

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

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 19/02/2014	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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
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