

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	<input type="checkbox"/> Prospectively registered
Registration date 13/02/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/08/2011	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

Contact name

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/09/2002

Eligibility

Key inclusion criteria

Volunteers

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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)

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

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