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External Validation of a Clinical Prediction Rule to Predict Full Recovery and Ongoing Moderate/Severe Disability Following Acute Whiplash Injury

● **STUDY DESIGN:** Retrospective secondary analysis of data.

● **OBJECTIVES:** To investigate the external validity of the whiplash clinical prediction rule (CPR).

● **BACKGROUND:** We recently derived a whiplash CPR to consolidate previously established prognostic factors for poor recovery from a whiplash injury and predicted 2 recovery pathways. Prognostic factors for full recovery were being less than 35 years of age and having an initial Neck Disability Index (NDI) score of 32% or less. Prognostic factors for ongoing moderate/severe pain and disability were being 35 years of age or older, having an initial NDI score of 40% or more, and the presence of hyperarousal symptoms. Validation is required to confirm the reproducibility and accuracy of this CPR. Clinician feedback on the usefulness of the CPR is also important to gauge acceptability.

● **METHODS:** A secondary analysis of data from 101 individuals with acute whiplash-associated disorder who had previously participated in either a randomized controlled clinical trial or prospective cohort study was performed using accuracy statistics. Full recovery was defined as NDI score at 6 months of 10% or less, and ongoing moderate/

severe pain and disability were defined as an NDI score at 6 months of 30% or greater. In addition, a small sample of physical therapists completed an anonymous survey on the clinical acceptability and usability of the tool.

● **RESULTS:** The positive predictive value of ongoing moderate/severe pain and disability was 90.9% in the validation cohort, and the positive predictive value of full recovery was 80.0%. Surveyed physical therapists reported that the whiplash CPR was simple, understandable, would be easy to use, and was an acceptable prognostic tool.

● **CONCLUSION:** External validation of the whiplash CPR confirmed the reproducibility and accuracy of this dual-pathway tool for individuals with acute whiplash-associated disorder. Further research is needed to assess prospective validation, the impact of inclusion on practice, and to examine the efficacy of linking treatment strategies with predicted prognosis.

● **LEVEL OF EVIDENCE:** Prognosis, level 1b.
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● **KEY WORDS:** Neck Disability Index, prediction, prognosis, whiplash-associated disorder

Understanding recovery pathways and optimal treatment for individuals with whiplash-associated disorder (WAD) continues to be a challenge. Over the past few decades, recovery rates have remained unchanged, with approximately 50% of individuals continuing to experience ongoing pain and disability at 6 months postinjury.^{14,32} Given the personal and societal burden associated with this long-term disability, research has focused on early identification of individuals at risk of ongoing symptoms to help understand recovery pathways and subsequently develop more targeted treatments.³²

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We recently derived a whiplash clinical prediction rule (CPR) that consolidated previously identified prognostic factors for poor recovery following whiplash injury to predict 2 recovery pathways: full recovery and ongoing moderate/severe pain and disability (FIGURE).²⁶

Prognostic CPRs quantify the contributions of relevant prognostic factors to calculate a numeric predicted probability of the outcome of interest.^{1,19} Clinical prediction rules are developed using rigorous methodological standards and have the potential to help clinicians make decisions regarding prognosis and researchers develop targeted therapies.^{12,25} Methodological standards exist for each step in the development of a CPR: derivation, validation, and impact analysis.^{1,19,25} While the derivation of a CPR provides information about potentially important prognostic factors, the validation process ensures that associations between given predictors and outcomes are not primarily due to chance and are not unique to the derivation population.^{19,29} Without validation, the decision to incorporate CPRs into clinical practice is potentially premature.^{2,11,29} Many statistically derived CPRs are not validated, and those that are validated often fail to perform adequately when tested in a new population.^{1,2,11,29}

