ISRCTN73847754 https://doi.org/10.1186/ISRCTN73847754

# Randomised controlled trial comparing voltarol and paracetamol analgesia following oocyte collection in in vitro fertilisation (IVF) patients.

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|---|---|---|
| **Submission date**<br>30/09/2004 | **Recruitment status**<br>No longer recruiting | [_] Prospectively registered<br>[_] Protocol<br>[_] Statistical analysis plan<br>[_] Results<br>[_] Individual participant data<br>[_] Record updated in last year |
| **Registration date**<br>30/09/2004 | **Overall study status**<br>Completed | |
| **Last Edited**<br>19/02/2014 | **Condition category**<br>Pregnancy and Childbirth | |

**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**
N0016137178

# Study information

### Scientific Title

### Study objectives
A randomised controlled trial comparing the use of voltarol or paracetamol analgesia at oocyte retrieval to determine if voltarol users have a lower pregnancy rate and higher pregnancy loss rate.

### 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)
Not Specified

### Participant information sheet

### Health condition(s) or problem(s) studied
Pregnancy and Childbirth: Analgesia

### Interventions
Randomised controlled trial.

### Intervention Type
Other

### Phase
Not Specified

### Primary outcome measure
To determine if a relatively simple intervention (alternative analgesia to voltarol) could improve success rates.

### Secondary outcome measures

Not provided at time of registration

### Overall study start date
11/11/2003

### Completion date
10/11/2007

## Eligibility

### Key inclusion criteria
1. Women starting an IVF cycle
2. Ages 20-45

### Participant type(s)
Patient

### Age group
Adult

### Sex
Female

### Target number of participants
Not provided at time of registration

### Key exclusion criteria
Not provided at time of registration

### Date of first enrolment
11/11/2003

### Date of final enrolment
10/11/2007

## Locations

### Countries of recruitment
England

United Kingdom

### Study participating centre
**Reproductive Medicine**
London
United Kingdom
W12 0HS

# 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**
Hammersmith Hospital NHS Trust (NHS R&D Support Funding) - UK

**Funder Name**
IVF Unit Funds

# 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