

The control of itching due to wound healing by gabapentin in children

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 02/11/2009	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0234117600

Study information

Scientific Title

Study objectives

Does gabapentin reduce itching when given to children with healing wounds - primary burn wounds?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised placebo 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

Itching

Interventions

50 children with itching causing distress following a burn will be randomized and receive placebo or 5 mg/kg gabapentin.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Gabapentin

Primary outcome measure

Reduction of itching.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2002

Completion date

01/11/2003

Eligibility

Key inclusion criteria

Children admitted to the Childrens Ward with burns who are distressed by itching.

Participant type(s)

Patient

Age group

Child

Sex

Both

Target number of participants

50

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/11/2002

Date of final enrolment

01/11/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Anaesthetics
Bristol
United Kingdom
BS16 1LE

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

North Bristol NHS Trust (UK)

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	01/12/2004		Yes	No