The process for the derivation of the dual-pathway whiplash CPR followed methodological recommendations.²⁶ The positive predictive value (PPV) for ongoing moderate/severe pain and disability was 71% in patients who were 35 years or older, reported initially higher levels of neck disability (Neck Disability Index [NDI] score of 40% or greater), and reported hyperarousal symptoms of 6 or greater on the hyperarousal subscale of the Posttraumatic Stress Diagnostic Scale (PDS), a substantial increase from the estimated 25% indicated in previous research.^{7,15,27,28} In addition, meeting the simple criteria of an NDI score of 32% or lower and an age of less than 35 years in the acute phase provided an estimated probability of full recovery of 71%, an increase from 50% indicated in previous

research.^{3,15,27} To ensure that the CPR is accurate and reproducible, the next step in the development of the whiplash CPR is external validation.^{1,4,25,29}

Methodological standards for the validation of a CPR include the use of a series of studies across varied environments.^{11,19,25} Initial external validation includes application of the CPR to a new patient population. Subsequent validation studies may include prospective application in new clinical settings by clinicians different from those involved in the derivation study.^{1,19,29} Evaluation of the CPR's clinical usefulness and acceptability is also important.^{1,17,33} The primary aim of the present study was to investigate the external validity of the whiplash CPR. The secondary aim was to gather preliminary feedback from clinicians about the acceptability and usability of the tool.

METHODS

A SECONDARY ANALYSIS OF DATA from 2 cohorts independent of the derivation cohort was performed to investigate the external validity of the recently developed dual-pathway whiplash CPR (FIGURE).²⁶ Both cohorts were recruited in Brisbane, Australia, ethical approval was gained from the University of Queensland Medical Research Ethics Committee, and all participants provided signed informed consent.^{7,14}

Participants With WAD

Information regarding inclusion and exclusion criteria and procedures for the 2 primary cohort studies have been published previously.^{7,14} All participants for the primary cohort studies were recruited from accident and emergency departments, through referral by primary care practitioners (eg, general practitioners and physical therapists), and through advertising in the popular press. Volunteers for the primary cohort studies were eligible if they were between 18 and 65 years of age and had acute neck pain (less than 4 weeks in duration) as a result of a motor vehicle collision that was classifi-

able as WAD grade II. Volunteers were ineligible if their neck pain was not related to a motor vehicle collision or was classifiable as WAD grades I, III, or IV; if there was a previous history of whiplash or other neck pain for which treatment had been sought; or if they lacked fluency in spoken and written English for questionnaire completion.

Cohort 1 was from a prospective longitudinal study that examined prognostic factors for poor functional recovery following a whiplash injury.⁷ The design of the study adhered to STROBE criteria for cohort studies.³⁰ Cohort 2 comprised participants from the control intervention (usual care) of a randomized controlled clinical trial.¹⁴ The trial was registered with the Australian New Zealand Clinical Trials Registry (ACTRN12605000109606). The randomized controlled clinical trial evaluated the effectiveness of individualized multiprofessional management (medical, physiotherapeutic, psychological) compared with usual care. Although the pragmatic multidisciplinary interventions tested in this trial produced no advantage over usual care, to avoid the potential influence of targeted therapy, only the participants receiving usual care (control group) were included within the current validation study.

Whiplash CPR

The whiplash CPR predicts 2 recovery pathways following an acute whiplash injury: ongoing moderate/severe pain and disability were predicted in the presence of baseline NDI scores of 40% or greater, age of 35 years or older, and a hyperarousal subscale symptom score on the PDS of 6 or greater. Full recovery was predicted in individuals less than 35 years of age with a baseline NDI score of 32% or less.²⁶

The method used for the derivation of the whiplash CPR has been explained previously.²⁶ Briefly, the NDI was the dependent variable. The NDI is a valid, reliable, and responsive measure.³¹ An overall score (out of 100) is calculated by

totaling responses to 10 questions, each with 6 potential Likert-type responses (0 is no disability and 5 is total disability), and multiplying the sum by 2 to yield a percentage. Previous research has suggested that an NDI score of 30% or greater is indicative of moderate to severe levels of pain-related disability and that an NDI score of 10% or less indicates full recovery.^{18,28} In the present study, these criteria were used with the NDI measured at 6 months to classify participants as experiencing ongoing moderate/severe pain and disability (NDI, 30% or greater), continued mild to moderate pain and disability (NDI, 11%-29%), or fully recovered (NDI, 10% or less).

