# Improving stroke diagnosis: A study using blood tests and clinical information to quickly identify stroke types in suspected stroke patients

<b>Submission date</b> 27/01/2025	<b>Recruitment status</b> Recruiting	[X] Prospectively registered		
2170172023		Protocol		
Registration date	Overall study status	Statistical analysis plan		
04/02/2025	Ongoing	☐ Results		
Last Edited	<b>Condition category</b> Circulatory System	☐ Individual participant data		
10/09/2025		[X] Record updated in last year		

#### Plain English summary of protocol

Background and study aims

Stroke is a leading cause of death and disability worldwide. Early and accurate diagnosis of stroke subtypes, such as bleeding in the brain (intracerebral haemorrhage, ICH) or a blocked blood vessel (large vessel occlusion, LVO), is critical to ensuring timely treatment and better outcomes. Current tools like the Face Arm Speech Test (FAST) are unable to reliably identify these subtypes, leading to delays in care. This study aims to develop and test a new diagnostic tool that combines clinical observations with a quick blood test (the LVOne test) to improve the diagnosis of stroke subtypes shortly after hospital arrival.

#### Who can participate?

Adults aged 18 or older who are brought to the hospital by ambulance with symptoms of a possible stroke that started within the last six hours are eligible to participate. Patients who have had recent injuries or conditions that could affect the test results may not be eligible.

#### What does the study involve?

Participants will have a small blood sample taken through a finger prick and, when possible, an additional sample from a routine blood draw. The blood samples will be tested using the LVOne test to measure levels of specific markers associated with stroke subtypes. Clinical data and brain imaging results will also be collected. The results of the blood test will not influence the patient's treatment but will be compared to final diagnoses to evaluate the tool's accuracy.

What are the possible benefits and risks of participating?

There are no direct benefits to participants, but the study may improve future stroke diagnosis and care. Risks are minimal, with a small chance of discomfort or bruising from the finger prick or additional blood samples.

Where is the study run from? University of Manchester (UK)

When is the study starting and how long is it expected to run for? May 2024 to December 2026

Who is funding the study?

The study is funded by the Translation Manchester Accelerator Awards and the University of Manchester (UK)

Who is the main contact?

Prof Adrian Parry-Jones, adrian.parry-jones@manchester.ac.uk

# Contact information

#### Type(s)

Public, Scientific, Principal investigator

#### Contact name

Prof Adrian Parry-Jones

#### **ORCID ID**

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#### Contact details

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# Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

351335

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

MR/X502868/1

# Study information

#### Scientific Title

DIAGnosis using NOvel technology for Subtypes In Stroke (DIAGNOSIS study): performance evaluation

#### Acronym

**DIAGNOSIS** 

#### **Study objectives**

Combining clinical features with point-of-care biomarker tests (GFAP and D-dimer) can improve the prehospital differentiation of stroke subtypes, specifically intracerebral hemorrhage (ICH) and large vessel occlusion (LVO), from other suspected stroke cases. This approach has the potential to enable more targeted and timely interventions, improving patient outcomes and resource allocation.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Approved 31/07/2025, Yorkshire & The Humber - Leeds West Research Ethics Committee (NHSBT Newcastle Blood Donor Centre, Holland Drive, Newcastle upon Tyne, NE2 4NQ, Newcastle upon Tyne, NE2 4NQ, United Kingdom; 0207 1048053, 0207 104 8272, 02071048100; leedswest.rec@hra.nhs.uk), ref: 25/YH/0102

#### Study design

Observational diagnostic accuracy study

#### Primary study design

Observational

#### Study type(s)

Diagnostic

## Health condition(s) or problem(s) studied

Differentiation of intracerebral hemorrhage (ICH) and large vessel occlusion (LVO) from other suspected stroke cases in adult patients presenting with stroke-like symptoms.

#### **Interventions**

This is an observational study with no interventions. The study methodology involves:

- 1. Collecting clinical features from suspected stroke patients upon hospital arrival, as documented in ambulance and emergency department records.
- 2. Blood sample collection:
- Fingerprick capillary blood sample for the LVOne test.
- Venous blood sample (< 4 ml in EDTA tube) for processing and storage.
- 3. Testing using UpFront Diagnostics' GFAP and D-dimer lateral flow tests (LVOne test), with results available within 15 minutes.
- 4. Processing of venous blood samples:
- Centrifugation and plasma aliquoting within 4 hours of collection
- Freezing of plasma at -80°C for later analysis
- 5. Comparing diagnostic results from the LVOne test and biomarker analyses with final diagnoses determined through clinical evaluation and brain imaging (CT and/or MRI).

The study will be conducted at three major stroke centres in Greater Manchester: Salford Royal Hospital, Fairfield General Hospital, and Stepping Hill Hospital. Participants will be followed up prior to hospital discharge to collect information on final diagnosis and treatment.

