

RECAST-3

SITE INVESTIGATOR MEETING

Remote Ischaemic Conditioning After Stroke 3 (RECAST-3)

A multicentre randomised controlled trial

16/05/2024

AGENDA

1. Recap of study flow
2. Site and recruitment update
3. Recent FAQs
4. Other important updates
 - i. Device training videos
 - ii. Ward staff training
 - iii. Amendment in progress
 - iv. Screening logs
 - v. Sub-study
5. Questions

RECAST 3

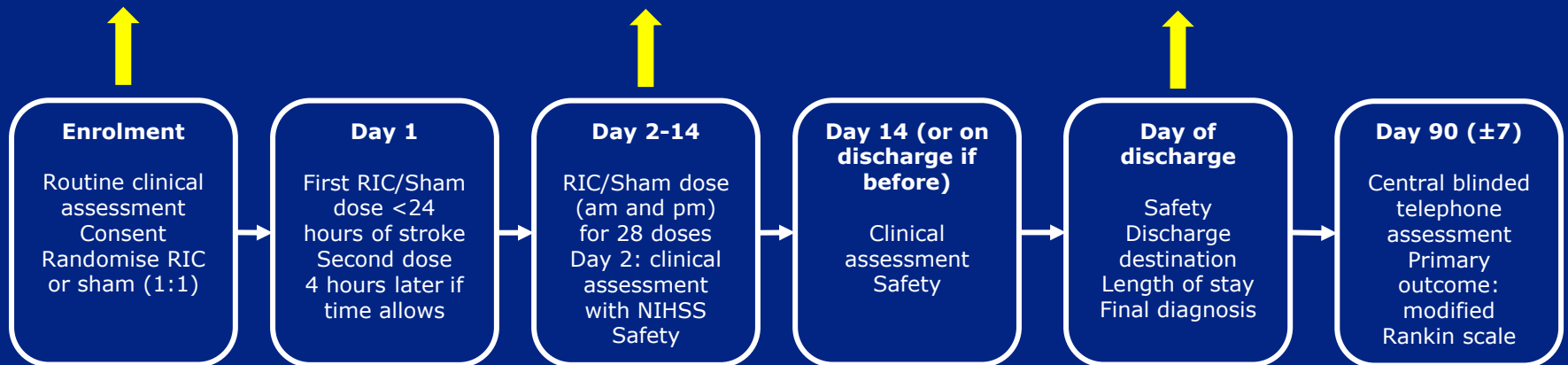
REMOTE ISCHAEMIC CONDITIONING AFTER STROKE TRIAL 3

STUDY FLOW

Randomisation form

Day 2 follow-up form

Death/discharge form



Day 1 follow-up form

End of treatment form

Day 90 telephone f/up by Nottingham

SCANS:

- Baseline CT scan prior to enrolment

RECAST 3

REMOTE ISCHAEMIC CONDITIONING AFTER STROKE TRIAL 3

SITE & RECRUITMENT UPDATE

Sites open (n=5)
Participants (n=3)

Open sites:

	Site	Participants
1	Royal Derby Hospital	2
2	Queen's Medical Centre, Nottingham	1
3	Fairfield General Hospital, Bury	0
4	Aberdeen Royal Infirmary	0
5	UCLH	0

*Twenty-one other sites are reviewing feasibility or have previously expressed an interest to take part but haven't yet started set up

Sites in set up having received the LIP

1	Morrison Hospital, Swansea
2	Bronglais General Hospital, Aberystwyth
3	St. George's Hospital, London
4	King's College Hospital, London
5	Princess Royal University Hospital, London
6	Salford Royal Hospital
7	James Cook University Hospital, Middlesbrough
8	Leighton Hospital, Crewe
9	Charing Cross, London
10	Royal Hallamshire Hospital, Sheffield
11	Bradford Royal Infirmary
12	Southampton General Hospital
13	Royal Stoke University Hospital
14	Leicester Royal Infirmary
15	Sunderland Royal Hospital

Recruitment target: 1-2 per site per month

RECENT FAQS

- ▲ **SAE reporting period:** Sites should report all SAEs for 20 days. Thereafter, please report all fatal SAEs and safety outcomes through until day 90.
- ▲ **Accepted adjustments to the intervention to encourage continued treatment:** Acceptable to change to unilateral and/or once daily treatments if necessary. Document on a filenote and ensure treatment log CRF completed in full. This will not be a protocol violation.
- ▲ **24-hour inclusion window:** Taken from symptom discovery/wake up time.
- ▲ **Previous breast cancer or axillary node clearance:** Not an exclusion criteria - down to the discretion of the site investigator.
- ▲ **No radiological evidence of stroke on baseline scan but are treated in accordance with a clinical diagnosis of stroke:** Considered eligible for the trial if all other criteria are met.

OTHER IMPORTANT UPDATES

- ▲ Device training videos available on the trial website
- ▲ Ward staff training
- ▲ Associate PI scheme
- ▲ Amendment in progress to gain approval for Research Practitioners/Coordinators/Assistants to obtain consent where local policy allows this
- ▲ Screening logs – please complete and send over to us monthly.
- ▲ Sub-study

End slide

Questions?