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The efficacy of Chinese herbal recipe versus diclofenac in symptomatic treatment of osteoarthritis of the knee

Submission date
29/11/2004

Recruitment status
No longer recruiting

Registration date
29/11/2004

Overall study status
Completed

Last Edited
07/08/2007

Condition category
Musculoskeletal Diseases

[ ] 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
67/2002

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
Osteoarthritis (OA) of the knee

Interventions
1st Group: Chinese herbal medicine (6 capsules after meal, 3 times a day) plus placebo of diclofenac (1 capsule after meal, 3 times a day)
2nd Group: Placebo of Chinese herbal medicine (6 capsules after meal, 3 times a day) plus diclofenac (1 capsule after meal, 3 times a day)

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Chinese herbal recipe versus diclofenac

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
01/01/2004

Eligibility

Key inclusion criteria
Patients who had been suffering from unilateral or bilateral OA of the knee according to the criteria of the American College of Rheumatology for more than 3 months

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
200

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2002

Date of final enrolment
01/01/2004

Locations

Countries of recruitment
Thailand

Study participating centre

Department of Pharmacology
Chiang Mai
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Sponsor information

Organisation
Chiang Mai University - Faculty of Medicine (Thailand)

Sponsor details
-
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Sponsor type
University/education

ROR
https://ror.org/05m2fqn25

Funder(s)

Funder type
University/education

Funder Name
Chiang Mai University (Thailand) - Endowment Fund for Medical Research, Faculty of Medicine

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	13/12/2004		Yes	No