INcidence of VENous Thromboembolism after acute stroke in China

Submission date	Recruitment status	Prospectively registered
28/04/2007	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
06/06/2007	Completed	Results
Last Edited	Condition category	Individual participant data
19/10/2021	Circulatory System	[] Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number INVENT-10

Study information

Scientific Title

INcidence of VENous Thromboembolism after acute stroke in China

Acronym

INVENT-China

Study objectives

- 1. Incidence of venous thromboembolism after acute stroke in China was lower than in western countries
- 2. Risk factors are different between Venous Thromboembolism (VTE) and non-VTE
- 3. Anticoagulation and neurological rehabilation can reduce the incidence

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approval received from the Tiantan Hospital Ethics Committee on the 29th November 2006 (ref: 7).

Study design

Multicentre observational, prospective, nested case-control study.

Primary study design

Observational

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Venous thromboembolism

Interventions

Group A: stroke inpatients with Deep Venous Thrombosis (DVT) during follow up Group B: stroke inpatients without DVT during follow up

DVT will be determined by complete-comprehensive ultrasound sonography two weeks (14 \pm 3 days) after stroke onset.

Possible risk factors will be compared between group A and group B, these will include:

- 1. stroke subtype
- 2. NIHSS
- 3. Medical history
- 4. Complications
- 5. High Density Lipoprotein (HDL)
- 6. Low Density Lipoprotein (LDL)
- 7. Triglycerides (TG)

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Incidence of Deep Venous Thrombosis after acute stroke in China, measured at two weeks (14 \pm 3 days) after stroke onset.

Key secondary outcome(s))

Predict model of VTE in acute stroke in China, measured at two weeks (14 \pm 3 days) after stroke onset.

Completion date

08/10/2007

Eligibility

Key inclusion criteria

- 1. Older than 18
- 2. Acute stroke patients within seven days
- 3. Identified by Computed Tomography (CT) or Magnetic Resonance Imaging (MRI)
- 4. National Institutes of Health Stroke Scale (NIHSS) item-six more than one

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Key exclusion criteria

- 1. Transient Ischaemic Attacks (TIAs)
- 2. Sub-Arachnoid Haemorrhage (SAH)
- 3. Tumour
- 4. Medical history of VTE

Date of first enrolment

08/05/2007

Date of final enrolment

08/10/2007

Locations

Countries of recruitment

China

Study participating centre

Tiantan Hospital

Beijing China 100050

Sponsor information

Organisation

Beijing Tiantan Hospital (China)

ROR

https://ror.org/003regz62

Funder(s)

Funder type

Industry

Funder Name

Glaxosmithkline (China)

Alternative Name(s)

GlaxoSmithKline plc., GSK plc., GlaxoSmithKline plc, GSK

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

United Kingdom

Funder Name

Beijing Municiple Science and Technology Commission (China)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary