Effect of polymorphic CYP2C19 genotype on the pharmacokinetics and pharmacodynamics of clopidogrel in healthy subjects

Submission date Recruitment status Prospectively registered 31/07/2007 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 17/09/2007 Completed [X] Results [] Individual participant data Last Edited Condition category Circulatory System 24/05/2019

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Effect of polymorphic CYP2C19 genotype on the pharmacokinetics and pharmacodynamics of clopidogrel in healthy subjects

Study objectives

To evaluate the pharmacogenetic effect of CYP2C19 gene on the Pharmacokinetics (PK) / Pharmacodynamics (PD) of clopidogrel. The original target was the patients with cardiovascular diseases taking clopidogrel. However, due to the limitation of enrollment the trial was conducted with healthy subjects with different CYP2C19 genotype to demonstrate that the effect of clopidogrel varies according to the patient's CYP2C19 genotype.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Institutional Review Board of Anam Hospital, Korea University College of Medicine, Seoul, Korea, approved on 12 February 2007 (ref: AN-06151-001).

Study design

Open-label, parallel, multiple-dose comparative study.

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular diseases

Interventions

After a loading dose of clopidogrel (300 mg; oral), patients will take a standard dose of clopidogrel 75 mg once a day for 6 days.

The following will be carried out:

- 1. Assessment of PK of clopidogrel and its metabolite SR26334
- 2. Measurement of the inhibition of ADP induced platelet aggregation by clopidogrel for 15 days
- 3. Evaluation of CYP2C19 phenotyping test using omperazole hydroxylation as a CYP2C19 probe

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

CYP2C19 gene

Primary outcome measure

Genetic association with biological effect of clopidogrel.

Secondary outcome measures

PK/PD relationship.

Overall study start date

01/03/2007

Completion date

31/08/2007

Eligibility

Key inclusion criteria

- 1. Healthy male subjects aged between 19 to 55
- 2. Wish to participate in the study
- 3. Informed consent for the trial

Participant type(s)

Patient

Age group

Adult

Sex

Male

Target number of participants

24

Total final enrolment

24

Key exclusion criteria

- 1. A history of or currently active clinically significant cardiac (including clinically significant abnormalities on Electrocardiogram [ECG] according to Principal Investigator [PI]), pulmonary, gastrointestinal, hepatic, renal, pancreatic, or neurological disease
- 2. Heavy smoker and alcohol consumer
- 3. Use of anticoagulants or medication within the last 1 month

Date of first enrolment

01/03/2007

Date of final enrolment

31/08/2007

Locations

Countries of recruitment

Korea, South

Study participating centre

126-1

Seoul Korea, South 136-705

Sponsor information

Organisation

Korea University (South Korea)

Sponsor details

126-1 Anam-dong 5-ga Sungbuk-gu Seoul Korea, South 136-705 +82 2 920 5114 webmaster@kumc.or.kr

Sponsor type

University/education

Website

http://www.korea.ac.kr/~eng/main.htm

ROR

https://ror.org/047dqcg40

Funder(s)

Funder type

Government

Funder Name

Anam Hospital, Korea University College of Medicine

Funder Name

The Korea Health 21 Research and Development Project, Ministry of Health and Welfare (South Korea)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/08/2008	24/05/2019	Yes	No