



OPEN ACCESS

EDITED AND REVIEWED BY
Manuela Mengozzi,
Brighton and Sussex Medical School,
United Kingdom

*CORRESPONDENCE

Shireen Mohammad
✉ s.mohammad@qmul.ac.uk
Christoph Thiemermann
✉ c.thiemermann@qmul.ac.uk

RECEIVED 24 July 2025
ACCEPTED 08 August 2025
PUBLISHED 21 August 2025

CITATION

Mohammad S, O'Riordan CE, Verra C,
Aimaretti E, Alves GF, Dreisch K, Evenäs J,
Gena P, Tesse A, Rützler M, Collino M,
Calamita G and Thiemermann C (2025)
Correction: RG100204, a novel Aquaporin-9
inhibitor, reduces septic cardiomyopathy and
multiple organ failure in murine sepsis.
Front. Immunol. 16:1672460.
doi: 10.3389/fimmu.2025.1672460

COPYRIGHT

© 2025 Mohammad, O'Riordan, Verra,
Aimaretti, Alves, Dreisch, Evenäs, Gena, Tesse,
Rützler, Collino, Calamita and Thiemermann.
This is an open-access article distributed under
the terms of the [Creative Commons Attribution
License \(CC BY\)](#). The use, distribution or
reproduction in other forums is permitted,
provided the original author(s) and the
copyright owner(s) are credited and that the
original publication in this journal is cited, in
accordance with accepted academic
practice. No use, distribution or reproduction
is permitted which does not comply with
these terms.

Correction: RG100204, a novel Aquaporin-9 inhibitor, reduces septic cardiomyopathy and multiple organ failure in murine sepsis

Shireen Mohammad^{1*}, Caroline E. O'Riordan¹, Chiara Verra¹,
Eleonora Aimaretti², Gustavo Ferreira Alves³, Klaus Dreisch⁴,
Johan Evenäs⁴, Patrizia Gena⁵, Angela Tesse⁶,
Michael Rützler^{7,8}, Massimo Collino³, Giuseppe Calamita⁵
and Christoph Thiemermann^{1*}

¹William Harvey Research Institute, Queen Mary University of London, London, United Kingdom,

²Department of Clinical and Biological Sciences, University of Turin, Turin, Italy, ³Department of Neurosciences "Rita Levi Montalcini", University of Turin, Turin, Italy, ⁴Red Glead Discovery Akiebolag (AB), Lund, Sweden, ⁵Department of Biosciences, Biotechnologies and Biopharmaceutics, University of Bari "Aldo Moro", Bari, Italy, ⁶Nantes Université, Institut National de la Santé et de la Recherche Médicale (INSERM), Centre National de la Recherche Scientifique (CNRS), l'institut du Thorax, Nantes, France, ⁷Department of Biochemistry and Structural Biology, Lund University, Lund, Sweden, ⁸Apoglyx Akiebolag (AB), Lund, Sweden

KEYWORDS

aquaporin (AQP), sepsis, cecal ligation and puncture, inflammation, multiple organ failure

A Correction on

RG100204, a novel Aquaporin-9 inhibitor, reduces septic cardiomyopathy and multiple organ failure in murine sepsis

By Mohammad S, O'Riordan CE, Verra C, Aimaretti E, Alves GF, Dreisch K, Evenäs J, Gena P, Tesse A, Rützler M, Collino M, Calamita G and Thiemermann C (2022) *Front. Immunol.* 13:900906. doi: 10.3389/fimmu.2022.900906

There was an error in **Figure 13** as published. The representative images related to Panel A have been inadvertently duplicated in Panel E. The corrected **Figure 13** and its caption appear below.

The original article has been updated.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

