ISRCTN36233495 https://doi.org/10.1186/ISRCTN36233495

S-Adenosyl methionine (SAMe) versus celecoxib for the treatment of osteoarthritis symptoms: a double-blind cross-over trial

Submission date
12/02/2004

Recruitment status
No longer recruiting

Registration date
13/02/2004

Overall study status
Completed

Last Edited
16/08/2011

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

Contact name
Dr Wadie Najm

Contact details
101 The City Drive
Bldg 200 #512, Rt81
Orange
United States of America
92868

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
HS#2000-1046

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
Arthritis

Interventions
Participants were randomised to receive SAMe or Celebrex over two months. Pain, quality of life and musculoskeletal measures were assessed before, during and after the study.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
S-Adenosyl methionine (SAMe), celecoxib

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2000

Completion date
01/09/2002

Eligibility

Key inclusion criteria
Volunteers

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
61

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/09/2000

Date of final enrolment
01/09/2002

Locations

Countries of recruitment
United States of America

Study participating centre
101 The City Drive
Orange
United States of America
92868

Sponsor information

Organisation
University of California, Irvine, Medical Center (UCIMC) (USA)

Sponsor details
300 University Tower
Irvine
United States of America
92697

Sponsor type
Hospital/treatment centre

ROR
https://ror.org/00cm8nm15

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University of California, Irvine, Medical Center (UCIMC) (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	26/02/2004		Yes	No