

Evaluating the Quality of Reports About Randomized Controlled Trials of Acupuncture for Low Back Pain

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Objective: This study aims to improve the reporting quality of randomized controlled trials (RCTs) by evaluating RCTs of acupuncture for low back pain (LBP) based on the CONSORT and STRICTA statements.

Methods: Literature from the Cochrane Library, Medline, Embase, Ovid, China National Knowledge Infrastructure (CNKI), WanFang database, and Chongqing Weipu (VIP) was systematically searched from 2010 to 2020. The general characteristics and the overall quality score (OQS) of the literature were evaluated by two investigators. The agreement between investigators was calculated using Cohen's kappa statistics.

Results: A total of 31 RCTs were extracted in the final analysis. Based on the CONSORT statement, the items "title and abstract", "background and objectives", "intervention", "outcomes", "statistical methods", "baseline data", "outcomes and estimation" and "interpretation" have a positive rate of greater than 80%. The items "implementation", "generalizability" and "protocol" have a positive rate of less than 30%. Based on the STRICTA statement, the items "style of acupuncture", "needle retention time", "number of treatment sessions", "frequency and duration of treatment" and "precise description of the control or comparator" have a positive rate of greater than 80%. The item "extent to which the treatment was varied" has a positive rate of less than 30%. The agreements among most items are determined to be moderate or good.

Conclusion: The reporting quality of RCTs of acupuncture for LBP is moderate. Researchers should rigidly follow the CONSORT and STRICTA statements to enhance the quality of their studies.

Keywords: acupuncture, quality of reporting, low back pain, CONSORT, STRICTA

Introduction

Low back pain (LBP), which is defined by an area of pain that is typically localized between the edge of the ribs and the crease of the hips,¹ is a problematic symptom. Once someone has problems with any part of the spine or part attached to the spine, such as lumbar intervertebral discs, ligaments, fascia, and muscles, LBP can occur.² The incidence of LBP was 7.3% globally in 2015,¹ meaning that approximately 540 million people suffered from LBP. In addition, it has been reported that one out of every six patients suffering from musculoskeletal problems is diagnosed with LBP.³ Although the number of LBP patients is large, LBP is usually tolerable. The prognosis represents a threat to LBP patients. Disability, for example, is the worst prognosis of LBP, representing a heavy burden to families and society.⁴ Due to poor

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medical conditions and harsh working conditions, the incidence of LBP in low-income countries or regions is rather high.⁵ According to the guidelines of LBP,⁶ nonsteroidal anti-inflammatory drugs (NASIDs) are recommended as the first-line therapy. Acupuncture is also recommended. However, NASIDs might cause bleeding⁷ and lead to gastrointestinal damage.⁸ Thus, patients have to pay extra money for other medicines that can protect the gastrointestinal tract. Compared with NASIDs, acupuncture seems safer because it has been reported to have fewer adverse events.⁹ Currently, acupuncture is increasingly accepted as a complementary or alternative therapeutic method. In America, approximately 54% of the population uses complementary or alternative therapy, especially for LBP.¹⁰ In China, the use of acupuncture is even higher. Therefore, acupuncture for LBP has attracted the attention of researchers, so many trials have been performed.

Established in 1996 and updated in 2017, the Consolidated Standards for Reporting Trials (CONSORT)^{11,12} has the goals of improving the transparency of trials and avoiding resource waste. The Standards for Reporting Interventions in Controlled Trials of Acupuncture (STRICTA)^{13,14} was established in 2002 and updated in 2010. As an extension of CONSORT for acupuncture, STRICTA has the goal of increasing the rigor of acupuncture trial design. To the best of our knowledge, there is no article evaluating the quality of acupuncture for LBP based on CONSORT and STRICTA statements. Based on the above two statements, this study evaluates the reporting quality of acupuncture for LBP.

