



● LETTER TO THE EDITOR

Calcitonin gene-related peptide and traumatic brain injury

Dear editor,

It is with great interest that we read the article “Relationship of calcitonin gene-related peptide with disease progression and prognosis of patients with severe traumatic brain injury” (Chen et al., 2018). In this study, the authors evaluated 121 patients who were divided into mild/moderate traumatic brain injury (TBI) ($n = 61$), severe TBI ($n = 35$) and control ($n = 25$) groups, and measured serum levels of calcitonin gene-related peptide (CGRP) and serum endothelin-1 (ET-1). They found that low levels of CGRP and high levels of ET-1 were associated with high mortality at 6 months.

Identification of morphological abnormalities on CT scans is very important for evaluating patients with TBI because different diagnoses are made based on different imaging findings (Maas et al., 2005).

We found that the study lacked the information about the severity of lesions that the patients presented, and about the treatment performed. Furthermore, there was no record on whether the patients presented with extracranial lesions, which could be associated with a worse prognosis (van Leeuwen et al., 2012).

CGRP is known to be a potent vasodilator and also to grant protective mechanisms in physiological and pathological conditions for the cardiovascular system and wound healing (Russell et al., 2014). Although the findings show an important relationship between serum CGRP reduction and increased mortality, the study fails to elaborate a pathophysiological explanation to justify such findings.

Finally, we would like to congratulate the authors for the interesting work. It is of highest importance to develop more studies regarding this subject. We expect that, as they are able to replicate the data presented, they may come to promote more understanding of the pathophysiology related to the reduction of CGRP.

Saul Almeida da Silva*, **Almir F. de Andrade**,
Robson Luis Oliveira de Amorim, **Wellingson S. Paiva**

Division of Neurological Surgery, Hospital das Clinicas
HCMFUSP, Universidade de Sao Paulo, SP, Brazil

*Correspondence to: Saul Almeida da Silva, MD,
alm.saul@gmail.com.

orcid: 0000-0003-3766-9990 (Saul Almeida da Silva)

Received: September 20, 2018

Accepted: October 1, 2018

doi: 10.4103/1673-5374.244801

da Silva SA, de Andrade AF, de Amorim RLO, Paiva WS (2019) Calcitonin gene-related peptide and traumatic brain injury. *Neural Regen Res* 14(4):736.

Conflicts of interest: The authors declare no conflicts of interest.

References

Chen LX, Zhang WF, Wang M, Jia PF (2018) Relationship of calcitonin gene-related peptide with disease progression and prognosis of patients with severe traumatic brain injury. *Neural Regen Res* 13:1782-1786.

Maas AI, Hukkelhoven CW, Marshall LF, Steyerberg EW (2005) Prediction of outcome in traumatic brain injury with computed tomographic characteristics: a comparison between the computed tomographic classification and combinations of computed tomographic predictors. *Neurosurgery* 57:1173-1182.

Russell FA, King R, Smillie SJ, Kodji X, Brain SD (2014) Calcitonin gene-related peptide: physiology and pathophysiology. *Physiol Rev* 94:1099-1142.

van Leeuwen N, Lingsma HF, Perel P, Lecky F, Roozenbeek B, Lu J, Shakur H, Weir J, Steyerberg EW, Maas AI (2012) International Mission on Prognosis and Clinical Trial Design in TBI Study Group; Corticosteroid Randomization After Significant Head Injury Trial Collaborators; Trauma Audit and Research Network. Prognostic value of major extracranial injury in traumatic brain injury: an individual patient data meta-analysis in 39,274 patients. *Neurosurgery* 70:811-818.