

A randomised double-blind placebo-controlled trial to investigate the efficacy and side-effects of pregabalin for the treatment of neuropathic pain in humans following severe burn injury

Submission date 03/11/2004	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 09/12/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 12/01/2021	Condition category Signs and Symptoms	<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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

2004/142

Study information

Scientific Title

A randomised double-blind placebo-controlled trial to investigate the efficacy and side-effects of pregabalin for the treatment of neuropathic pain in humans following severe burn injury

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)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Neuropathic pain following burn injury may occur in the acute or chronic state.

Interventions

Randomised trial of pregabalin versus placebo

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

pregabalin

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2005

Completion date

31/01/2006

Eligibility

Key inclusion criteria

Patients who have severe burn injuries with elements of neuropathic pain.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2005

Date of final enrolment

31/01/2006

Locations

Countries of recruitment

Australia

Study participating centre

Multidisciplinary Pain Centre

Herston, QLD

Australia

4029

Sponsor information

Organisation

Royal Brisbane and Women's Hospital (Australia)

Sponsor details

Multidisciplinary Pain Centre
Royal Brisbane and Women's Hospital
Herston, QLD
Australia
4029

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Sponsor type

Not defined

ROR

<https://ror.org/05p52kj31>

Funder(s)

Funder type

Charity

Funder Name

Royal Brisbane and Women's Hospital Research Foundation

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	01/06/2011	12/01/2021	Yes	No