Materials and Methods

Search Strategy

To identify all articles that studied the efficacy of acupuncture on LBP, the following databases were searched from January 2010 to December 2020: Cochrane Library, Medline, Embase, Ovid, China National Knowledge Infrastructure (CNKI), WanFang database, and Chongqing Weipu (VIP). In particular, we found that the trend of published articles about LBP was obviously increasing in the pilot search. Therefore, we aimed to search articles in the last 10 years. The following search terms were used in Chinese and English: (Acupuncture OR Acupuncture therapy OR Electro-acupuncture OR Manual acupuncture OR Warming acupuncture OR Auricular acupuncture OR Ear acupuncture OR Thread embedding acupuncture OR Motion style acupuncture) AND (Low Back

Pain OR Back Pain, Low OR Back Pain, Low OR Low Back Pain OR Pain, Low Back OR Pain, Low Back OR Lumbago OR Lower Back Pain OR Back Pain, Lower OR Back Pain, Lower OR Lower Back Pains OR Pain, Lower Back OR Pains, Lower Back OR Low Back Ache OR Ache, Low Back OR Aches, Low Back OR Back Ache, Low OR Back Aches, Low OR Low Back Aches OR Low Backache OR Backache, Low OR Backaches, Low OR Low Backaches OR Low Back Pain, Postural OR Postural Low Back Pain OR Low Back Pain, Posterior Compartment OR Low Back Pain, Recurrent OR Recurrent Low Back Pain OR Low Back Pain, Mechanical OR Mechanical Low Back Pain OR low back pain).

Included and Excluded Criteria

Types of Studies

We searched for randomized controlled trials (RCTs) that compared acupuncture with at least one control strategy for the treatment of LBP. The intervention of the control group can be another form of acupuncture or conventional treatment. RCTs without available data for extraction were excluded.

Types of Participants

All LBP patients of any age, gender and ethnicity were eligible. The clinical diagnosis of LBP was followed by expert consensus based on the site, duration, frequency and severity of the pain, excluding pain from feverish illness or menstruation.¹⁵ Patients who suffered from LBP for at least 6 months with or without lumbar disc protrusion screened by computed tomography or magnetic resonance imaging were included.

Types of Intervention

Different forms of acupuncture techniques or needles, such as manual acupuncture, electroacupuncture, warming acupuncture, auricular acupuncture, thread embedding acupuncture, and motion style acupuncture, were included. In particular, acupuncture plus cupping, moxibustion and Chinese medicine were not included in this research. The intervention in the experimental group was acupuncture alone or acupuncture combined with medication, which is similar to the control group. The control group used placebo acupuncture, sham acupuncture, no treatment or conventional treatment.

Placebo acupuncture means that a semiblunt retractable needle did touch but did not pierce the skin.¹⁶ Sham acupuncture refers to a needle set on nonacupuncture

points or acupuncture points not related to LBP.¹⁷ A modified nonfunctioning electroacupuncture stimulator was used to contrast with real electroacupuncture.

Selection of Reports

First, two investigators (HLL and DLZ) preliminarily searched RCTs according to the title and abstract on their own. Second, the investigators read the full text of reports for further selection following rigid inclusion and exclusion criteria. Third, after inspecting the selected reports for consistency, the studies were moved into the specified folders with different labels (included, excluded, undecided). Another magisterial investigator (LXZ) made a final decision regarding whether the reports in the “undecided” folder were included.

Data Extraction

Two investigators (HLL and DLZ) used Microsoft Excel 2019 to record study information, including author, publication year, language of the article, number of participants, intervention and course of treatment, from the final selection of RCTs. If the information was missing, then “no mention” was recorded. The investigators checked the sheet for consistency. Another magisterial investigator (ZLX) determined how to resolve any discrepancies noted in the sheets.

Assessment of Reporting Quality

Two investigators (ZQX and YTW) scored the reporting quality of RCTs of acupuncture for LBP individually based on the CONSORT and STRICTA statements. Before assessment, two investigators had a complete understanding of these two standards. Each item was scored 1 if it was reported and 0 if it was not mentioned or unclear (see the details in the [Supplementary Materials](#)). To assess the agreement between two investigators, Cohen’s κ -statistic¹⁸ was calculated using IBM SPSS Statistics version 26 (IBM SPSS Inc., Chicago, USA). According to Cohen’s definition, agreement was evaluated as perfect if κ was >0.8 , good if $0.6 < \kappa \leq 0.8$, moderate if $0.4 < \kappa \leq 0.6$, fair if $0.2 < \kappa \leq 0.4$, and poor if $\kappa \leq 0.2$.

