

A pilot study to determine the effects of corticotrophin releasing hormone (CRH) on myocardial function

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 07/07/2017	Condition category Circulatory System	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0295132800

Study information

Scientific Title

A pilot study to determine the effects of corticotrophin releasing hormone (CRH) on myocardial function

Study objectives

What is the effect of a single intravenous dose of CRH on myocardial contractility in normal, healthy subjects?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double-blind randomised crossover pilot study

Primary study design

Interventional

Secondary study design

Randomised cross over trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Cardiovascular: Myocardial function

Interventions

Single intravenous dose of CRH

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Corticotrophin releasing hormone

Primary outcome measure

Changes in measure of cardiac contractility/function as assessed by echocardiography: ejection fraction, fractional shortening, stroke volume, diastolic left ventricular filling, myocardial velocity, strain rate and strain.

Secondary outcome measures

Not provided at time of registration

Overall study start date

08/12/2003

Completion date

30/04/2004

Eligibility

Key inclusion criteria

Healthy men or women, 18-60 years of age.

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

60 Years

Sex

Both

Target number of participants

10 in total

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

08/12/2003

Date of final enrolment

30/04/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Coventry & Warwickshire Hospital

Coventry

United Kingdom

CV1 4FH

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

University Hospitals Coventry and Warwickshire NHS Trust (UK)

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