

### 104 patients now recruited!

#### The STITCH (Trauma) trial now has over 100 patients!

**Congratulations to the MM Institute of Medical Sciences & Research, Haryana who recruited the 100th patient on 14th September 2011.**

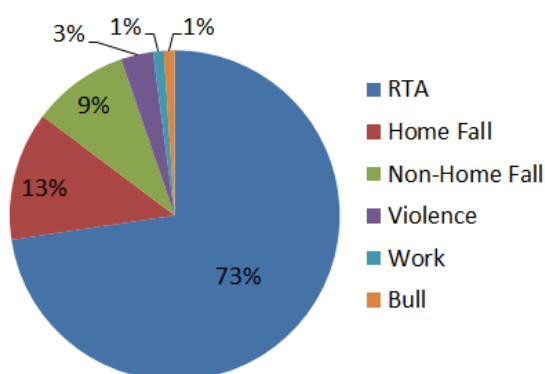
**Many thanks also to Neuro Hospital, Biratnagar, Beijing Tiantan Hospital, CSM Medical University, Lucknow and Klinikum Bogenhausen, Munich who have all recruited since, taking our total recruitment to 104 patients.**

**We now have a total of 45 registered centres participating in the trial.**

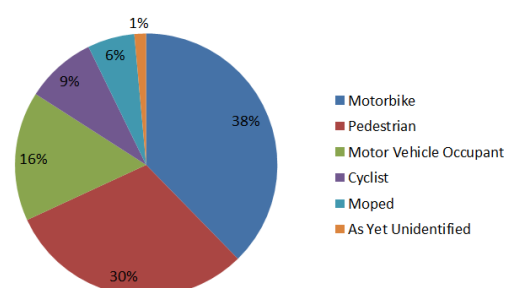
| Centre              | Total | Change since last newsletter |
|---------------------|-------|------------------------------|
| Lucknow             | 21    | 6                            |
| Beijing             | 20    | 6                            |
| Dehradun            | 12    | 0                            |
| Biratnagar          | 12    | 2                            |
| Visakhapatnam       | 7     | 0                            |
| Ludhiana            | 3     | 0                            |
| Newcastle           | 3     | 0                            |
| Tianjin             | 3     | 0                            |
| Wardha              | 3     | 0                            |
| Dundee              | 2     | 1                            |
| Kaunas              | 2     | 1                            |
| Mysore              | 2     | 0                            |
| Peshawar            | 2     | 1                            |
| Shanghai            | 2     | 0                            |
| Timisoara           | 2     | 2                            |
| Bangalore           | 1     | 0                            |
| Brno                | 1     | 0                            |
| Chennai             | 1     | 0                            |
| Haryana             | 1     | 1                            |
| Heidelberg          | 1     | 0                            |
| Munich              | 1     | 1                            |
| Southampton         | 1     | 0                            |
| Valladolid          | 1     | 1                            |
| Calcutta            | 0     | 0                            |
| Cluj                | 0     | 0                            |
| Haywards Heath      | 0     | 0                            |
| Hyderabad           | 0     | 0                            |
| Jena                | 0     | 0                            |
| Johor Bahru         | 0     | 0                            |
| Kerala              | 0     | 0                            |
| Klaipeda            | 0     | 0                            |
| Kubang Kerian       | 0     | 0                            |
| Leeds               | 0     | 0                            |
| London (St Georges) | 0     | 0                            |
| New York            | 0     | 0                            |
| Pennsylvania        | 0     | 0                            |
| Portland            | 0     | 0                            |
| Rome                | 0     | 0                            |
| Santander           | 0     | 0                            |
| Szeged              | 0     | 0                            |
| Toronto             | 0     | 0                            |
| Ulm                 | 0     | 0                            |
| Ulm—Günzburg        | 0     | 0                            |
| Yerevan             | 0     | 0                            |
| Zagazig             | 0     | 0                            |

Centres who have joined the study since the last newsletter are highlighted in green.

#### Injury Type: Data From 95 Patients



#### RTA Type: Data From 69 RTA Patients



#### Inclusion Criteria

- Adults aged 14 or over.
- Evidence of 1 or 2 TICHs or contusions with a confluent volume of attenuation significantly raised above that of the background white and grey matter that has a total volume greater than 10mls (calculated by width x height x length / 2 in cm).
- Within 48 hours of head injury.
- Clinical equipoise: only patients for whom the responsible neurosurgeon is uncertain about the benefits of either treatment are eligible.

#### Exclusion Criteria

- A significant surface haematoma (EDH or SDH) requiring surgery.
- Three or more separate haematomas fulfilling inclusion criteria.
- If surgery cannot be performed within 12 hours of randomisation.
- If the haemorrhage/ contusion is located in the cerebellum.
- Severe pre-existing physical or mental disability or severe co-morbidity which would lead to a poor outcome even if the patient made a full recovery from the head injury.
- Permanent residence outside a study country preventing follow up.
- Patient and/or relative has a strong preference for one treatment modality.

#### Is your centre's recruitment on target?

**We have an interim target of 110 patients including 12 UK patients by November 2011.**

**It is important that we reach this target and that ALL centres aim to find between 1 & 3 suitable patient a year.**

**So far in 2011, 15 centres have recruited at least 1 patient and 3 centres have recruited more than 10 patients. Check the recruitment leader board above to see how your centre is doing.**