Results

A total of potentially relevant RCTs were identified from 7 databases. After reading the title, abstract and full text, 31 RCTs were extracted for the final analysis. The whole selection process is depicted in [Figure 1](#), and the general

characteristics of the 31 included RCTs are summarized in [Table 1](#).

Year of Publication

In total, 31 RCTs were published in the last 10 years from 2010 to 2020. The average scores of each year based on the CONSORT and STRICTA statements are presented in the line chart ([Figure 2](#)).

Publication Language and Nationality of Authors

Among the 31 RCTs, 22 RCTs (71%) were published in English, whereas 9 (29%) were published in Chinese. The authors of these articles were from America, Australia, Brazil, China, France, Germany, India, Korea, New Zealand and Spain.

Invention

In the final extracted studies, different forms of acupuncture, including usual acupuncture (58.1%), electroacupuncture (12.9%), ear acupuncture (6.5%), internal heating acupuncture (6.5%), thread embedding acupuncture (6.5%), bee venom acupuncture (3.2%), floating acupuncture (3.2%) and motion style acupuncture (3.2%), were used.

Funding Resources

Sixteen studies (51.2%) reported their sources of funding. Nine (56.3%) received national funding, 2 (12.5%) received university funding, 4 (25.0%) received regional funding and 1 (6.3%) received personal funding. None of the included studies received funding from pharmaceutical companies.

Quality of Reporting

Reporting Quality Score Based on CONSORT Items

Based on the CONSORT statement, the data of overall quality of reporting are listed in [Table 2](#). Among all included studies, the median overall quality score (OQS) was 20, ranging from 8 to 27. Good reporting included terms of “title and abstract”, “background and objectives”, “intervention”, “outcomes”, “statistical methods”, “baseline data”, “outcomes and estimation” and “interpretation” with a positive rate of greater than 80%. Nevertheless, poor reporting was noted for terms of “implementation”, “generalizability” and “protocol” with a positive rate of less than 30%. All items have moderate, good or perfect

