Prevention of cerebrovascular and cardiovascular Events of ischaemic origin with teRutroban in patients with a history of ischaemic strOke or tRansient ischaeMic attack

Submission date	Recruitment status No longer recruiting	Prospectively registered		
22/03/2006		☐ Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
13/06/2006		[X] Results		
Last Edited	Condition category	[] Individual participant data		
25/01/2018	Circulatory System			

Plain English summary of protocol

Not provided at time of registration and not expected to be available in the future

Contact information

Type(s)

Scientific

Contact name

Prof Marie-Germaine Bousser

Contact details

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Additional identifiers

Clinical Trials Information System (CTIS)

2005-003700-10

Protocol serial number

CL3-18886-012

Study information

Scientific Title

Prevention of cerebrovascular and cardiovascular Events of ischaemic origin with teRutroban in patients with a history oF ischaemic strOke or tRansient ischaeMic attack

Acronym

PERFORM

Study objectives

To demonstrate the superiority of terutroban over a marketed drug (antithrombotic agent) in reducing the number of cerebrovascular and cardiovascular events in patients with cerebrovascular diseases

Ethics approval required

Old ethics approval format

Ethics approval(s)

First Ethics Committee approval in Italia on 06/12/2005, reference number: 42

Study design

Randomized double-blind parallel-group study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Ischaemic stroke

Interventions

S 18886 (terutroban) versus antithrombotic agent

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Terutroban

Primary outcome(s)

Cerebrovascular and cardiovascular events

Key secondary outcome(s))

Safety criteria

Completion date

31/03/2010

Eligibility

Key inclusion criteria

Male or female aged at least 55 years with recent history of stroke or transient ischaemic attack

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Haemorrhagic stroke

Date of first enrolment

22/02/2006

Date of final enrolment

31/03/2010

Locations

Countries of recruitment

Austria

Belgium

Bulgaria

Canada

Chile

China

Croatia

Czech Republic

Finland



France

Greece

India

Ireland

Hong Kong

Organisation

Institut de Recherches Internationales Servier (France)

ROR

https://ror.org/034e7c066

Funder(s)

Funder type

Industry

Funder Name

Institut de Recherches Internationales Servier (France)

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be available upon request from https://clinicaltrials.servier.com/ if a Marketing Authorisation has been granted after 2014.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created Date added	Peer reviewed?	Patient-facing?
Results article	results	11/06/2011	Yes	No
Results article	results	01/08/2014	Yes	No
Results article	results	01/04/2016	Yes	No
Results article	results	01/04/2017	Yes	No
Basic results			No	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes