152)
% Total (N)	84.8(379)	15.2 (68)	100 (447)

532

Initial Diagnosis	% NL (N)	% BE (N)	% Total (N)
<i>Mild TBI (GCS 13-15)</i>	74.2 (267)	13.6 (49)	87.8 (316)
<i>Moderate TBI (GCS 9-12)</i>	5.0 (18)	.8 (3)	5.8 (21)
<i>Severe TBI (GCS 3-8)</i>	4.7 (17)	1.7 (6)	6.4 (23)
Total (N)	302	58	360
Missing (N)	77	10	87

533

Recovery Status at six months post TBI	% NL (N)	% BE (N)	% Total (N)
<i>Good recovery (GOSE 7-8)</i>	65.1 (235)	44.8 (13)	60.3 (248)
<i>Moderate Disabilities (GOSE 5-6)</i>	28.3 (102)	34.5 (10)	26.2 (112)
<i>Severe Disabilities (GOSE 3-4)</i>	6.6 (24)	20.7 (6)	7.7 (30)
Total (N)	361	29	390
Missing (N)	18	39	57

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537 **Table 3b Patient recovery status at six months TBI by type of hospital stay and initial diagnosis**
538

	Initial Diagnosis (N)	Good Recovery at six months post TBI % (N)	Moderate disability at six months post TBI % (N)	Severe disability at six months post TBI % (N)
ER	<i>Mild TBI (53)</i>	14.7 (46)	2.2 (7)	-
Admission	<i>Mild TBI (180)</i>	44.4 (139)	9.3 (29)	3.8 (12)
	<i>Moderate TBI (5)</i>	.6 (2)	1.0 (3)	-
	<i>Severe TBI (2)</i>	.6 (2)	-	-
ICU	<i>Mild TBI (43)</i>	5.4 (17)	7.3 (23)	1.0 (3)
	<i>Moderate TBI (13)</i>	4.2 (13)	4.2 (13)	.03 (1)
	<i>Severe TBI (17)</i>	1.6 (5)	3.2 (10)	.6 (2)
Total	313	214	81	18

539 Note: percentages are based on the 313 respondents of which all three datatypes could be retrieved.
540

541 **Table 4** individual item descriptives and characteristics
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<i>RPQ item</i>	<i>Mean (SD)</i>	<i>Trimmed Mean</i>	<i>Skewness (Kurtosis)</i>	<i>Item-Total correlation</i>	<i>Alpha if item deleted</i>
1. <i>Headaches</i>	.93 (1.23)	.81	1.13 (.075)	.57	.93
2. <i>Dizziness</i>	.88 (1.18)	.76	1.15 (.242)	.57	.93
3. <i>Nausea/ Vomiting</i>	.28 (1.24)	.8	3.1 (9.5)	.46	.93
4. <i>Noise sensitivity</i>	.92 (1.24)	.8	1.18 (.28)	.66	.93
5. <i>Sleep disturbance</i>	1.06 (1.28)	.95	.94 (-.37)	.60	.93
6. <i>Fatigue</i>	1.76 (1.35)	1.73	.11 (-1.23)	.73	.93
7. <i>Irritable</i>	1.04 (1.219)	.93	.96 (-.075)	.73	.93
8. <i>Depressed</i>	1.0 (1.18)	.89	.98 (-.09)	.72	.93
9. <i>Frustrated</i>	1.1 (1.23)	1.0	.87 (-.32)	.77	.93
10. <i>Poor memory</i>	1.4 (1.3)	1.3	.49 (-.93)	.74	.93
11. <i>Poor concentration</i>	1.3 (1.25)	1.24	.62 (-.68)	.79	.92
12. <i>Taking longer to think</i>	1.4 (1.28)	1.29	.45 (-.98)	.75	.93
13. <i>Blurred vision</i>	.67 (1.05)	.55	1.57 (1.63)	.55	.93
14. <i>Light sensitivity</i>	.55 (.97)	.42	1.92 (3.08)	.60	.93
15. <i>Double vision</i>	.4 (.9)	.25	2.57 (6.18)	.43	.93
16. <i>Restlessness</i>	.95 (1.17)	.84	1.06 (.107)	.78	.93

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 544 *Note: N=439; the number of missing values was 8, the minimum value 0, and the maximum value 4 for all items*
 545