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

Submission date	Recruitment status	Prospectively registered		
29/11/2004	No longer recruiting	☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
29/11/2004	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
07/08/2007	Musculoskeletal Diseases			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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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

-

Chiang Mai Thailand 50200 nimit@mail.med.cmu.ac.th

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