



# Health Research Authority

## London - Surrey Borders Research Ethics Committee

Research Ethics Committee (REC) London Centre  
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30 March 2020

Dr Farah Al-Beidh  
ICNARC  
Napier House  
24 High Holborn  
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Dear Dr Al-Beidh

**Study title:** Randomized, Embedded, Multifactorial, Adaptive Platform  
trial for Community-Acquired Pneumonia  
**REC reference:** 18/LO/0660  
**EudraCT number:** 2015-002340-14  
**Amendment number:** 1.0  
**Amendment date:** 25 March 2020  
**IRAS project ID:** 237150

Thank you for submitting the above amendment, which was received on 25 March 2020.

<i>Research site</i>	<i>Principal Investigator / Local Collaborator</i>
Guys and St Thomas's	Dr Manu Shankar-Hari
South Tyneside Hospital,	Dr Christain Frey
Royal United Bath Hospital	Dr Ian Kerslate
City Hospital, Birmingham	Dr Jonathan Hulme
Southend Hospital	Dr Mark Saville
Royal Brompton Hospital	Dr Darius Armstrong-James
Royal Marsden Hospital	Dr Kate Tatham
Northwick Park Hospital,	Dr Ashley Whittington
Central Middlesex Hospital	Dr Ashely Whittington
Ealing Hospital	Dr Ashley Whittington
Wexham Park Hospital	Dr Omar Touma
Royal Gwent Hospital	Dr Tamas Szakmany
Kings Mill Hospital	Dr Paul Pulak
Southampton Hospital	Dr A Dushianthan
Salford Royal Hospital	Dr Paul Dark
Royal Liverpool Hospital	Dr Ingeborg Welters
Wythenshawe Hospital	Dr Peter Alexander

The Cumberland Infirmary	Dr Tim Smith
West Cumberland Hospital	Dr Tim Smith
Frimley Park Hospital	Dr George Evetts
Bolton Hospital	Dr Madhu Balasubramaniam
Belfast City Hospital, Belfast	Dr Johnathan Silversides
Royal Victoria Hospital, Belfast	Dr Jonathan Silversides
Mater Hospital, Belfast	Dr Jonathan Silversides

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.

**18/LO/0660**

**Please quote this number on all correspondence**

Yours sincerely



**Barbara Cuddon**  
**Approvals Specialist**

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*Copy to: Prof Anthony Gordon, Imperial College London*