The derivation of the whiplash CPR included analysis of 8 biopsychosocial

variables shown in previous research to be predictive of recovery following a whiplash injury: initial NDI, age, cold pain threshold, initial neck pain (visual analog scale [VAS]), neck range of movement, posttraumatic stress symptoms, sex, and presence of headache.^{15,27,28,32} Predictor variables were measured at baseline.²⁶

Briefly, measurement methods were age measured in years, cold pain thresholds measured over the mid cervical spine using the THERMOTEST system (Somedic Production AB, Sollentuna, Sweden),²⁸ neck pain measured using an 11-point VAS with anchors of 0 (no pain) and 10 (worst pain imaginable), and range of movement calculated as total neck rotation (the sum of left and right neck rotation, flexion, and extension)

from measurements using an electromagnetic motion-tracking device (FAS-TRAK; Polhemus Inc, Colchester, VT). Sex failed to meet statistical significance within the derivation study, and presence of headache was measured differently in the validation cohorts, hence both of these variables were omitted from the present study.

Posttraumatic stress symptoms were measured using the valid and reliable 49-item PDS.⁸ Following a standard multiple regression, only the hyperarousal subscale of the PDS (5 items) was found to be significantly predictive and hence was the only subscale included in the derivation analyses and, subsequently, in the present study. The hyperarousal subscale includes questions about trouble sleeping, irritability, difficulties concentrating, being overly alert, and being easily startled. Frequency of symptoms is assessed for each question using a Likert-type format, with options ranging from 0 (not at all or only 1 time) to 3 (5 or more times a week/almost always), resulting in a subscale score of 0 to 15.

Preliminary Feedback: Clinician Acceptability

Practitioners who might decide to use this tool in the future were asked to give preliminary feedback about its clinical acceptability and usefulness. Although not totally representative of clinicians who might treat individuals with an acute whiplash injury, a few groups were selected that represented clinicians who had and had not been involved in clinical research trials involving whiplash injuries. These groups included physical therapists practicing in Queensland, Australia who were titled “musculoskeletal physiotherapists” by the Australian Physiotherapy Association and whose contact e-mail details were available to the public (n = 74); chiropractors registered with the Chiropractors’ Association of Australia practicing in Queensland, Australia whose contact details were available to the public (n = 61); and physical therapists identified as experts in the area of

whiplash research from Perth and Melbourne, Australia, and Canada (n = 10).

A written format of the whiplash CPR with instructions for use and an anonymous survey were sent via e-mail. The survey collected background information on number of patients with whiplash seen each month and included Likert-type questions about the acceptability and usability of the whiplash CPR. Open-ended questions collected information on preferred mode of delivery, benefits and disadvantages of the whiplash CPR, and potential difficulties in using this type of tool. A reminder e-mail was sent 10 days after the initial request.

It was understood that interpretation of feedback would be limited because clinicians had not yet used the tool with patients. Despite this, we felt it important to seek some feedback before embarking on the next stage of CPR development, which was prospective evaluation followed by evaluation of impact on practice. If there were concerns about the content, acceptability, and/or usability, changes to the tool would be needed. Ethical approval was granted by the University of Queensland Medical Research Ethics Committee (number 2013000671).

