

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

<b>Submission date</b> 13/06/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 01/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 21/05/2008	<b>Condition category</b> 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

### 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

**Study type(s)**

Treatment

**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(s)**

Regional myocardial function 6 months after revascularisation

**Key secondary outcome(s)**

Infarct size, left ventricular ejection fraction

**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

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Upper age limit**

80 years

**Sex**

All

**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

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**Sponsor information****Organisation**

German Research Foundation (Deutsche Forschungsgemeinschaft) (DFG)

**ROR**

<https://ror.org/018mejw64>

## **Funder(s)**

### **Funder type**

Research organisation

### **Funder Name**

German Research Foundation (Deutsche Forschungsgemeinschaft) (DFG)

## **Results and Publications**

### **Individual participant data (IPD) sharing plan**

#### **IPD sharing plan summary**

Not provided at time of registration