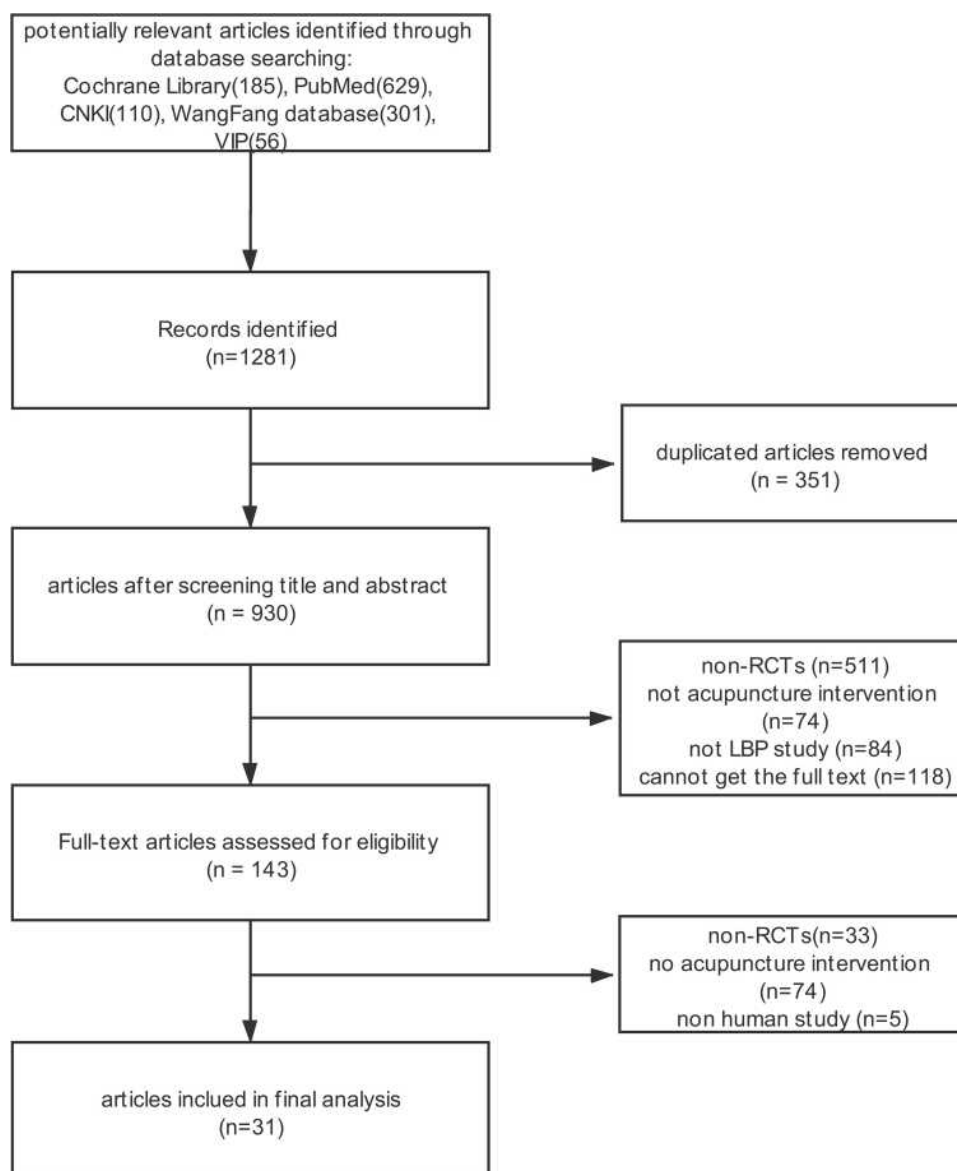


Figure 1 Flow chart of selection process. 31 RCTs were extracted for the final analysis.

agreement, except for the items of “background and objectives-2.a” and “generalizability”, which have fair agreement. OQS details are listed in [Table 2](#).

Reporting Quality Score Based on STRICTA Items

Based on the STRICTA statement, the data of overall quality of reporting are listed in [Table 3](#). Among all included studies, the median OQS was 12, ranging from 8 to 16. Good reporting was noted for the terms of “style of acupuncture”, “needle retention time”, “number of treatment sessions”, “frequency and duration of treatment” and “precise description of the control or comparator” with a positive rate of greater than 80%. Nevertheless, poor

reporting was noted for the term “extent to which treatment was varied” with a positive rate of less than 30%. All items had moderate, good or perfect agreement. OQS details are listed in [Table 3](#).

Discussion

This study first showed the reporting quality of RCTs that assessed the efficacy of acupuncture for LBP, adhering strictly to the CONSORT and STRICTA statements. Good quality RCTs not only reduce the bias of the trial but also contribute to the development of guidelines.⁴⁹ Therefore, a good quality study has positive meaning.

Table I General Characteristics of the Included 31 Studies

No.	Included Trials	Publication Language	No. of Participants		Intervention		Course of Treatment
			Trial	Control	Trial	Control	
1	Wasan 2010 ¹⁹	English	21	19	Acupuncture	Sham acupuncture	21d
2	Chen 2010 ²⁰	Chinese	50	50	Acupuncture	Transcutaneous electrical nerve stimulation	5w
3	Su 2010 ²¹	Chinese	30	30	Acupuncture	Sham acupuncture	1 session
4	Shankar 2011 ²²	English	30	30	Electro-acupuncture	Valdecoxib	3w
5	Hunter 2012 ²³	English	24	28	Ear acupuncture + exercise	Exercise	12w
6	Yun 2012 ²⁴	English	124	63	Acupuncture	Usual care	7w
7	Vas 2012 ²⁵	English	210	70	Acupuncture	Conventional treatment	4w
8	Cho 2013 ¹⁷	English	65	65	Acupuncture	Sham acupuncture	6w
9	Shin 2013 ²⁶	English	29	29	Motion style acupuncture	NSAIDs injection	1 session
10	Wand 2013 ²⁷	English	25	25	Sensory discrimination acupuncture	Usual acupuncture	1 session
11	Weib 2013 ²⁸	English	79	77	Acupuncture + standard rehabilitation programme	Standard rehabilitation programme	21d
12	Hasegawa 2014 ²⁹	English	40	40	Acupuncture	Sham acupuncture	5 sessions
13	Seo 2017 ³⁰	English	27	27	Bee venom acupuncture + Loxonin	Sham bee venom acupuncture + Loxonin	3w
14	Kizhakkeveetil 2017 ³¹	English	34	67	Acupuncture	Spinal manipulative treatment/ integrative care	60d
15	Liu 2017 ³²	English	30	15	4/7 sessions acupuncture	10 sessions acupuncture	12w
16	Zhang 2017 ³³	Chinese	30	60	Acupuncture	Sham acupuncture	1 session
17	Wu 2017 ³⁴	Chinese	30	30	Internal heating acupuncture	Warm acupuncture	3w
18	Zheng 2018 ³⁵	English	63	27	Electroacupuncture/sham acupuncture	Pain medication management (opioid medications)	10w
19	Lee 2018 ³⁶	English	20	20	Thread embedding acupuncture	Acupuncture	8w
20	Heo 2018 ³⁷	English	18	21	Electroacupuncture + usual care	Usual care	4w
21	Liu 2018 ³⁸	Chinese	42	42	Internal heating acupuncture	Warm acupuncture	10d
22	Luo 2019 ³⁹	English	104	48	Acupuncture	Usual care	7w
23	Vas 2019 ⁴⁰	English	165	55	Ear acupuncture + standard obstetric care	Standard obstetric care	2w
24	Nicolian 2019 ⁴¹	English	96	103	Acupuncture	Standard care	4w
25	Li 2019 ⁴²	Chinese	49	49	Tiaoshen acupuncture	Usual acupuncture	2w
26	Comachio 2020 ⁴³	English	33	33	Electro-acupuncture	Manual acupuncture	6w

