

## East of England - Essex Research Ethics Committee

The Old Chapel  
Royal Standard Place  
Nottingham  
NG1 6FS

30 April 2019

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

Dear Miss Boota

<b>Study title:</b>	<b>STRESS-L: STudy into the REversal of Septic Shock with Landiolol (Beta Blockade)</b>
<b>REC reference:</b>	<b>17/EE/0368</b>
<b>Protocol number:</b>	<b>STRESS-L</b>
<b>EudraCT number:</b>	<b>2017-001785-14</b>
<b>Amendment number:</b>	<b>9</b>
<b>Amendment date:</b>	<b>18 April 2019</b>
<b>IRAS project ID:</b>	<b>213669</b>

Thank you for submitting the above amendment, which was received on 18 April 2019.

<i>Research site</i>	<i>Principal Investigator / Local Collaborator</i>
Northampton General Hospital NHS Trust, Cliftonville, Northampton NN1 5BD	Jonny Wilkinson
Brighton and Sussex University Hospitals NHS Trust, Eastern Rd, Brighton, BN2 5BE	Barbara Phillips
East and North Hertfordshire NHS Trust, Lister Hospital, Stevenage, SG1 4AB	Sunil Jamadarkhana

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

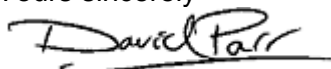
This Committee is recognised by the United Kingdom Ethics Committee Authority under the Medicines for Human Use (Clinical Trials) Regulations 2004 and is authorised to carry out the ethical review of clinical trials of investigational medicinal products.

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: [NRESCCommittee.EastofEngland-Essex@nhs.net](mailto:NRESCCommittee.EastofEngland-Essex@nhs.net)

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