Data Analysis

The predictive ability of the whiplash CPR to identify those likely to fully recover and those likely to experience ongoing moderate/severe pain and disability was assessed using accuracy statistics for the positive case (specificity, sensitivity, likelihood ratios [LRs], PPV) with 95% confidence intervals (CIs). Validation accuracy for identification purposes was assessed using revised estimates of the probability (PPV) of participants from the validation cohort who were likely to experience ongoing moderate/severe pain and disability or likely to fully recover, using the CPR.²⁶ The Hosmer-Lemeshow statistic was used to evaluate the appropriateness of the whiplash CPR models, in which a non-significant test indicates good model fit.²⁰

Several analyses were also performed to compare the validation study popula-

TABLE 1		
ACCURACY STATISTICS FOR ONGOING MODERATE/ SEVERE PAIN AND DISABILITY FOR THE VALIDATION AND DERIVATION POPULATIONS*		
	Validation Population	Derivation Population
Sensitivity	43.5 (22.9, 65.1)	43.5 (31.8, 54.9)
Specificity	98.7 (92.9, 99.9)	93.8 (89.1, 96.6)
Positive likelihood ratio	33.9 (4.6, 251.2)	7.0 (3.8, 12.9)
Negative likelihood ratio	0.6 (0.4, 0.8)	0.6 (0.5, 0.7)
Positive predictive value	90.9 (58.7, 98.5)	71.4 (55.1, 83.8)
*Values in parentheses are 95% confidence interval.		

TABLE 2		
ACCURACY STATISTICS FOR THE LIKELIHOOD OF FULL RECOVERY FOR THE VALIDATION AND DERIVATION POPULATIONS*		
	Validation Population	Derivation Population
Sensitivity	54.9 (40.5, 68.6)	45.3 (35.1, 54.4)
Specificity	86.0 (72.6, 93.7)	84.5 (77.3, 89.8)
Positive likelihood ratio	3.9 (1.9, 8.1)	2.9 (1.9, 4.5)
Negative likelihood ratio	0.5 (0.4, 0.7)	0.6 (0.5, 0.8)
Positive predictive value	80.0 (62.5, 91.7)	70.7 (58.9, 80.3)
*Values in parentheses are 95% confidence interval.		

tion with the derivation study population. First, *t* tests and chi-square analyses for independent samples were performed to identify any statistically significant differences in terms of 6 of the 8 possible predictor variables investigated for the CPR in the derivation study (eg, initial NDI, neck pain VAS, age, range of movement, cold pain threshold, hyperarousal). An additional chi-square analysis was performed to determine whether possible differences in recovery rates existed between the 2 study populations: specifically, differences in the proportion of participants who were classified at 6 months as being fully recovered (NDI, 10% or less), continuing to experience mild to moderate levels of disability (NDI, 11%-29%), and having ongoing moderate/severe disability (NDI, 30% or greater).

Second, *t* tests and chi-square analyses for independent samples were

conducted to investigate potential differences between the 2 cohorts within the validation populations, with regard to both the whiplash CPR variables as well as additional baseline and background variables. Chi-square analysis was also performed to examine the possible differences in recovery rates between the 2 cohorts.

Finally, a descriptive analysis was used to summarize feedback from the acceptability survey.

Sample Size

If intervals based on a 95% confidence level were constructed from at least 71 samples drawn from a population with a PPV proportion of 55%, where the distance from 55% to the lower limit was 10.5%, then 95% of these intervals would contain the actual population proportion. The calculations were based on PPV percent statistics reported for the whiplash

TABLE 3

COMPARISONS BETWEEN PARTICIPANTS IN THE VALIDATION AND DERIVATION STUDIES

Factor	Validation Participants (n = 101)*	Derivation Participants (n = 262)*	P Value
Initial NDI (0-100)	31.5 ± 18.5	32.9 ± 17.5	.475
Age, y	34.2 ± 10.7	37.1 ± 14.2	.034
PDS hyperarousal subscale (0-15)	4.3 ± 3.9	4.8 ± 3.8	.271
Neck range of motion, deg†	191.2 ± 53.5	201.4 ± 63.4	.169
Initial visual analog scale for pain (0-10)	3.5 ± 2.1	4.2 ± 2.1	.005
Cold pain threshold, °C	13.2 ± 7.7	14.4 ± 7.7	.274

Abbreviations: NDI, Neck Disability Index; PDS, Posttraumatic Stress Diagnostic Scale.