(Continued)

Table 1 (Continued).

No.	Included Trials	Publication Language	No. of Participants		Intervention		Course of Treatment
			Trial	Control	Trial	Control	
27	Bishop 2020 ⁴⁴	English	69	41	Acupuncture + standard care	Standard care	6w
28	Sung 2020 ⁴⁵	English	19	19	Thread embedding acupuncture + acupuncture	Acupuncture	8w
29	Li 2020 ⁴⁶	Chinese	30	30	Six-directions acupuncture	Usual acupuncture	13d
30	Wang 2020 ⁴⁷	Chinese	34	34	Acupuncture at tendon lesions	Usual acupuncture	4w
31	Yang 2020 ⁴⁸	Chinese	99	99	Floating acupuncture	Usual acupuncture	10d

The OQS of RCTs based on the CONSORT statement was not satisfactory enough. Almost every included study prominently illustrated that the study was an RCT in the title. However, the positive rate of the items of randomization was low, and the positive rate for the item “implementation” was even less than 30%. Most of the studies stated that they allocated patients randomized but without details. The studies did not mention the method used to generate the random allocation sequence or the mechanism used to implement the random allocation sequence. Moreover, only 8 RCTs (25.81%) reported the person who generated the random allocation sequence. Accurate randomization eliminates the selection bias to the maximum extent and ascertains how well the randomization materials are performed.⁵⁰ Therefore, researchers should pay more attention to the randomization methodology to ensure that their study is randomized correctly and to improve the quality of the study.

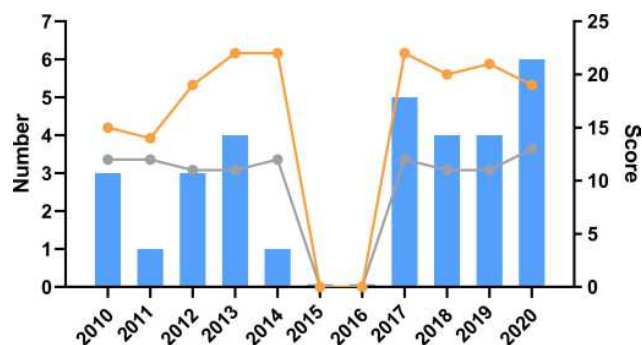


Figure 2 Year of publication. The blue bar is the number of RCTs published each year. The orange line is the median OQS with CONSORT statement of each year. The gray line is the median OQS with STRICTA statement of each year.

Given the particularities of acupuncture, many problems still need to be solved when considering blindness. Although advanced placebo acupuncture needles⁵¹ that broke the stereotype that only naïve acupuncture studies can be double-blinded⁵² were invented, it is difficult to completely and simultaneously blind patients and intervenors.⁵³ Therefore, most of the included RCTs blinded patients or investigators who analyzed the data. Despite the difficulties in blinding, researchers should make efforts employ blindness in the trials to eliminate the expectation bias that seems inevitable at present.

