Health Research Authority

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

07 December 2018

Miss Nafisa Boota
Warwick Clinical Trials Unit
University of Warwick, Gibbet Hill Campus
Coventry
CV4 7AL

Dear Miss Boota

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:	**8**
Amendment date:	**06 December 2018**
IRAS project ID:	**213669**

Thank you for submitting the above amendment, which was received on 07 December 2018.

Research site	*Principal Investigator / Local Collaborator*
Royal Victoria Infirmary, The Newcastle upon Tyne Hospitals NHS Foundation Trust, Queen Victoria Rd, NE1 4LP	Ian Clement
Freeman Hospital, Freeman Rd, High Heaton, Newcastle upon Tyne NE7 7DN	Ian Clement
The York Hospital, Wigginton Road, York, North Yorkshire, YO31 8HE	Joseph Carter
Amersham Hospital, Whielden Street, Amersham, Buckinghamshire, HP7 0JD	Raha West
Rotherham Hospital, Moorgate Road, Rotherham, S60 2UD	Anil Hormis
Addenbrooke's Hospital, Hills Road, Cambridge, Cambridgeshire, CB2 0QQ	James Varley
Leeds Teaching Hospitals NHS Trust, St. James's University Hospital, Beckett Street , Leeds, LS9 7TF	Andy Breen

The amendment relates solely to the addition of new sites and investigators 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 sites and investigators, subject to management permission being given by the relevant host organisation prior to the study starting at the site.

Statement of compliance

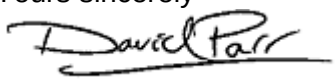
This Committee is recognised by the United Kingdom Ethics Committee Authority under the Medicines for Human Use (Clinical Trials) Regulations 2004, and is authorised to carry out the ethical review of clinical trials of investigational medicinal products.

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



David Parr

Email: NRESCommittee.EastofEngland-Essex@nhs.net

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