

<b>PATIENT DEMOGRAPHICS</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
Age at time of ICH		100%
Gender		100%
<b>COMORBIDITIES PRESENT BEFORE ICH</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
Pre-morbid Rankin	Modified Rankin Scale score before the index ICH.	99%
Angina pectoris within 30 days before admission	Angina pectoris as defined by ICD-10 (International Statistical Classification of Diseases and Related Health Problems) codes that begin with I20 occurring within 30 days before admission.	99%
Peripheral vascular disease within 30 days before admission	Peripheral vascular diseases as defined by ICD-10 codes that begin with I73 occurring within 30 days before admission.	99%
Deep venous thrombosis within 30 days before admission	Deep venous thrombosis as defined by ICD-10 codes that begin with I80 or with I82 occurring within 30 days before admission.	99%
Myocardial infarction within 30 days before admission	Myocardial infarction as defined by ICD-10 codes that begin with I21 occurring within 30 days before admission.	99%
Cerebral infarction 30 days before admission	Cerebral infarction as defined by ICD-10 codes that begin with I63 occurring within 30 days before admission.	99%
Any history of a malignant neoplasm	Any forms of a malignant neoplasm as defined by any ICD-10 codes that begin with C00 through to C43 and C45 through to C97 diagnosed at any time before admission.	99%
Malignant neoplasm within 2 years before admission	Any forms of a malignant neoplasm as defined by any ICD-10 codes that begin with C00 through to C43 and C45 through to C97 diagnosed within 2 years before admission.	99%
Any history of ischaemic events	Any history prior to admission for the index ICH of a vaso-occlusive/ischemic event (e.g. myocardial infarction, angina pectoris, TIA, cerebral infarction, peripheral vascular disease, pulmonary embolism) .	99%
Any history of deep venous thrombosis	Any history prior to admission for the index ICH of deep venous thrombosis.	99%
Any history of heart failure	Any history prior to admission for the index ICH of heart failure.	99%
Any history of ischaemic stroke	Any history prior to admission for the index ICH of ischaemic stroke.	99%
Any history of pulmonary embolism	Any history prior to admission for the index ICH of pulmonary embolism.	99%
Any history of peripheral vascular disease	Any history prior to admission for the index ICH of peripheral vascular disease.	99%
Any history of	Any history prior to admission for the index ICH of transient	99%

transient ischaemic attacks	ischaemic attacks.	
<b>MEDICATION AT TIME OF ICH</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
On antihypertensives at time of ICH	Includes any medication commonly prescribed as an antihypertensive (not furosemide, since this is usually prescribed for heart failure.)	99%
On anticoagulants at time of ICH	Includes warfarin, enoxaparin (both long term, short term – prophylactic and therapeutic doses if given immediately prior to the ICH.)	100%
On antiplatelets at time of ICH	Includes aspirin, dipyridamol and clopidogrel.	99%
<b>FIRST ASSESSMENT AFTER ICH</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
Present in the hospital	Patient presented to the hospital for their ICH.	100%
Nature of symptom onset	The nature of symptom onset was divided into three categories: <ul style="list-style-type: none"> <li>▪ Awake at onset</li> <li>▪ Awoke from sleep</li> <li>▪ Last seen well</li> </ul>	87%
Delay in hours from onset to CT scan (rounded up)	<ul style="list-style-type: none"> <li>▪ Delay in hours from the nature of onset until the CT scan that diagnosed the ICH was performed.</li> <li>▪ Completeness varies depending on the search mode: for “Equal or less” and “Less than” the completeness is 100%. However over-estimations have been applied in 32%.</li> <li>▪ For the other search modes the completeness is 68%.</li> <li>▪ Also note that the nature of onset determines the true delay from ICH symptom onset to when a CT scan was made.</li> </ul>	68 or 100%
Systolic BP on admission	Systolic blood pressure measured at the first assessment in hospital after ICH.	88%
Diastolic BP on admission	Diastolic blood pressure measured at the first assessment in hospital after ICH.	87%
GCS Eye score on admission	Glasgow Coma Scale Eye score at the first assessment in hospital after ICH.	96%
GCS Verbal score on admission	Glasgow Coma Scale Verbal score at the first assessment in hospital after ICH.	97%
GCS Motor score on admission	Glasgow Coma Scale Motro score at the first assessment in hospital after ICH.	96%
GCS Total score on admission	Glasgow Coma Scale Total score at the first assessment in hospital after ICH.	96%
Total prognostic ICH score	Total prognostic ICH score derived from the first assessments in hospital after ICH.	95%
<b>RADIOGRAPHIC CHARACTERISTICS</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
First-ever ICH	The intracerebral haemorrhage that made the patient part of this study was the first one in the patient’s lifetime.	100%
Total ICH volume (ml)	Intracerebral haemorrhage volume (excluding subarachnoid or intraventricular blood) was measured with the ABC/2 method.	98%
Number of bleeds	Single intracerebral haemorrhage versus multiple intracerebral haemorrhages.	100%