Due to the small sample size of the included RCTs, the positive rate of generalizability was less than 30%. This item is relatively subjective, and the agreement of the two investigators is low. Therefore, multicenter, large-scale RCTs are needed to improve generalizability.

To our surprise, only 6 RCTs (19.35%) reported that the protocol was accessed, whereas 18 RCTs (58.06%) reported their registration number. The importance of a protocol is that it describes the entire process and the details of a study. If one small step in the entire trial goes wrong, the whole trial may be worthless or might need to be performed again, wasting time and money. Magisterial experts can judge the feasibility of a study by reading the protocol and agree to the ethical assessment of the study. However, according to the results, the number of registered trials and trials with protocols that can be assessed is not equal. This finding indicates that some of the included trials might not have a rigorous design.

The OQS of RCTs based on the STRICTA statement do not reach satisfactory levels, especially for the item “extent to which the treatment was varied”. According to traditional Chinese medical theory,⁵⁴ different syndromes

Table 2 Details of OQS Assessed with CONSORT Statement (n=31)

Items	Items No.	Items Details	No. of Positive RCTs	%	Cohen's κ	95% CI
Title and abstract						
	1.a	Identification as a randomized trial in the title	29	93.55	0.78	0.38 to 1.00
	1.b	Structured summary of trial design, methods, results, and conclusions; for specific guidance see CONSORT for Abstracts	27	87.10	0.87	0.62 to 1.00
Introduction						
Background and objectives	2.a	Scientific background and explanation of rationale	25	80.65	0.37	0.05 to 0.69
	2.b	Specific objectives or hypotheses	29	93.55	0.48	0 to 1.00
Methods						
Trial design	3	Description of trial design including allocation ratio	26	83.87	0.76	0.46 to 1.00
Participants	4.a	Eligibility criteria for participants	31	100.00	0.89	0.69 to 1.00
	4.b	Settings and locations where the data were collected	21	67.74	0.50	0.19 to 0.81
Interventions	5	The interventions for each group with sufficient details to allow replication	31	100.00	0.89	0.69 to 1.00
Outcomes	6	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed	31	100.00	0.89	0.69 to 1.00
Sample size	7	How sample size was determined	17	54.84	0.81	0.61 to 1.00
Randomization						
Sequence generation	8.a	Method used to generate the random allocation sequence	16	51.61	0.49	0.18 to 0.79
	8.b	Type of randomization; details of any restriction	12	38.71	0.59	0.31 to 0.88
Allocation concealment	9	Mechanism used to implement the random allocation sequence, describing any steps taken to conceal the sequence until interventions were assigned	12	38.71	0.52	0.21 to 0.83
Implementation	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions	8	25.81	0.41	0.07 to 0.74
Blinding	11	If done, who was blinded after assignment to interventions and how	14	45.16	0.62	0.41 to 0.84
Statistical methods	12	Statistical methods used to compare groups for primary and secondary outcomes	28	90.32	0.64	0.18 to 1.00
Results						
Participant flow	13	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analyzed for the primary outcome	23	74.19	0.67	0.39 to 0.95
Implementation of intervention						
Recruitment	14	Dates defining the periods of recruitment and follow-up	18	58.06	0.74	0.49 to 0.98
Baseline data	15	A table showing baseline demographic and clinical characteristics for each group	26	83.87	0.67	0.34 to 1.00

(Continued)

Table 2 (Continued).

Items	Items No.	Items Details	No. of Positive RCTs	%	Cohen's κ	95% CI
Numbers analyzed	16	For each group, number of participants included in each analysis and whether the analysis was by original assigned groups	22	70.97	0.40	0.09 to 0.71
Outcomes and estimation	17	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision	31	100.00	1.00	–
Ancillary analyses	18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	23	74.19	0.67	0.37 to 0.96
Harms	19	All important harms or unintended effects in each group; for specific guidance see CONSORT for Harms	12	38.71	0.54	0.27 to 0.81
Discussion						
Limitations	20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	20	64.52	0.45	0.04 to 0.85
Generalizability	21	Generalizability (external validity, applicability) of the trial findings	8	25.81	0.38	0.09 to 0.67
Interpretation	22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	30	96.77	0.73	0.46 to 1.00
Registration	23	Registration number and name of trial registry	18	58.06	0.68	0.44 to 0.92
Protocol	24	Where the full trial protocol can be accessed, if available	6	19.35	0.89	0.64 to 1.00
Funding	25	Sources of funding and other support; role of funders	20	64.52	0.87	0.70 to 1.00