*Values are mean ± SD.

†Sum of left and right neck rotation, flexion, and extension.

TABLE 4

PARTICIPANTS CLASSIFIED BY RECOVERY CATEGORY AT 6 MONTHS FOR EACH STUDY POPULATION*

	Validation Participants (n = 101)	Derivation Participants (n = 262)
Fully recovered (NDI, ≤10%)	51 (50)	120 (46)
Continued mild to moderate pain and disability (NDI, 11%-29%)	27 (27)	73 (28)
Ongoing moderate/severe pain and disability (NDI, ≥30%)	23 (23)	69 (26)

Abbreviation: NDI, Neck Disability Index.

*Values are n (%).

The PPV for the fully recovered pathway with the validation population was 80.0%, slightly better than the PPV for the derivation cohort (70.7%)²⁶ (TABLE 2). The Hosmer-Lemeshow test resulted in a *P* value of .081 ($\chi^2 = 14.042$, *df* = 8), indicating support for the model. The very high specificity indicates that most of those who had some disability did not meet the 2-factor criteria. As a result, an individual who met these criteria was likely to recover. The moderate sensitivity, however, indicates that some individuals who did not meet the criteria had varied recovery pathways, including full recovery. A positive LR between 2 and 5 represents a small but sometimes meaningful shift in probability.¹³

Validation and Derivation Populations

The results of independent *t* tests showed that the derivation and validation populations differed significantly at baseline in terms of age and initial pain levels as assessed with a VAS (TABLE 3). However, eta-squared showed that the magnitudes of the differences in the means for both age and initial VAS were small and therefore unlikely to be clinically significant. Chi-square analysis showed no difference between the proportion of women in the derivation (*n* = 169/262) and the validation populations (*n* = 67/101) ($\chi^2 = 0.108$, *df* = 1, *P* = .74).

The validation population had recovery rates similar to those of the established prevalence^{15,27} and to the derivation cohort²⁶ at 6 months: 50% (*n* = 51) were fully recovered, 27% (*n* = 27) continued to experience milder levels of disability, and 23% (*n* = 23) experienced ongoing moderate/severe pain and disability (TABLE 4). Chi-square analysis showed that there were no significant differences in the proportion of participants who fully recovered or those who experienced ongoing moderate/severe pain and disability between the validation and derivation study participants ($\chi^2 = 0.74$, *df* = 2, *P* = .69).

Validation Cohorts

The results of independent-sample *t* tests showed no significant differences

CPR derived by Ritchie et al,²⁶ where PPV was estimated to range from 55% (the lowest reported 95% confidence value for a PPV) and the smallest reported distance of 10.5% to a lower 95% confidence limit. The calculation allows for the worst PPV outcome with high expectations of accuracy. If the PPV were greater than 55% or accuracy less stringent, then a smaller sample size would suffice. Thus, 101 participants available for the current study met this requirement.

RESULTS

DATA WERE ANALYZED FROM 101 individuals involved in 2 independent cohort studies. Six-month data applicable to the validation of the whiplash CPR were available for 95% (*n* = 53) of cohort 1 participants and for 92% (*n* = 48) of cohort 2 participants.^{7,14}

Whiplash CPR External Validation

The PPV for the pathway of ongoing moderate/severe pain and disability for the validation population was 90.9%, which is greater than the derivation PPV of 71.4%²⁶ (TABLE 1). The Hosmer-Lemeshow test resulted in a *P* value of .649 ($\chi^2 = 5.980$, *df* = 8), indicating support for the model. The very high specificity indicates that almost all of those who did not have moderate/severe disability at 6 months did not meet the 3-factor criteria. As a result, an individual who met the 3-factor criteria was likely to experience ongoing moderate/severe pain and disability. However, the lower sensitivity indicates that some of the individuals who had ongoing moderate/severe pain and disability did not meet the criteria. The positive LR of 33.9 indicates a very large shift in probability¹³; however, the large CI indicates a lower degree of precision.

