

Acupuncture for diabetic neuropathic pain A protocol for systematic review and meta analysis

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Abstract

Background: Diabetic neuropathic pain (DNP) is a common complication of diabetes mellitus, it severely affects the quality of life of Diabetic patients. Acupuncture is proofed to have favorable effects in treating DNP, however, evidence needs to be gathered and interpreted. We will make a comprehensive review of clinical trials concerning acupuncture in treating DNP and do meta-analysis if possible.

Method: The following databases will be searched from the inception to September 2020: PubMed, Embase, Web of Science, China National Knowledge Infrastructure, Wan-Fang Database, and Chinese Scientific Journal Database. RCTs that evaluated acupuncture for patients with DNP will be included. The primary outcome will be patient-reported pain intensity using validated scales or verbal reporting. The secondary outcomes including the Toronto clinical scoring system, Sensory Nerve Conduction Velocity, Motor Nerve Conduction Velocity, and quality of life. The study selection, data extraction, and study quality evaluation will be performed independently by 2 researchers. A meta-analysis will be performed using RevMan V5.3 statistical software if possible; otherwise, descriptive analysis or subgroup analysis will be conducted. The quality of evidence for outcomes will be assessed with the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach.

Results: This study will evaluate the effect and safety of acupuncture in treating DNP.

Conclusions: The evidence we generated from the present study will provide more options for DNP management in clinical practice.

Systematic review registration: INPLASY202090043.

Abbreviations: CI = confidence interval, CNKI = China National Knowledge Infrastructure, DNP = diabetic neuropathic pain, GRADE = Grading of Recommendations Assessment Development and Evaluation, MNCV = motor nerve conduction velocity, PRISMA-P = Preferred Reporting Items for Systematic Reviews and Meta-Analysis Protocol, RCT = randomized controlled trials, SF-36 = 36-Item Short Form Health Survey, SNCV = sensory nerve conduction velocity, TCSS = Toronto clinical scoring system, VAS = visual analogue scales.

Keywords: acupuncture, diabetic neuropathic pain, meta-analysis, systematic review

1. Introduction

Diabetic neuropathic pain (DNP) is one of the most common complications of diabetes mellitus, affecting approximately

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Received: 9 October 2020 / Accepted: 20 October 2020 http://dx.doi.org/10.1097/MD.000000000023244 30%-50% of diabetes patients.^[1–3] The main symptoms of DNP include spontaneous pain and irritation-induced pain.^[4,5] Spontaneous pain can be manifested as constant burning, sharp, shooting, or even as electric shock sensations. Stimulus-induced pain includes hyperalgesia and allodynia. Hyperalgesia refers to stimuli that can cause pain under normal conditions, and the pain is more severe than normal; allodynia refers to stimuli that do not cause pain under normal conditions (such as tactile) leads to pain. DNP is usually considered moderate to severe and often worse at night, causing sleeping disturbs.^[4,5] The pain can be last for years, restrict daily activity, and affect the quality of life.^[6]

DNP is hard to treat. Its management and treatment require a series of approaches including early recognition, glycemic control, psychological therapy, and agents for symptomatic pain relief.^[7] Oral treatment is most commonly used as pain relief; however, it is associated with the risk of adverse effects since some DNP patients also suffer from multiple comorbidities and altered pharmacokinetics and pharmacodynamics that may alter drug metabolism. Also, pain reliefs currently used such as duloxetine, pregabalin, tapentado, and norepinephrine are unsatisfied for most DNP patients, new managements for DNP are needed.^[8,9]

Complementary and alternative medicine interventions are being used to treat DNP. Acupuncture and electroacupuncture are proved to be effective in pain management in many conditions There are several small sample size randomized controlled trials concerning acupuncture in treating DNP, however, to the best of our knowledge, there is no systematic review to summaries the results.^[11–13] Therefore, a comprehensive review of acupuncture in treating DNP is needed, and the evidence could provide more options for DNP management.

2. Methods

2.1. Study registration

This systematic review protocol has been registered on INPLASY. The registration number was INPLASY202090043. The protocol followed the Preferred Reporting Items for Systematic Reviews and Meta-Analysis Protocol (PRISMA-P) statement guidelines.^[14]

2.2. Eligibility criteria

2.2.1. Types of studies. We will include all randomized controlled trials (RCTs) regarding acupuncture in treating DNP. Nonrandomized clinical studies, cluster randomized trials, and quasi-randomized trials will be excluded. The language is limited to English and Chinese.

2.2.2. Types of participants. Patients diagnosed with DNP will be included regardless of sex, age, race, education, and economic status.

2.2.3. Type of interventions. Acupuncture is defined as needle stimulation of acupoints, we will include studies using body acupuncture, scalp acupuncture, manual acupuncture, auricular acupuncture, electro-acupuncture, fire needling et al. Studies using acupressure, moxibustion, laser acupuncture, pharmaco-acupuncture, transcutaneous electrical nerve stimulation will be excluded. Studies regarding acupuncture as adjunctive treatment compared with other treatments will also be included.

