Simplifying the use of prognostic information in patients with traumatic brain injury.

Citation for published version:

Digital Object Identifier (DOI):
10.3171/2018.5.JNS181386

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published In:
Journal of Neurosurgery

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
Simplifying the use of prognostic information in patients with traumatic brain injury

TO THE EDITOR: We read with great interest the articles recently published by Brennan et al.² and Murray et al.⁴ concerning the Glasgow Coma Scale-Pupils (GCS-P) score, and we commend the authors on the efforts undertaken to improve traumatic brain injury (TBI) prognostication (Brennan PM, Murray GD, Teasdale GM: Simplifying the use of prognostic information in traumatic brain injury. Part 1: The GCS-Pupils score: an extended index of clinical severity. J Neurosurg 128:1612–1620, June 2018; Murray GD, Brennan PM, Teasdale GM: Simplifying the use of prognostic information in traumatic brain injury. Part 2: Graphical presentation of probabilities. J Neurosurg 128:1621–1634, June 2018). In this regard, we would like to raise some issues that could contribute to the refinement of the new proposed tool.

The basic assessment of a prognostic score should comprise at least its 1) calibration (the agreement between observed outcomes and predictions); 2) discrimination (ability to distinguish between those with or without the outcome); and 3) overall performance (global accuracy). The GCS-P models were adequately calibrated, as depicted on the graphics; however, no discrimination statistics were reported. It would be interesting to report the area under the receiver operating characteristic curve (AUC-ROC) for the GCS-P model and the GCS-P plus Age & CT models. This is the most widely used discriminatory capacity measure in the medical literature and its interpretation is broadly understood, which facilitates comparisons between models.¹

The Nagelkerke R² (similarly to all pseudo R² statistics) is an overall performance measure. Although valid, it lacks intuitive understanding and its direct interpretation as “the proportion of variability in outcome that is explained by the logistic regression model” is highly vulnerable to criticism and statistical reasoning (most pseudo R² statistics do not have 1.0 as the maximum value, and even the adjusted Nagelkerke R² is not adequately scaled).³

The Nagelkerke R² differences between the GCS-P and GCS (for modeling death: 18.4 vs 15.5, difference 2.9; for favorable outcome: 22.2 vs 19.8, difference 2.4) are similar to those between the Corticosteroid Randomisation After Significant Head Injury (CRASH) CT model and the GCS-P/Age/CT chart (death: 41.9 vs 39.7, difference 2.2; favorable outcome: 42.1 vs 39.7, difference 2.4). Thus, some may find it difficult to understand why the GCS-P was considered superior to the GCS but the GCS-P/Age/CT chart was considered sufficiently non-inferior to the CRASH-CT model. The AUC-ROC analysis could further elucidate this question.

Considering that the incidence of pupil alteration is higher in patients with severe TBI, we could be losing valuable prognostic information for mild and moderate TBI if the GCS-P model were to be routinely recommended over the CRASH model. On the general pooled sample reported in the paper, the CRASH model was at least marginally superior to the new proposed tool. Could we hypothesize that this superiority would be higher for nonsevere TBI? It would be enlightening to see a stratified analysis by TBI severity.

Nowadays medical apps are widely used for instant prognostic score calculations, and the CRASH/International Mission for Prognosis and Clinical Trials in Traumatic Brain Injury (IMPACT) models can be assessed as fast as any chart.⁵ Many prognostic scores and decision rules are already routinely used in the emergency department and intensive care unit by other specialties.

In conclusion, although it may be too early to endorse the GCS-P/Age/CT model over the CRASH/IMPACT CT models for regular TBI management, it is indeed an interesting alternative approach and could be a step forward to advance prognostic reasoning and more rational decision making.

Davi J. Fontoura Solla, MD
Manoel Jacobsen Teixeira, MD, PhD
Wellington Silva Paiva, MD, PhD
Universidade de São Paulo, São Paulo, Brazil

References
4. Murray GD, Brennan PM, Teasdale GM: Simplifying the


Disclosures
The authors report no conflict of interest.

Correspondence
Davi J. Fontoura Solla: davisolla@hotmail.com.

