

# Heparin in the treatment of ulcerative colitis

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 06/11/2014	<b>Condition category</b> Digestive 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**  
N0278078802

## Study information

## **Scientific Title**

### **Study objectives**

Is intravenous heparin of clinical benefit to patients with moderate to severe ulcerative colitis?

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Not provided at time of registration

### **Study design**

Double-blind randomised placebo-controlled study

### **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**

Digestive System: Ulcerative colitis

### **Interventions**

Intravenous heparin vs placebo

### **Intervention Type**

Other

### **Phase**

Not Applicable

### **Primary outcome measure**

Time to remission and time to next relapse.

### **Secondary outcome measures**

Not provided at time of registration

### **Overall study start date**

01/07/1997

**Completion date**

31/12/2005

## Eligibility

**Key inclusion criteria**

Patients with known acute, moderate/severe ulcerative colitis, with a Powell-Tuck score  $\geq 12$  on relapse, requiring inpatient treatment with intravenous steroids

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

Not provided at time of registration

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/07/1997

**Date of final enrolment**

31/12/2005

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Winchester Gastrointestinal Unit

Winchester

United Kingdom

SO22 5DG

## Sponsor information

**Organisation**

Department of Health (UK)

**Sponsor details**

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

**Sponsor type**

Government

**Website**

<http://www.doh.gov.uk>

**Funder(s)****Funder type**

Government

**Funder Name**

Winchester and Eastleigh Healthcare 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