

Prevention by epidural Injection of postherpetic Neuralgia in the Elderly

Submission date 23/10/2003	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 24/11/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 31/03/2010	Condition category Infections and Infestations	<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
NWO 945-02-009

Study information

Scientific Title

Acronym

PINE

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)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Herpes zoster/postherpetic neuralgia

Interventions

Index group: Standard treatment (antiviral and analgesic medication) plus epidural injection of local anaesthetics and corticosteroids.

Control group: Standard treatment (antiviral and analgesic medication)

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Methylprednisolone-acetate, bupivacaine

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2004

Completion date

01/01/2005

Eligibility

Key inclusion criteria

Elderly patients (age >50 years) with acute herpes zoster (skin lesions <7 days) below dermatome C6 in primary care

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

550

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2004

Date of final enrolment

01/01/2005

Locations

Countries of recruitment

Netherlands

Study participating centre

Julius Center for Health Sciences and Primary Care
Utrecht
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Sponsor information

Organisation

Julius Center for Health Sciences and Primary Care (The Netherlands)

Sponsor details

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Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/0575yy874>

Funder(s)

Funder type

Research organisation

Funder Name

The Netherlands Organisation for Scientific Research (The Netherlands) (ref: 945-02-009)

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?
Protocol article	Protocol	26/01/2004		Yes	No
Results article	Results	21/01/2006		Yes	No