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Clinical Trial on the Therapy of the Reperfusion Injury during an Acute Myocardial Infarction

Submission date 13/06/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 01/09/2005	**Overall study status** Completed	
Last Edited 21/05/2008	**Condition category** Circulatory System	

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
N/A

Study information

Scientific Title

Acronym
Myoprotect II

Study objectives
Not provided at time of registration

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Acute myocardial infarction

Interventions
Randomised controlled trial:
A. Glutathione (GSH)/Cariporide
B. Placebo

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Glutathione (GSH), Cariporide

Primary outcome measure
Regional myocardial function 6 months after revascularisation

Secondary outcome measures
Infarct size, left ventricular ejection fraction

Overall study start date
01/07/2001

Completion date
31/05/2006

Eligibility

Key inclusion criteria
Age: 18-80, 1-6 hours after onset of chest pain, clinic and electrocardiogram (ECG) of an acute myocardial infarction, written informed consent

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Upper age limit
80 Years

Sex
Both

Target number of participants
100

Key exclusion criteria
1. Myocardial infarction within 28 days
2. Lysis therapy within 24 hours
3. Cardiogenic shock

Date of first enrolment
01/07/2001

Date of final enrolment
31/05/2006

Locations

Countries of recruitment
Germany

Study participating centre
Klinikum Großhadern
Munich
Germany
81377

Sponsor information

Organisation
German Research Foundation (Deutsche Forschungsgemeinschaft) (DFG)

Sponsor details
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Sponsor type
Research organisation

Website
http://www.dfg.de

ROR
https://ror.org/018mejw64

Funder(s)

Funder type
Research organisation

Funder Name
German Research Foundation (Deutsche Forschungsgemeinschaft) (DFG)

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