The clinical study of intercostal nerve pulsed radiofrequency in postherpetic neuralgia

Submission date	Recruitment status No longer recruiting	Prospectively registered		
10/06/2010		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
21/06/2010	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
26/07/2013	Nervous System Diseases			

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

Protocol serial number

N/A

Study information

Scientific Title

The clinical study of intercostal nerve pulsed radiofrequency in postherpetic neuralgia: A randomised controlled trial.

Study objectives

- 1. That pulsed radiofrequency of intercostals nerve was effective and safe in chronic refractory postherpetic neuralgia (PHN) patients
- 2. This pulsed radiofrequency can improve the quality of life and reduce the oral pain-related drugs dosage of PHN patients

Ethics approval required

Old ethics approval format

Ethics approval(s)

- 1. The ethics committee of Xinhua Hospital (affiliated to Shanghai Jiaotong Universty School of Medicine) approved on the 16th of March 2010 (ref: 2010-002)
- 2. The ethics committee of Shanghai Sixth People's Hospital approved on the 28th of October 2009 (ref: 2009-96)

Study design

Prospective double blind randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Postherpetic neuraglia (PHN)

Interventions

Patients were randomised to receive one of the following treatments, once a week for 3 weeks:

- 1. Pulsed radiofrequency (42 degree, 120 second) of the intercostal nerve (Thoracic 1 to T12 level)
- 2. Sham procedure, apparatus was left in test mode without the appropriate power output

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

- 1. Visual Analogue Scale (VAS) score, measured at baseline, days 3, 7, 14, 30 and 60, 6 months and 1 year
- 2. Flare pain per day during days 3, 7, 14, 30, 60 and at 6 months
- 3. Neurometer CPT/C (Neurotron, Inc) to measure three fiber function
- 4. SF-36 at baseline, 2 months, 6 months and 1 year following-up

Key secondary outcome(s))

- 1. Need to PRF again or other invasive operation at 2 day, 6 month or 1 year following-up
- 2. Dosage of rescue drug consumed per week at the end of treatment and after 4 weeks follow-up including Tramadol and/or acetaminophen and the dosage of combined drug, anti-

depressive drug (amitriptyline)

3. Presence, frequency and duration of adverse effects at 7, 14, 30, and 60 days

Completion date

28/05/2011

Eligibility

Key inclusion criteria

- 1. Age greater than 60 years
- 2. PHN history is longer than 3 months
- 3. Pain on Visual Analogue Scale (VAS) >3 (0-10 VAS scale)
- 4. PHN affected nerve sectors were thoracic nerves from T1 to T12 level
- 5. Refractory to formal treatment such as antiepileptic medicine, antidepressants, opioids and physical treatments and epidural block

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

All

Key exclusion criteria

- 1. Exclusion criteria included withdraws from the study
- 2. Poor effect, intolerant to the study
- 3. Uncooperative and unable to finish the self evaluation (VAS, QOL and SF-36)
- 4. Coagulation disturbances
- 5. Allergies to local anaesthetic
- 6. Malignancy

Date of first enrolment

22/02/2008

Date of final enrolment

28/05/2011

Locations

Countries of recruitment

China

Study participating centre

Department of Anesthesiology Shanghai China 200092

Sponsor information

Organisation

Xinhua Hospital (China)

ROR

https://ror.org/04dzvks42

Funder(s)

Funder type

Government

Funder Name

Shanghai Education Funding Committee (China)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created Date adde	d Peer reviewed?	Patient-facing?
Results article	results	01/01/2013	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/202	.5 No	Yes