Control intervention will include any other treatments used to manage DNP other than acupuncture.

2.2.4. Types of outcome measures. The primary outcome measure will be patient-reported pain intensity using validated scales (e.g., visual analogue scales (VAS), numerical rating scales), or verbal reporting. The secondary outcomes including the Toronto clinical scoring system (TCSS), Sensory Nerve Conduction Velocity (SNCV), Motor Nerve Conduction Velocity(MNCV), and quality of life(36-Item Short Form Health Survey (SF-36)).

2.3. Search strategy

The reviewers will conduct a systematic literature search in the following electronic databases: PubMed, Embase, Web of Science, China National Knowledge Infrastructure (CNKI), Wan-Fang Database and Chinese Scientific Journal Database (VIP database). The search dates will be set from the inception to September 2020. The sample of the search strategy for PUBMED is presented in Table 1.

2.4. Data selection and extraction

2.4.1. Study selection. Paired investigators (ZG and XN) will independently screen all titles and abstracts to get qualified

Table	1		
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Number	Search items	
1	Acupuncture	
2	Acupuncture therapy	
3	Electroacupuncture	
4	Auricular acupuncture	
5	Needling	
6	Acupoints	
7	Moxbustion	
8	1 or2-7	
9	Diabetic neuropathies	
10	Autonomic Neuropathies, Diabetic	
11	Diabetic Neuralgias	
12	Diabetic Neuropathy, Painful	
13	Symmetric Diabetic Proximal Motor Neuropathy	
14	Diabetic Asymmetric Polyneuropathy	
15	Polyneuropathies, Diabetic	
16	9 or 10–15	
17	Randomized controlled trial	
18	Randomized	
19	Randomly	
20	Clinical trial	
21	17 or 18–20	
22	8 and 16 and 21	

studies, and then excluded duplications. After that, the full text of all potential studies will be checked for further screening. We will record all removed studies with specific reasons. Discrepancies will be resolved via referencing the original article and via group discussions or in consultation with the principal investigator (LW). The whole process of study selection is summarized as a flowchart in Figure 1.

2.4.2. Data extraction. Two independent investigators (ZG and MY) will extract and tabulated all data using a standardized data extraction form. Discrepancies will be resolved via referencing the original article and in consultation with the principal investigator (LW).

The following data will be extracted including leading author, year of publication, journal, country or region, study design, sample size, patients age, diagnostic criteria of DNP, type of intervention, controls, treatment frequency, outcome measures, adverse effects, and any other relevant information.

2.4.3. Assessment of risk of bias in included studies. The risk of bias of included study will be assessed using the Cochrane collaborative tool. We will evaluate the following aspects of the studies including sequence generation, assignment sequence hiding, blindness of participants and staff, outcome evaluators, incomplete result data, selective result reporting, and other sources of bias. The risk of bias is evaluated at 3 levels, namely, low risk, high risk, and ambiguity. If the information is not clear enough, we will try to contact the author of the article for further information.

2.5. Statistical analysis.

We will use Review Manager Software (RevMan) V.5.3 for data synthesis, meta-analysis. Mean difference or standardized mean difference and 95% confidence intervals (CIs) will be used to calculate quantitative data, and dichotomous data will be exerted as risk ratio and 95% CIs. Statistical heterogeneity across studies will be done with I^2 statistic. $I^2 \leq 50$ indicates homogeneity



among studies, and a fixed-effects model will be employed for pooled analysis. $I^2 > 50\%$ suggests obvious heterogeneity, and a random-effects model will be employed for synthesized analysis.

When there is homogeneity of the merged outcome results across sufficient studies, meta-analysis will be conducted.

2.6. Subgroup analysis

Subgroup analysis will be conducted based on the difference of interventions, controls, outcome measurements, and so on if necessary.

2.7. Sensitivity analysis

Sensitivity analysis will be undertaken to check the stability of merged outcome results by excluding studies with high risk of bias if significant heterogeneity exists.

2.8. Quality of evidence.

The quality of evidence for outcomes will be assessed with the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach.^[15]

The evaluation included bias risk; heterogeneity; indirectness; imprecision; publication bias. And each level of evidence will be made as very low, low, moderate, and high.

3. Ethics and dissemination

Since this is a protocol of systematic review and meta-analysis, ethics approval is not required. We will report our findings of this systematic review and meta-analysis in a peer-reviewed journal in the future.

Author contributions

Conceptualization: Liqin Wang, Yan Li, Fei Wang, Chuang Guo, Zhen Ren. Data curation: Zhaohong Gao, Xiangru Niu, Meiqi Yuan. Formal analysis: Zhaohong Gao, Xiangru Niu, Meiqi Yuan. Funding acquisition: Liqin Wang. Methodology: Yan Li. Project administration: Liqin Wang. Writing – original draft: Yan Li.

Writing - review & editing: Liqin Wang, Zhaohong Gao, Yan Li.

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