ISRCTN52633712 https://doi.org/10.1186/ISRCTN52633712

A randomized, placebo controlled, trial of preoperative sustained release betamethasone plus non-controlled intraoperative ketorolac or fentanyl on pain after diagnostic laparoscopy or laparoscopic tubal ligation

Submission date
18/08/2003

Recruitment status
No longer recruiting

Registration date
19/08/2003

Overall study status
Completed

Last Edited
05/09/2007

Condition category
Signs and Symptoms

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Roger Carroll

Contact details
UT Graduate School of Medicine
Department of Anesthesiology
1924 Alcoa Highway
Knoxville
United States of America
37920
+1 865 544 9469
rccarrol@mc.utmck.edu

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
Betamethasone ARF#1212

Study information

Scientific Title

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)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Postoperative pain

Interventions
Sustained release betamethasone versus placebo.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)

Betamethasone, ketorolac, fentanyl

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2002

Completion date
31/12/2002

Eligibility

Key inclusion criteria
Patients undergoing laparoscopic surgery at University of Tennessee Medical Centre Knoxville, Tennessee (USA)

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
74

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2002

Date of final enrolment
31/12/2002

Locations

Countries of recruitment
United States of America

Study participating centre

UT Graduate School of Medicine
Knoxville
United States of America
37920

Sponsor information

Organisation
University of Tennessee Anesthesiology Research Fund (USA)

Sponsor details
University of Tennessee Graduate School of Medicine
Department of Anesthesiology
1924 Alcoa Highway
Knoxville
United States of America
37920
+1 865 544 9220
prinehar@mc.utmck.edu

Sponsor type
University/education

ROR
https://ror.org/00xzqjh13

Funder(s)

Funder type
Research organisation

Funder Name
University of Tennessee Anesthesiology Research Fund (USA)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	21/08/2003		Yes	No