require different treatments. This principle is also applicable in acupuncture. Therefore, LBP patients need personalized acupuncture therapeutic schedules. However, only 3 RCTs (9.68%) reported changes in acupuncture, and the remaining researchers spent more time on the acupuncture rationale. We also found that the OQS values of Chinese RCTs were greater than those of English RCTs based on the STRICTA statement. However, the OQS values of Chinese RCTs were lower than those of English RCTs based on the CONSORT statement. Given that acupuncture can be traced back over 3000 years in China, the Chinese formed a relatively perfect therapeutic system. Chinese researchers might think more apprehensively when making acupuncture schedules. However, regarding the standard trial design, Chinese researchers were less thoughtful than foreign researchers. We found that the OQS trend of each year was quite flat (Figure 2), and the

average score of the included RCTs with STRICTA was 12 of 17, indicating that the reporting quality was always greater than moderate. Thus, we believe that spending more time on the acupuncture rationale is the key to enhancing the quality of acupuncture trials.

Although we systematically analyzed 31 RCTs, there are still some limitations in this study. Due to language barriers, we only included RCTs published in English and Chinese and excluded RCTs published in Korean or Japanese. Acupuncture is widely used in Korea and Japan, resulting in the loss of valuable data about the use of acupuncture for the treatment of LBP.

Conclusions

This study indicates that the reporting quality of RCTs that assessed the efficacy of acupuncture for LBP was moderate and needs further improvement to increase the

Table 3 The Details of OQS Assessed with STRICTA Statement (n=31)

Items	Item Details	No. of Positive RCTs	%	Cohen's κ	95% CI
1. Acupuncture rationale	1a) Style of acupuncture	31	100.00	1.00	–
	1b) Reasoning for treatment provided	24	77.42	1.00	–
	1c) Extent to which treatment was varied	4	12.90	0.43	0 to 0.89
2. Details of needling	2a) Number of needle insertions per subject per session	23	74.19	0.83	0.61 to 1.00
	2b) Names of points used	23	74.19	0.67	0.39 to 0.95
	2c) Depth of insertion	17	54.84	0.74	0.52 to 0.97
	2d) Response sought	16	51.61	0.81	0.61 to 1.00
	2e) Needle stimulation	19	61.29	0.49	0.23 to 0.75
	2f) Needle retention time	30	96.77	0.65	0.02 to 1.00
	2g) Needle type	24	77.42	0.74	0.46 to 1.00
3. Treatment regimen	3a) Number of treatment sessions	28	90.32	0.52	0.04 to 0.99
	3b) Frequency and duration of treatment sessions	29	93.55	0.48	0 to 1.00
4. Other components of treatment	4a) Details of other interventions administered to the acupuncture group	22	70.97	0.53	0.21 to 0.86
	4b) Setting and context of treatment	12	38.71	0.79	0.57 to 1.00
5. Practitioner background	5) Description of participating acupuncturists	13	41.94	0.48	0.18 to 0.78
6. Control or comparator interventions	6a) Rationale for the control or comparator in the context of the research question, with sources that justify this choice	20	64.52	0.86	0.67 to 1.00
	6b) Precise description of the control or comparator	25	80.65	0.59	0.23 to 0.95

level of evidence and guide clinical treatment better. In particular, the randomization methodology and acupuncture rationale should be explicitly explained in the article. These findings emphasize the need to improve the standard of operation. Therefore, we recommend that researchers draft acupuncture protocols rigidly based on the CONSORT and STRICTA statements to enhance the OQS of their studies, thereby convincing more people that acupuncture has good efficacy in the treatment of LBP.

Data Sharing Statement

The data used to support the findings of this study are included within the article and [Supplementary Information](#).

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work. In particular, Nanbu Wang, who has the overseas experience in Massachusetts General Hospital, polished up this article and edited the layout of this article. Thanks for her contribution.

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Disclosure

The authors declare that they have no conflicts of interest.

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