ISRCTN84121379 https://doi.org/10.1186/ISRCTN84121379

Randomised controlled trial of gabapentin in Complex Regional Pain Syndrome type 1

Submission date 07/06/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 08/06/2004	**Overall study status** Completed	
Last Edited 12/09/2011	**Condition category** Musculoskeletal Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Wim E.J. Weber

Contact details
Department of Neurology
University Hospital Maastricht
PO Box 5800
Maastricht
Netherlands
6202 AZ
+31 (0)43 3875117
wweb@neurologie.azm.nl

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

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
Complex Regional Pain Syndrome (CRPS)

Interventions
Gabapentin versus placebo

Intervention Type
Drug

Phase
Not Specified

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

Primary outcome measure
Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date
19/11/1998

Completion date
02/12/1999

Eligibility

Key inclusion criteria
1. Patients with Complex Regional Pain Syndrome (CRPS)
2. With pain Visual Analogue Scale (VAS) greater than 3
3. Have undergone sympathetic blocks, mannitol infusions and Transcutaneous Electrical Nerve Stimulation (TENS) without success

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
58

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
19/11/1998

Date of final enrolment
02/12/1999

Locations

Countries of recruitment
Netherlands

Study participating centre
Department of Neurology
Maastricht
Netherlands
6202 AZ

Sponsor information

Organisation
University Hospital Maastricht (The Netherlands) - Pain Centre

Sponsor details
PO Box 5800
Maastricht
Netherlands
6202 AZ
+31 (0)43 3877498
sre@sane.azm.nl

Sponsor type
Hospital/treatment centre

ROR
https://ror.org/02d9ce178

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University Hospital Maastricht (The Netherlands) - Pain Centre

Funder Name
Parke-Davis Nederland (The Netherlands)

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	29/09/2004		Yes	No