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Evaluation of effectiveness of pain control in patients after total hip replacement using the anaesthetic continuous-infusion device with bupivacaine

Submission date
03/03/2007

Recruitment status
No longer recruiting

Registration date
13/04/2007

Overall study status
Completed

Last Edited
09/05/2019

Condition category
Musculoskeletal Diseases

[X] 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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title
Evaluation of effectiveness of pain control in patients after total hip replacement using the anaesthetic continuous-infusion device with bupivacaine

Study objectives
The Anaesthetic Continuous-Infusion Device (ACID) with bupivacaine is effective for post-Total Hip Replacement (THA) pain control.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Approval received from the Institutional Review Board of the Chang Gung Memorial Hospital on the 25th December 2006 (ref: 95-1192B).

Study design
Rrandomised double-blind placebo-controlled study

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
Osteoarthritic hip

Interventions
Group I (25 patients): ACID intra-articular (above lessor trochanter) and normal saline 100 ml
Group II (25 patients): ACID inter-muscular (between tensor facia lata and gluteal medius) and normal saline 100 ml
Group III (25 patients): ACID intra-articular (above lessor trochanter) and bupivacaine 0.5% 100 ml
Group IV (25 patients): ACID intra-articular (between tensor facia lata and gluteal medius) and bupivacaine 0.5%100 ml

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Bupivacaine

Primary outcome measure
Pain control

Secondary outcome measures
Improve post-operative hip function and rehabilitation programs

Overall study start date
01/08/2007

Completion date
31/07/2009

Eligibility

Key inclusion criteria
Consecutive 100 patients with indications for total hip replacement.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
100 patients

Total final enrolment
92

Key exclusion criteria
Patients who refuse to join the clinical trial.

Date of first enrolment
01/08/2007

Date of final enrolment
31/07/2009

Locations

Countries of recruitment
Taiwan

Study participating centre
Orthopaedic Department
Taoyuan
Taiwan
333

Sponsor information

Organisation
Republic of China National Science Council (Taiwan)

Sponsor details
106 Ho-Ping East Road
Section 2
Taipei
Taiwan
106

Sponsor type
Government

ROR
https://ror.org/02kv4zf79

Funder(s)

Funder type
Government

Funder Name
Republic of China National Science Council (Taiwan)

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	01/05/2010	09/05/2019	Yes	No