### Signature and Delegation list

**Responsibility Key:**

1. Register patients in SITS Registry/SITS Open CRF
2. Assess inclusion, exclusion criteria and confirm eligibility
3. Make registration call
4. Obtain informed consent
5. Perform NIHSS assessment
6. Make study related medical decisions
7. Sign the Physician’s Verification Form
8. Perform mRS assessment
9. Obtain study measurements/collection data according to protocol
10. Source documentation entry (eg. Medical Notes)
11. Assess adverse events
12. Assess adverse events and Serious Adverse Events
13. Source documentation entry (eg. Medical Notes)
14. Maintain study files
15. Archive study material

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<tr>
<th>FULL NAME</th>
<th>EMAIL ADDRESS</th>
<th>PHONE NUMBER</th>
<th>POSITION IN THE STUDY</th>
<th>RESPONSIBILITIES</th>
<th>DATE OF RESPONSIBILITIES</th>
<th>INITIALS</th>
<th>SIGNATURE DELEGATES</th>
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To be signed and dated at end of study:

Principal Investigator: ___________________ Date: ______________

2016-02-18, version 2.0