



# Health Research Authority

## NRES Committee London - South East

Bristol Research Ethics Committee Centre  
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17 April 2014

Mr Paul Cartledge  
Research Innovation Services, University of Nottingham  
Kings Meadow Campus, Lenton Lane  
Nottingham  
NG7 2NR

Dear Mr Cartledge

**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:** AM29: SA02/14

**Amendment date:** 11 March 2014

Thank you for submitting the above amendment, which was received on 31 March 2014.

| Research Site  | Principal Investigator / Local Collaborator |
|--|---|
| Great Western Hospitals NHS Trust, Great Western Hospital, Marlborough Road, Swindon, Wiltshire SN3 6BB  | Dr Dr Gopinath Ramadurai                    |
| Bedford Hospital NHS Trust, Kempston Road, Bedford, MK42 9DL   | Dr Mohammad Wasil                           |
| Salford Royal NHS Foundation Trust, North West Hyperacute Stroke Research Centre, E Room b210, Clinical Science Building, Stott Lane, Salford M6 8HD | Dr Chris Douglass                           |
| Heart of England NHS Foundation Trust, Good Hope Hospital, rectory Road, Sutton Coldfield, Birmingham B91 2JL  | Dr Arumugam Ravindrane                      |
| Heart of England NHS Foundation Trust, Solihull  | Dr Khaled Elfandi                           |

|   |                      |
|---|----------------------|
| Hospital, Lode Lane, Birmingham B91 2JL   |                      |
| United Lincolnshire Hospitals NHS Trust, Greetwell Road, Lincoln, Lincolnshire LN2 5QY  | Dr Jagdish Sharma    |
| North Tees & Hartepool NHS Foundation Trust, Stroke Unit, University Hospital of North Tees, Hardwick Road, Stockton on Tees TS19 8PE | Dr Balakrishma Kumar |

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.

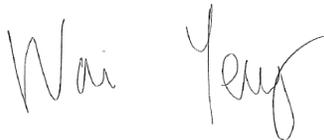
The Committee is fully compliant with the Regulations as they relate to ethics committees and the conditions and principles of good clinical practice.

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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Copy to: *Professor Philip M. W. Bath, University of Nottingham*