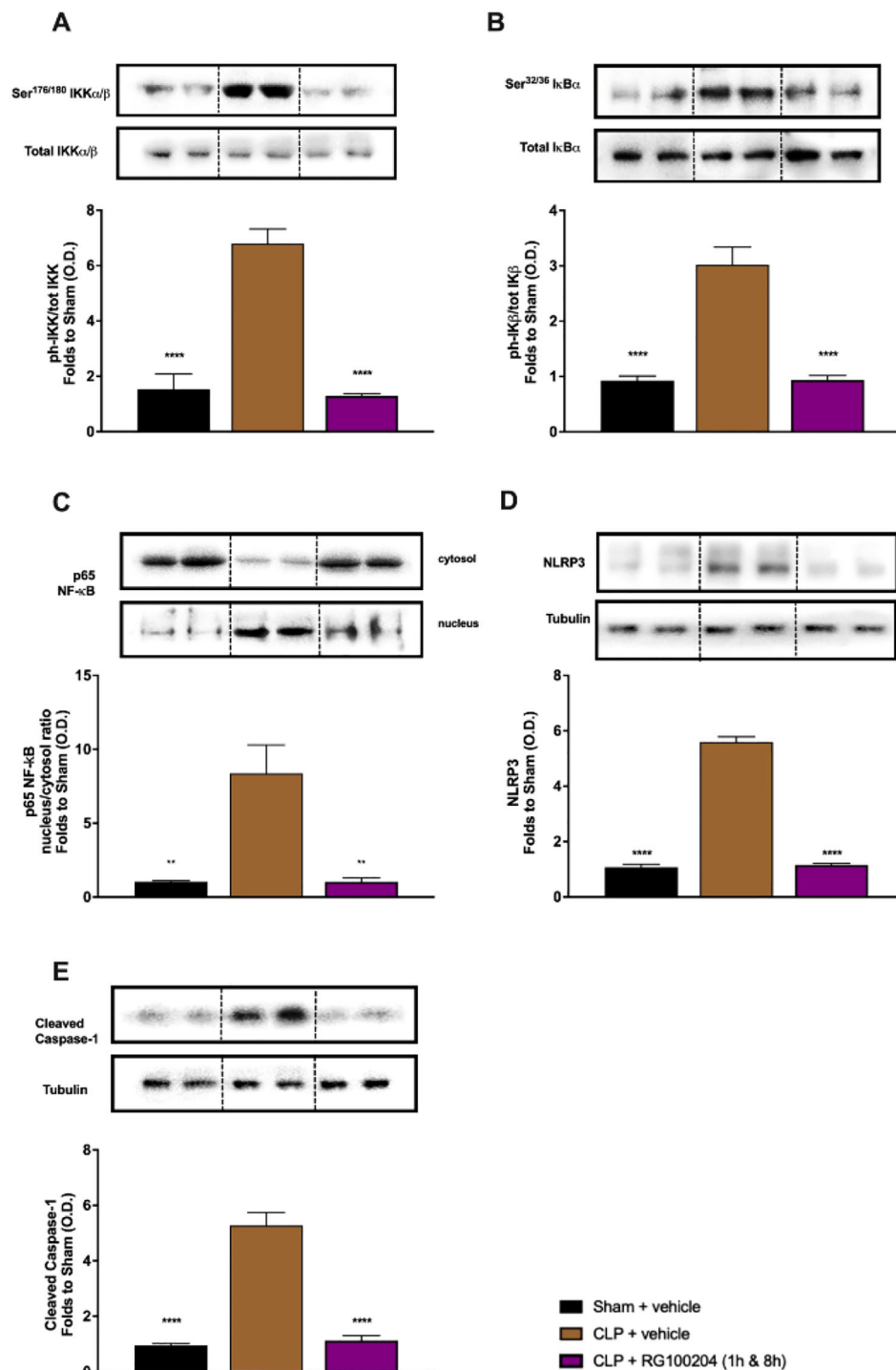


FIGURE 13

Effect of post-treatment (therapeutic administration) with RG100204 on the NF- κ B signalling pathway and the activation NLRP3 inflammasome in the heart. Heart samples were collected at the end of the experiment and the NF- κ B signalling pathway, as well as the activation of the NLRP3 inflammasome. Densitometry analysis of the bands is expressed as relative optical density (O.D.) of the (A) phosphorylation of IKK α/β at Ser178/180 corrected for the corresponding total IKK α/β content and normalized using the related sham band; (B) phosphorylation of I κ B α at Ser32/36 corrected for the corresponding total I κ B α content and normalized using the related sham band; (C) NF- κ B p65 subunit levels in both, cytosolic and nuclear fractions expressed as a nucleus/cytosol ratio normalized using the related sham bands; (D) NLRP3 activation, corrected against tubulin and normalized using the related sham bands; and (E) proteolytic cleavage of pro-caspase-1 to activated caspase-1 and normalized using the related sham band. The following groups were studied: sham + vehicle (n = 5), CLP + vehicle (n = 10), CLP +RG100204 (1 h & 8 h) (n = 10). All data were analyzed by one-way ANOVA, followed by a Bonferroni's post-hoc test. Data are expressed as mean \pm SEM. **P < 0.01 and ****P < 0.0001 vs. the respective sham-operated group.