

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	<input type="checkbox"/> Prospectively registered
Registration date 29/11/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 07/08/2007	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> 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
Thailand
50200

Sponsor information

Organisation

Chiang Mai University - Faculty of Medicine (Thailand)

Sponsor details

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Sponsor type

University/education

ROR

<https://ror.org/05m2fq25>

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