#### Intervention Type

Other

#### Primary outcome(s)

- 1. Diagnostic accuracy of models for determining intracerebral haemorrhage (ICH) is measured using point-of-care GFAP and D-dimer tests (LVOne test) at baseline and prior to hospital discharge
- 2. Diagnostic accuracy of models for determining large vessel occlusion (LVO) is measured using point-of-care GFAP and D-dimer tests (LVOne test) at baseline and prior to hospital discharge
- 3. Final diagnoses of intracerebral haemorrhage (ICH) are measured using CT/MRI prior to hospital discharge
- 4. Final diagnoses of large vessel occlusion (LVO) are measured using CT/MRI prior to hospital discharge

#### Key secondary outcome(s))

- 1. Sensitivity, specificity, positive predictive value, and negative predictive value for identifying intracerebral haemorrhage (ICH) and large vessel occlusion (LVO) are measured using the LVOne test (GFAP and D-dimer levels) compared with final diagnoses determined by brain imaging (CT /MRI) at baseline (on hospital arrival) and prior to hospital discharge
- 2. Diagnostic accuracy metrics of the developed models compared with those of the Face Arm Speech Test (FAST) and other standard diagnostic tools are measured using a retrospective review of ambulance and hospital patient records at baseline (on hospital arrival) to study completion
- 3. Evidence of the impact of diagnostic models on clinical decision-making, such as changes in treatment pathways or triage decisions, is measured using a retrospective analysis of clinical pathways and decisions documented in patient notes from baseline (on hospital arrival) to study completion

#### Completion date

15/12/2026

# **Eligibility**

#### Key inclusion criteria

- 1. The patient arrived at the study hospital via emergency ambulance.
- 2. The patient is 18 years of age or older.
- 3. Ambulance staff suspected a new acute stroke prior to arrival at the hospital.
- 4. Stroke symptoms began within 6 hours of sample collection (if the onset was not witnessed, the patient was last known to be well less than 6 hours ago).
- 5. Blood samples can be collected prior to the administration of any treatment.
- 6. Urgent brain imaging is planned as part of the patient's diagnostic pathway.

# Participant type(s)

Patient

#### Healthy volunteers allowed

No

#### Age group

Adult

#### Lower age limit

18 years

#### Sex

All

#### Key exclusion criteria

- 1. The patient was assessed at another hospital and transferred for ongoing care.
- 2. The patient had a recent diagnosis (within the last 4 weeks) of deep vein thrombosis, pulmonary embolism, arterial embolism, stroke, transient ischaemic attack, long bone fracture, major trauma, or surgery under general anaesthesia, as these conditions may increase D-dimer levels.
- 3. The patient had a recent head injury (within the last 4 weeks) requiring hospital care, as this may increase GFAP levels.
- 4. Stroke symptoms started more than 6 hours prior to sample collection.

#### Date of first enrolment

01/10/2025

#### Date of final enrolment

31/10/2026

## Locations

#### Countries of recruitment

United Kingdom

England

# Study participating centre Northern Care Alliance NHS Foundation Trust

Salford Royal Stott Lane Salford United Kingdom M6 8HD

Study participating centre
Stockport NHS Foundation Trust
Stepping Hill Hospital
Poplar Grove

Stockport United Kingdom SK2 7JE

# Study participating centre Fairfield General Hospital

Fairfield General Hospital Rochdale Old Road Bury United Kingdom BL9 7TD

# Sponsor information

#### Organisation

University of Manchester

#### **ROR**

https://ror.org/027m9bs27

# Funder(s)

#### Funder type

Other

#### **Funder Name**

Translation Manchester Accelerator Awards, Confidence for Translation (C4T) 2024 scheme

#### **Funder Name**

University of Manchester

#### Alternative Name(s)

The University of Manchester, University of Manchester UK, University of Manchester in United Kingdom, UoM

#### Funding Body Type

Government organisation

#### **Funding Body Subtype**

Universities (academic only)

#### Location

United Kingdom

# **Results and Publications**

#### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be available upon reasonable request from Prof. Adrian Parry-Jones (adrian.parry-jones@manchester.ac.uk).

#### Details of the data sharing plan:

Type of data to be shared: Anonymised clinical data and biomarker test results collected during the study, including diagnostic model outputs and final diagnosis imaging results.

Data availability: Data will become available after the study results are published and will remain accessible for a minimum of 5 years.

Access criteria: Data will be shared with researchers affiliated with academic or healthcare institutions for the purpose of conducting ethically approved research into stroke diagnosis and management. A data-sharing agreement will be required.

Mechanism: Researchers can request access to the data by contacting the primary investigator. Requests will be reviewed by the study team, and approval will be based on alignment with ethical guidelines and data-sharing agreements.

Consent and anonymisation: All shared data will be fully anonymised, ensuring no participant can be identified. Participant consent has been obtained for the use and sharing of anonymised data.

Ethical and legal restrictions: Data sharing will comply with relevant ethical and legal requirements, including GDPR and the study's ethical approvals. Anonymisation will ensure privacy and data security.

#### IPD sharing plan summary

Available on request

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient- facing?
Other files	independent consultee information version 1.0	13/01/2025	29/01 /2025	No	No
Participant information sheet	version 1.0	13/01/2025	29/01 /2025	No	Yes
Participant information sheet	Participant information sheet	11/11/2025	11/11 /2025	No	Yes