

Clopidogrel versus Aspirin in Chronic HEart failure

Submission date 11/01/2012	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 08/02/2012	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 24/04/2019	Condition category Circulatory System	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

<http://www.hta.ac.uk/2144>

Contact information

Type(s)

Scientific

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Additional identifiers

Clinical Trials Information System (CTIS)

2009-011637-27

Protocol serial number

HTA 08/53/36

Study information

Scientific Title

A randomised, open-label, study comparing the effects of aspirin and clopidogrel on outcomes in patients with chronic heart failure

Acronym

CACHE

Study objectives

Compared to clopidogrel, aspirin has an adverse effect on cardiovascular function in patients with heart failure leading to an increase in the symptoms of heart failure, reduced quality of life, an increase in hospitalisation and higher mortality.

Demonstrating whether such differences do or do not exist would be valuable and settle a substantial controversy in care, highlighted in international guidelines for the management of heart failure. The protocol designed to address this in a non-blinded, randomised controlled trial comparing aspirin and clopidogrel, conducted in 3,000 patients with heart failure over a period of approximately 5 years.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Northern & Yorkshire REC, 15/05/2012, ref: 10/H0903/13

Study design

Phase IV trial open-label interventional non-blinded randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Heart failure

Interventions

Clopidogrel vs aspirin both 75 mg/day per patient taken orally & daily.

This phase IV trial will compare the effects of prescribing aspirin 75 mg/day and clopidogrel 75 mg/day, primarily on all-cause mortality. Half of the patients will be randomised to each intervention. The investigator and the patient will know to which treatment the patient has been assigned. The duration of the study is determined by the event-rate although it could be stopped early for futility or because of a larger than anticipated treatment difference. The planned mean duration of follow-up is 3.8 years and some patients may be followed for more than five years before the required number of events has been attained. However, to allow for uncertainty we seek permission to follow patients for up to ten years.

Intervention Type

Drug

Phase

Phase IV

Drug/device/biological/vaccine name(s)

Clopidogrel, aspirin

Primary outcome(s)

All causes mortality

Key secondary outcome(s)

1. Cardiovascular death or hospitalisation for heart failure (time to first event)
2. Sudden death or a vascular event (myocardial infarction, stroke, peripheral embolism, requirement for angioplasty or vascular surgery) using a time to first event analysis
3. Total days lost to death or hospitalisation (all causes)
4. Quality-adjusted years alive (QALY) using repeated measurements of EQ5D
5. Cost per QALY

Completion date

31/08/2016

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility

Key inclusion criteria

1. Willing and able to provide written confirmation of informed consent
2. A clinical diagnosis of heart failure
3. Currently in sinus rhythm on clinical examination
4. Receiving diuretics for at least 6 weeks prior to inclusion
5. Patients must have a telephone
6. Patients must be willing to provide their personal contact details, those of their next of kin and those of their general practitioner (GP) and hospital to the national coordinating office and be willing to be contacted by telephone by these staff.
7. Patients must be willing to have hospitalisation and other serious events tracked through mechanisms including:
 - 7.1. In England:
 - 7.1.1. The National Health Service (NHS) Central Register (NHSCR)
 - 7.1.2. National Office of Statistics
 - 7.2. In Scotland:
 - 7.2.1. The Registrar Generals Office
 - 7.2.2. NHS Information Statistics Division
8. If non-English speaking, patients must have a friend or relative who can translate or who have other access to translation

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

87

Key exclusion criteria

1. Plasma (central lab) NT-proBNP <400pg/ml (47.3pmol/L) or MR-proANP <200pmol/L
2. Lack of an ECG within the previous 12 months documenting sinus rhythm. Patients who have had an episode of atrial fibrillation in the previous year may be enrolled provided the most recent ECG shows sinus rhythm and the treating doctor has decided not to prescribe anti-coagulants.
3. Severe valve disease in the investigators opinion (an echocardiogram report within the previous 12 months must be available)
4. Serum creatinine >250umol/L (local lab)
5. Intolerant of aspirin or clopidogrel or who have a contraindication to such treatment or who require anti-coagulation
6. Contraindications to aspirin or clopidogrel include:
 - 6.1. Substantial, in the investigators opinion, bleeding from an uncorrected source within the previous year
 - 6.2. Endoscopically proven peptic ulcer within the previous 3 months. Patients must be on treatment if peptic ulcer diagnosed in previous year
 - 6.3. Haemorrhagic stroke within the previous 3 months
 - 6.4. Known coagulation disorder (e.g. haemophilia)
 - 6.5. Full blood count suggesting iron deficiency (patients may be enrolled in the study after the cause of iron deficiency is or has previously been investigated and treatment has been initiated) (local lab)
7. Platelet count <100,000 (local lab)
8. Scheduled procedure that would require discontinuation of study medication for > 2 weeks (patient may be recruited after procedure)
9. History of uncontrolled seizures or high risks of falls
10. Regular use of non-steroidal anti-inflammatory agents > 3 times a week
11. Use of maintenance oral corticosteroids
12. Women of child-bearing potential or who are breast feeding
13. Patients with a history of asthma should not take part unless they have taken aspirin previously without ill-effect
14. Patients with an indication for oral anti-coagulation including:
 - 14.1 Current or recent (within 12 months) atrial fibrillation or flutter (evidence of an ECG documenting sinus rhythm must be provided)
 - 14.2. Prior embolic stroke
 - 14.3. Mechanical prosthetic heart valve
15. Patients requiring dual anti-platelet therapy including:
 - 15.1. Patients within 3 months of an acute coronary syndrome
 - 15.2. Transient ischaemic attack or vascular procedure or within one year of receiving a drug eluting coronary stent

16. Patients likely to die of something other than heart failure or sudden (cardiac) death
17. Inability to walk without the physical assistance of another person (patients with walking aids are permitted)
18. Other patients deemed unlikely to comply with the protocol
19. Women who are at pregnant or who could become pregnant
20. Women of child-bearing age should be taking reliable contraception (tubal ligation or implanted contraceptive)
21. Inability to communicate in English

Date of first enrolment

11/02/2011

Date of final enrolment

31/08/2016

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Castle Hill Hospital

Hull

United Kingdom

HU16 5JQ

Sponsor information

Organisation

Hull and East Yorkshire Hospitals NHS Trust (UK)

ROR

<https://ror.org/01b11x021>

Funder(s)

Funder type

Government

Funder Name

Health Technology Assessment Programme

Alternative Name(s)

NIHR Health Technology Assessment Programme, Health Technology Assessment (HTA), HTA

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	02/09/2014	18/04/2019	Yes	No
HRA research summary	Participant information sheet		28/06/2023	No	No
Participant information sheet		11/11/2025	11/11/2025	No	Yes