Health Research Authority

NRES Committee London - South East
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23 July 2014

Professor Philip M. W. Bath
Professor of Stroke Medicine
University of Nottingham
Division of Stroke Medicine - Research Dept
Clinical Sciences Building
City Hospital Campus
Hucknall Road
Nottingham NG5 1PB

Dear Professor Bath

Study title:	**Safety and tolerability of clopidogrel when added to aspirin and dipyridamole in high risk patients with recent ischaemic stroke: a randomised controlled trial**
REC reference:	**08/H1102/112**
Protocol number:	**31350**
EudraCT number:	**2007-006749-42**
Amendment number:	**Change of PI, New Sites dated 23 July 2014**
Amendment date:	**23 July 2014**

Thank you for submitting the above amendment, which was received on 23 July 2014.

Research site	*Principal Investigator / Local Collaborator*
St. Thomas' Hospital, Guy's and St. Thomas' NHS Foundation Trust	Dr Ajay Bhalla
Royal Preston Hospital, Lancashire Teaching Hospitals NHS Foundation Trust	Dr Hedley Emsley
University Hospital of North Durham, County Durham and Darlington NHS Foundation Trust	Dr David Bruce
York Hospital, York Teaching Hospitals NHS Foundation Trust	Dr Elizabeth Iveson

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. The site-specific assessment for the site(s) will therefore form part of the research governance review. The Site-Specific Information (SSI) Form for the site should be included with the application for R&D approval.

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 NHS/HSC R&D office(s) prior to the study starting at the site.

Statement of compliance

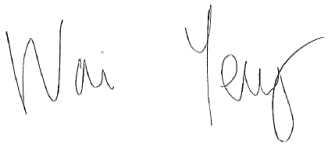
This Committee is recognised by the United Kingdom Ethics Committee Authority under the Medicines for Human Use (Clinical Trials) Regulations 2004, and is authorised to carry out the ethical review of clinical trials of investigational medicinal products.

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.

08/H1102/112	**Please quote this number on all correspondence**

Yours sincerely



Mr Wai Yeung
Research Ethics Committee (REC) Assistant

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