TABLE 5

COMPARISONS BETWEEN THE 2 COHORTS
OF PARTICIPANTS IN THE PRESENT STUDY

Factor	Cohort 1 (n = 53)*	Cohort 2 (n = 48)*	P Value
Initial NDI (0-100)	33.8 ± 19.6	29.0 ± 17.1	.20
Age, y	33.4 ± 9.4	35.2 ± 11.9	.393
PDS hyperarousal subscale (0-15)	3.9 ± 4.1	4.8 ± 3.7	.281
Neck range of motion, deg [†]	195.9 ± 50.3	186.0 ± 56.9	.356
Initial visual analog scale for pain (0-10)	3.6 ± 2.3	3.3 ± 1.9	.516
Cold pain threshold, °C	13.3 ± 7.9	13.1 ± 7.6	.893

Abbreviations: NDI, Neck Disability Index; PDS, Posttraumatic Stress Diagnostic Scale.

*Values are mean ± SD.

[†]Sum of left and right neck rotation, flexion, and extension.

between the 2 cohorts in the baseline variables examined (TABLE 5). Chi-square analysis showed that there were significantly more women in cohort 1 (n = 41/53) compared with cohort 2 (n = 26/48) ($\chi^2 = 6.07$, $df = 1$, $P = .014$).

Chi-square analysis showed that there were no significant differences in the proportion of participants who fully recovered and those who experienced ongoing moderate/severe pain and disability between the validation study cohorts ($\chi^2 = 2.10$, $df = 2$, $P = .35$). Hence, it was deemed appropriate to externally validate the whiplash CPR with these cohorts.

Preliminary Feedback: Clinician Acceptability

Twenty-six (31%) physical therapists and 2 (3%) chiropractors returned the clinician acceptability survey. Due to the very small number of responding chiropractors, these responses were not included in the descriptive summary of feedback. Of the responding physical therapists, 11 were men and 15 were women. Respondents reported that 1 to 30 patients with a whiplash injury were seen in their practice per month. Collation of data on clinical acceptability showed that all respondents agreed or strongly agreed that the whiplash CPR gathered relevant information and that it was understandable for clinicians. Ninety-six percent (n = 25) agreed or strongly agreed that the whiplash CPR would be easy to use, 88% (n =

23) agreed that it would be understandable for patients, and 85% (n = 22) agreed that it would help direct therapy. Ninety-six percent (n = 25) already clinically used the NDI to assess neck disability in patients with WAD. Clinicians were also asked about their preferred format for the administration of the whiplash CPR. The survey responses were reported in a written survey (73%, n = 19), followed by a smartphone application survey (42%, n = 11), web-based survey (42%, n = 11), and software-based survey (23%, n = 6).

DISCUSSION

EXTERNAL VALIDATION CONFIRMED the reproducibility and accuracy of the recently derived dual-pathway whiplash CPR for individuals seeking health care for neck pain following a motor vehicle collision.²⁶ The whiplash CPR consolidates existing evidence regarding prognostic factors for poor recovery and provides a tool to break down varied recovery patterns into more homogeneous subsets. The PPV of ongoing moderate/severe pain and disability was 90.9% for individuals who were 35 years of age and older with initially higher levels of neck disability and hyperarousal symptoms. In addition, the probability of full recovery was 80.0% for younger individuals with initially lower levels of neck disability. The current study is one of only a few to validate a CPR in the area of musculo-

skeletal pain and, to our knowledge, the only study to validate a CPR for patients with a whiplash injury.^{2,22} Application of this tool with similar patients may help clinicians to make decisions about prognosis and enable interventions to be designed to match patient subgroups.