Response
We thank Dr. Solla and colleagues for their interest in our papers. We agree that there are many different measures available to assess performance of a prognostic model, each with its own strengths and limitations. Many papers report detailed technical evaluations of prognostic models in TBI, and we have ourselves contributed to many such studies. We nonetheless very deliberately avoided the temptation of writing our two papers as statistical treatises, but instead aimed to focus on practical relevance, and we presented our results in terms of a simple overall measure of performance.

The fundamental thrust of our papers was to point out that utility and acceptability are at least as important as the “technical” performance of a prognostic model. A well-calibrated and powerfully discriminatory model is futile if its complexity deters its use in practice. In spite of the authors’ assertions to the contrary, our perception echoes previous views referred to in our papers and is supported by an informal survey of UK neurosurgical units, which found that statistical models are not widely used in TBI. Our suggestion is that a simple chart might make the breakthrough that leads to prognostic models for TBI becoming incorporated widely in clinical practice.

Moreover, we do not consider the performance of our charts to be “sufficiently non-inferior” to the CRASH models. What we do is present a quantification of the inevitable tradeoff in performance between complexity on the one hand and ease of use and interpretation on the other. It will be for clinicians caring for a patient to decide if the simpler, less-powerful approach is to be preferred on the grounds of utility.

We present in Table 1 the relevant data for the “area under the curve” for the receiver operating characteristic, or the c-statistic as it is also known. As might be expected, these closely mirror the results expressed in terms of Nagelkerke’s R². In particular, with both the IMPACT and CRASH models, the performance of the Age/GCS-P/CT chart lies between that of the simpler and more complex models, and is generally closer to that of the more complex models than to that of the simpler models.

We have severe reservations as to the wisdom of using measures of model performance within subsets stratified by severity, for if one wished to optimize the performance of a model within a subset of patients defined by severity then one would develop a model restricted to those patients. Nevertheless, in the interests of transparency, we present in Table 2 such an analysis with the CRASH data, stratifying patients as GCS ≤ 8 (a widely used definition of “severe” TBI) versus GCS > 8. This shows that the GCS-P/Age/CT chart still performs well relative to the CRASH models for the patients with a better prognosis.

Gordon D. Murray, MA, PhD
Usher Institute of Population Health Sciences and Informatics,
University of Edinburgh, United Kingdom

Paul M. Brennan, MBBChir, FRCS, PhD
Centre for Clinical Brain Sciences,
University of Edinburgh, United Kingdom

Graham M. Teasdale, MBBS, FRCP, FRCS
Institute of Health and Wellbeing,
University of Glasgow, United Kingdom

TABLE 1. Predictive yield for a range of models including GCS

<table>
<thead>
<tr>
<th>Model</th>
<th>Modeling Death</th>
<th>Modeling Favorable Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>GCS score as linear variable</td>
<td>0.717</td>
<td>0.731</td>
</tr>
<tr>
<td>GCS-P as linear variable</td>
<td>0.731</td>
<td>0.741</td>
</tr>
<tr>
<td>GCS-P as linear variable &amp; age as linear variable</td>
<td>0.778</td>
<td>0.787</td>
</tr>
<tr>
<td>GCS-P as linear variable &amp; age as linear variable &amp; CT findings in 3 groups</td>
<td>0.806</td>
<td>0.803</td>
</tr>
</tbody>
</table>

TABLE 2. Predictive yield for a range of models based on a stratified analysis of the CRASH data set

<table>
<thead>
<tr>
<th>Model</th>
<th>Modeling Death</th>
<th>Modeling Favorable Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>CRASH patient group w/ GCS score ≤8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CRASH basic model</td>
<td>0.742</td>
<td>0.746</td>
</tr>
<tr>
<td>CRASH CT model</td>
<td>0.790</td>
<td>0.780</td>
</tr>
<tr>
<td>Age/GCS-P/CT chart</td>
<td>0.772</td>
<td>0.767</td>
</tr>
<tr>
<td>CRASH patient group w/ GCS score &gt;8</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CRASH basic model</td>
<td>0.778</td>
<td>0.759</td>
</tr>
<tr>
<td>CRASH CT model</td>
<td>0.824</td>
<td>0.783</td>
</tr>
<tr>
<td>Age/GCS-P/CT chart</td>
<td>0.815</td>
<td>0.767</td>
</tr>
</tbody>
</table>
References


INCLUDE WHEN CITING
Published online August 3, 2018; DOI: 10.3171/2018.6.JNS181473.
©AANS 2018, except where prohibited by US copyright law