Health Research Authority

East of England - Essex Research Ethics Committee
The Old Chapel
Royal Standard Place
Nottingham
NG1 6FS

22 June 2018

Emma Skilton
Clinical Trial Manager
Warwick Clinical Trials Unit
Warwick Medical School
University of Warwick
Coventry
CV4 7AL

Dear Ms Skilton,

Study title:	**STRESS-L: STudy into the REversal of Septic Shock with Landiolol (Beta Blockade)**
REC reference:	**17/EE/0368**
Protocol number:	**STRESS-L**
EudraCT number:	**2017-001785-14**
Amendment number:	**4**
Amendment date:	**20 June 2018**
IRAS project ID:	**213669**

Thank you for submitting the above amendment, which was received on 21 June 2018.

Research site	*Principal Investigator / Local Collaborator*
King's College Hospitals NHS Foundation Trust	Phil Hopkins
Royal Cornwall Hospitals NHS Trust	Michael Spivey
University Hospitals Plymouth NHS Trust	Sam Waddy
Dorset Country Hospital NHS Foundation Trust	Ian Mew
The Dudley Group NHS Foundation Trust	Vikram Anumakonda
Royal Free London NHS Foundation Trust	Nasirul Ekbal
University Hospitals Coventry and Warwickshire NHS Trust	Thomas Billyard

The amendment relates solely to the addition of new site(s) and/or investigator(s) within the National Health Service (NHS) or Health and Social Care (HSC) in Northern Ireland. Site-specific assessment (SSA) for any site within the National Health Service (NHS) or Health and Social Care (HSC) in Northern Ireland will form part of the nation specific local processes for that site. Guidance on how to work with sites is provided in the IRAS help section at https://www.myresearchproject.org.uk/help/hlpnhshscr.aspx

On behalf of the Committee, I am pleased to confirm the extension of the favourable opinion to the new site(s) and/or investigator(s), subject to management permission being given by the relevant host organisation prior to the study starting at the site.

Statement of compliance

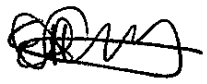
This Committee is recognised by the United Kingdom Ethics Committee Authority under the Medicines for Human Use (Clinical Trials) Regulations 2004, and is authorised to carry out the ethical review of clinical trials of investigational medicinal products.

The Committee is fully compliant with the Regulations as they relate to ethics committees and the conditions and principles of good clinical practice.

The Committee is constituted in accordance with the Governance Arrangements for Research Ethics Committees and complies fully with the Standard Operating Procedures for Research Ethics Committees in the UK.

17/EE/0368	**Please quote this number on all correspondence**

Yours sincerely



Ellena Stansbury
REC Assistant

Email: NRESCommittee.EastofEngland-Essex@nhs.net

Copy to: *Tony Whitehouse, Queen Elizabeth Hospital Birmingham NHS Foundation Trust*
Miss Nafisa Boota