There has been growing interest in CPRs for the assessment of patients with musculoskeletal pain, with the speculation that therapeutic benefits may be more likely to occur if subgroups of patients are identified and better matched to treatments.^{2,11,22} While, to our knowledge, there is only 1 other derivation-only CPR¹⁰ and 1 stratified risk assessment tool¹⁶ for individuals with a whiplash injury, there has been a surge of CPRs in the area of low back pain.^{9,11,12,23,24} Similar to acute whiplash, nonspecific low back pain is a heterogeneous condition with varied recovery rates, and optimal therapy remains equivocal.^{5,9,11,14,22}

Positive effects of matching therapeutic pathways to predict prognosis based on a CPR have been shown in patients with low back pain.¹² Hill and colleagues¹² subgrouped patients into low-, moderate-, and high-risk categories using a CPR-like prognostic screening tool. Treatments were matched to these categories and unnecessary treatment was avoided in low-risk patients, whereas medium- and high-risk patients were encouraged to access more concerted treatments, thereby improving clinical outcomes. This type of stratified treatment strategy, designed to target patient subgroups such as those identified in the whiplash CPR, may help to optimize recovery. Clinical treatment pathways need to be developed with clinical experts, particularly given the evidence suggesting that early intensive health care for individuals with acute whiplash injury may delay recovery.^{6,21} It may be that specific subsets of patients benefit from specific treatment strategies, whereas other patients may find the same therapies to be detrimental. To avoid unnecessary treatment and reduce practice variability, the recommendation for patients

who meet the full recovery criteria may be minimal treatment and reassurance, whereas a targeted treatment approach involving physical therapy, psychological treatment, and possibly medication may be recommended for patients who meet the criteria for ongoing moderate/severe pain and disability. Recommendations for individuals who do not meet the criteria for either pathway may be initial assessment with reassurance and follow-up as needed to ensure that recovery does take place, and to institute more targeted treatment if it does not. Randomized controlled treatment trials are needed to identify optimal treatment recommendations for each of the recovery pathways identified with the whiplash CPR.

Aligning treatment options to recovery pathways necessitates a CPR with high positive predictive probability and minimal false positives.^{2,18} Previous research suggests that, of individuals who experience an acute whiplash injury, approximately 25% are likely to experience ongoing moderate/severe pain and disability, and 50% are likely to fully recover.^{15,27} Application of the whiplash CPR provides a significantly better prediction of recovery, with 90.9% predictive probability for ongoing moderate/severe pain and disability (**TABLE 1**) and 80.0% predictive probability for full recovery (**TABLE 2**). The specificity of the whiplash CPR for the pathway of ongoing moderate/severe pain and disability was 98.7% (95% CI: 93%, 99%) (**TABLE 1**) and 86.0% (95% CI: 73%, 94%) for the pathway of full recovery (**TABLE 2**). These very high specificities indicate a low false-positive rate and help to rule in the condition. For example, if the patient meets the 3-factor criteria for ongoing moderate/severe pain and disability, the patient is likely to continue to experience moderate/severe pain and disability; or, if a patient meets the 2-factor criteria for full recovery, the patient is likely to fully recover. Likelihood ratios quantify both the direction and magnitude of the shift in probability that an individual will be positive or negative on the reference standard, in

this case the NDI.¹³ It is favorable to have higher LRs and to have a narrow CI (eg, 95% CI) to indicate that the estimate is precise.¹³ The positive LR of 33.9 for the ongoing moderate/severe pain and disability arm indicates a very large shift in probability, and the positive LR of 3.92 for the full recovery arm indicates a smaller shift in probability. The very large 95% CI for the ongoing moderate/severe pain and disability arm indicates a lower degree of precision, whereas the narrow 95% CI for the full recovery arm indicates higher precision. Despite these positive statistical results, the whiplash CPR is a predictive tool; it is not meant to provide mutually exclusive decisions on prognosis but may be an additional tool to support clinical judgment.