<p>ICH cause (primary /secondary)</p>	<p>Primary causes include:</p> <ul style="list-style-type: none"> <li>▪ Acquired small vessel disease</li> <li>▪ Cerebral amyloid angiopathy – without a detected genetic mutation;</li> <li>▪ Cerebral amyloid angiopathy – with a detected genetic mutation</li> <li>▪ Genetic small artery diseases – CADASIL</li> <li>▪ Genetic small artery diseases – COL4A1 mutation</li> <li>▪ Genetic small artery diseases – familial without an identified mutation</li> <li>▪ Unknown cause</li> </ul> <p>Secondary causes include:</p> <ul style="list-style-type: none"> <li>▪ Moya-moya phenomenon</li> <li>▪ Vasculitis, reversible cerebral vasoconstriction syndrome</li> <li>▪ Arterial aneurysm</li> <li>▪ Arteriovenous malformation</li> <li>▪ Dural arteriovenous fistula</li> <li>▪ Cavernous malformation</li> <li>▪ Acute leucoencephalopathy syndromes</li> <li>▪ Intracranial venous thrombosis</li> <li>▪ Malignancy</li> </ul>	<p>100%</p>
<p>Specific ICH cause</p>	<p>The following specific ICH causes were defined:</p> <ul style="list-style-type: none"> <li>▪ Acquired small vessel disease</li> <li>▪ Cerebral amyloid angiopathy – without a detected genetic mutation;</li> <li>▪ Cerebral amyloid angiopathy – with a detected genetic mutation</li> <li>▪ Genetic small artery diseases – CADASIL</li> <li>▪ Genetic small artery diseases – COL4A1 mutation</li> <li>▪ Genetic small artery diseases – familial without an identified mutation</li> <li>▪ Moya-moya phenomenon</li> <li>▪ Vasculitis, reversible cerebral vasoconstriction syndrome</li> <li>▪ Arterial aneurysm</li> <li>▪ Arteriovenous malformation</li> <li>▪ Dural arteriovenous fistula</li> <li>▪ Cavernous malformation</li> <li>▪ Acute leucoencephalopathy syndromes</li> <li>▪ Intracranial venous thrombosis</li> <li>▪ Malignancy Multiple – evidence of at least 2 of the previously mentioned causes</li> <li>▪ Unknown</li> </ul>	<p>100%</p>
<p>Strictly lobar ICH</p>	<p>Haemorrhage classified as lobar.</p>	<p>100%</p>
<p>Supratentorial ICH</p>	<p>ICH limited to the supratentorial compartment.</p>	<p>100%</p>
<p>Hydrocephalus</p>	<p>Dilated ventricles caused by an increase in volume of the cerebrospinal fluid.</p>	<p>98%</p>
<p>Intraventricular extension</p>	<p>Presence or absence of blood in the intraventricular spaces as determined from radiographic or pathological examination.</p>	<p>98%</p>

Subarachnoid extension	Presence or absence of blood in the subarachnoid spaces as determined from radiographic or pathological examination.	98%
Subdural extension	Presence or absence of blood in the subdural spaces as determined from radiographic or pathological examination.	98%
<b>LABORATORY RESULTS</b>		
<b>Variable name</b>	<b>Definition</b>	<b>Completeness</b>
Creatinine (mcmol/l) on admission	Serum creatinine determined on admission to the hospital.	96%
Glucose (mmol/l) on admission	Serum glucose determined on admission to the hospital.	78%
INR on admission	INR determined on admission to the hospital.	58%
Platelets (x 10 <sup>9</sup> /L) on admission	Platelet count determined on admission to the hospital.	95%