

Dutch Intracerebral Hemorrhage Surgery Trial

NCT05460793

Status	RECRUITING
Phase	Not Applicable
Sponsor	Radboud University Medical Center
Enrollment	600 participants

Key Eligibility Criteria

Inclusion (6)

- Age 18 years or older;
- NIHSS e 2;
- Supratentorial non-traumatic ICH confirmed by non-contrast CT, without a CT-angiography confirmed causative vascular lesion (e.g. aneurysm, arteriovenous malformation \[AVM\], dural arteriovenous fistula \[DAVF\], cerebral venous sinus thrombosis \[CVST\]), or other known underlying lesion (e.g. tumor, cavernoma);
- Minimal hematoma volume of 10 mL;
- Intervention can be started within 8 hours of symptom onset;

... and 1 more (see full listing online)

Exclusion (7)

- Considerable pre-stroke dependency in activities of daily living, defined as a pre-stroke mRS e3;
- ICH-GS score e11;
- Hemorrhage due to hemorrhagic transformation of an infarct;
- Untreated coagulation abnormalities, including INR $\gt;1.3$ (point of care measurement allowed), treatment with heparin and treatment with factor Xa inhibitors. Patients on vitamin K antagonist can be included after correction of the INR, and patients on dabigatran (direct thrombin inhibitor) can be included after reversal of dabigatran with idarucizumab;
- Moribund (e.g. coning, bilateral dilated unresponsive pupils), or progressively deteriorating clinical course with imminent death;

... and 2 more (see full listing online)

Locations (11 total)

Amsterdam University Medical Center, Amsterdam, Netherlands

Medisch Spectrum Twente, Enschede, Netherlands

University Medical Center Groningen, Groningen, Netherlands

... and 8 more locations

<https://clinicaltrials.gov/study/NCT05460793>

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