Finally, a CPR will only be used if it makes clinical sense, is easy to use, there are no obvious items missing, and the items seem appropriate for the purpose of the rule.^{17,25} The variables within the whiplash CPR fit a biopsychosocial model of recovery, thereby confirming current evidence about recovery from an acute whiplash injury. The simple, 3-step whiplash CPR is designed to be easily completed by the patient and treating clinician. The NDI, the first step, is an outcome measure that is readily available, and 96% (n = 25) of the responding clinicians were already using the NDI to assess neck disability in patients with a whiplash injury. In a written format, the clinician will need to calculate the score on the NDI. If the NDI is 40% or greater, the patient should then complete the 5-question PDS subscale to assess hyperarousal symptoms, and the clinician would simply calculate the total. The 5-question subscale used in the present study is part of the PDS, a valid instrument designed to help with both the detection and symptomatic evaluation of posttraumatic stress disorder symptoms.⁸ While there is a cost associated with using the entire instrument, the 5-question hyperarousal subscale asks generic questions about arousal symptoms and is very similar to other

available tools, for example, the hyperarousal items of the Impact of Event Scale-Revised,³⁴ a free and easily accessible questionnaire. It would thus not be difficult for clinicians to use the 5 hyperarousal symptom items independently of the entire PDS questionnaire.

Although the sample of clinicians surveyed was small, the majority agreed that the whiplash CPR gathered relevant information, would be easy to use, would be understandable for both clinicians and patients, and would help direct therapy. Future studies should gather additional information about acceptability from a wider range of clinicians, including general practitioners and those involved in the acute care of individuals following a whiplash injury, to ensure the acceptability of the whiplash CPR.

The current study was the first of a recommended series of studies needed to comprehensively validate a CPR. As such, there are several limitations to the study. First, although the data were collected prospectively in a new patient cohort, predictive validity was assessed through a retrospective secondary analysis of data. The possibility exists that the predictive validity of the whiplash CPR may be different when used prospectively.^{1,19} Second, this external validation included patients classified as WAD grade II, whereas the derivation study included patients classified as WAD grades I, II, and III. Hence, although most injured people are classified as WAD grade II, the validation sample is not fully representative of the population of patients seeking health care for neck pain following a motor vehicle collision. Third, the derivation and validation of the whiplash CPR involved data from participants enrolled in research studies in Brisbane, Australia. Participants recruited for research studies may respond differently from the general public with regard to components of the whiplash CPR, and the CPR may perform differently in other countries and jurisdictions. Prospective validation with a new cohort of patients fully representative of the

WAD population would be beneficial.¹⁹ Finally, the response rate to the clinician acceptability survey was low for physical therapists and almost nonexistent for chiropractors. Caution is needed when interpreting these data. Despite these limitations, the predictive probabilities found for each pathway through this external validation study were significantly higher than the estimated probabilities suggested in previous research and therefore provide a level of evidence sufficient for clinicians to consider using the whiplash CPR with patients similar to those in the derivation and validation studies.¹⁹

CONCLUSION

EXTERNAL VALIDATION CONFIRMED that the whiplash CPR is reproducible and accurate in predicting 2 prognostic pathways following an acute whiplash injury. Further research is needed to assess prospective validation, the impact of inclusion on practice, and to examine the efficacy of linking treatment strategies with predicted prognosis. ●

KEY POINTS

FINDINGS: The predictive probabilities found for both pathways of the whiplash CPR following external validation were significantly greater than the estimated probabilities suggested in previous research.

IMPLICATIONS: The whiplash CPR may be considered reproducible and accurate for individuals seeking health care for neck pain following a motor vehicle collision. Application of this tool with similar patients may help clinicians with decisions about prognosis and may help researchers develop targeted therapies to better match patient subgroups.

CAUTION: The whiplash CPR has not been prospectively validated with patients classified as WAD grades I, II, and III outside of Brisbane, Australia. Feedback on clinical acceptability of this tool is limited to a small group of physical therapists. The wide 95% CIs for the

ongoing moderate/severe pain and disability pathway indicate less-than-ideal precision.

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