ISRCTN18249834 https://doi.org/10.1186/ISRCTN18249834

Acupuncture versus metoprolol for interval treatment of migraine

Submission date	Recruitment status
12/01/2005	No longer recruiting
Registration date	**Overall study status**
29/07/2005	Completed
Last Edited	**Condition category**
10/09/2009	Nervous System 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
N/A

Study information

Scientific Title

Acronym
PEP-COMP

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)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Migraine

Interventions
Individualised acupuncture (up to 15 sessions) or 100 to 200 mg metoprolol

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Metoprolol

Primary outcome measure

Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/2002

Completion date
31/01/2005

Eligibility

Key inclusion criteria
Adult patients with migraine fulfilling the criteria of the International Headache Society (planned number of participants was 480; recruitment had to be terminated early, in summer 2004, due to recruitment problems and time limits of the project after recruiting 115 patients)

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
115

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/10/2002

Date of final enrolment
31/01/2005

Locations

Countries of recruitment
Germany

Study participating centre

Centre for Complementary Medicine Research
Munich
Germany
80801

Sponsor information

Organisation
Munich Technical University - Centre for Complementary Medicine Research (Germany)

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Sponsor type
University/education

ROR
https://ror.org/02kkvpp62

Funder(s)

Funder type
Other

Funder Name
The study is funded by a group of 10 German statutory sickness funds (Germany)

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	01/11/2006		Yes	No