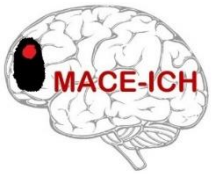

ISRCTN 15383301
IRAS 1004870

MAnnitol for Cerebral oEdema after IntraCerebral Haemorrhage (MACE-ICH): a feasibility trial

Eligibility Checklist V1.0

	Inclusion Criteria *(All must be 'YES' for the patient to take part)*	**Yes**	**No**
1	Adult (18 years and over)		
2	Spontaneous ICH confirmed by CT scan with estimated largest diameter > 2 cm		
3	≤72 hours of ictus (or from last seen healthy)		
4	Cerebral oedema with or without evidence of mass effect		
5	At risk of developing oedema (limited GCS <9 (eye opening and motor only) and NIHSS>8)		
6	Signed consent (patient, personal or professional representative or independent physician)		

	Exclusion Criteria *(All must be 'NO' for the patient to take part)*	**Yes**	**No**
1	GCS<5		
2	Premorbid mRS >3		
3	Isolated subarachnoid haemorrhage		
4	Haemorrhage known to be from: trauma or venous thrombosis or arteriovenous malformation or brain tumour or transformation of cerebral infarct or cerebral aneurysm or thrombolytic drug		
5	Known hypersensitivity to mannitol		
6	Severe renal failure (e-GFR<30ml/min or dialysis)		
7	Severe pulmonary oedema/ cardiac failure		
8	Hypotension at baseline (SBP <90 mm Hg)		
9	Anuria		
10	Patient unwilling to participate		
11	Geographical or other factors which prohibit follow-up		
12	Pre-existing comorbidity with pre-ictal life expectancy <6 months		
13	Severe dementia		
14	Planned for palliative care		
15	Severe hypernatremia (sodium >160 mmol)		
16	Severe hyponatremia (sodium <125 mmol)		
17	Women of child-bearing potential with a positive pregnancy test at the time of admission, or lactating		
18	Patients in whom peripheral intravenous cannula cannot be placed		
19	Planned neurosurgery		

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Eligibility must be signed off by a Medic

*(The medic does **not** have to be on the delegation log)*

(Name)	(Signature)
(Occupation)	(Date)