Quality of reporting of trial abstracts needs to be improved: using the CONSORT for abstracts to assess the four leading Chinese medical journals of traditional Chinese medicine

Ling Wang†1, Yulin Li†1, Jing Li*1, Mingming Zhang*1, Lin Xu1, Wenming Yuan2, Gang Wang3 and Sally Hopewell4

Abstract

Background: Due to language limitations, the abstract of journal article may be the only way for people of non-Chinese speaking countries to know about trials in traditional Chinese medicine (TCM). However, little is known about the reporting quality of these trial abstracts. Our study is to assess the reporting quality of abstracts of randomized controlled trials (RCT) published in four leading Chinese medical journals of TCM, and to identify any differences in reporting between the Chinese and English version of the same abstract publication.

Method: Two reviewers hand-searched the Chinese Journal of Integrated Traditional and Western Medicine, the Chinese Journal of Integrative Medicine, the China Journal of Chinese Materia Medica and the Chinese Acupuncture & Moxibustion for all abstracts of RCTs published between 2006 and 2007. Two reviewers independently assessed the reporting quality of the Chinese and English version of all eligible abstracts based on a modified version of the CONSORT for reporting randomised trials in journal and conference abstracts (CONSORT for abstracts).

Results: We identified a total of 345 RCTs of TCM with both a Chinese and English abstract. More than half of Chinese abstracts reported details of the trial participants (68%; 234/345), control group intervention (52%; 179/345), the number of participants randomized (73%; 253/345) and benefits when interpreting the trial results (55%; 190/345). Reporting of methodological quality or key features of trial design and trial results were poor; only 2% (7/345) included details of the trial design, 3% (11/345) defined the primary outcome, 5% (17/345) described the methods of random sequence generation, and only 4% (13/345) reported the number of participants analyzed. No abstracts provided details on allocation concealment and trial registration. The percentage agreement in reporting (between the Chinese and English version of the same abstract) ranged from 84% to 100% across individual checklist item.

Conclusion: The reporting quality of abstracts of RCTs published in these four TCM journals needs to be improved. Since none of the four journals adopted CONSORT for Abstracts, we hope that the introduction and adoption of CONSORT for Abstracts by TCM journals will lead to an improvement in reporting quality.
the only way for non-Chinese speaking countries to know about RCTs of TCM.

It is therefore important that abstracts of RCTs provide details of the trial participants, interventions, methodology (such as details of randomization, blinding, and intention-to-treat analysis) and the importance of the trial results (including estimates of benefits and harms, and their precision). To date, little is known about the quality of reporting of RCT abstracts in TCM. This study aims to assess and compare the reporting quality of both Chinese and English abstracts of RCTs published in four leading TCM Chinese medical journals using the recent CONSORT for reporting randomised trials in journal and conference abstracts (CONSORT for abstracts) [3].

Methods
Selection of journals
We selected four MEDLINE-indexed Chinese medical journals (Chinese Acupuncture & Moxibustion, Chinese Journal of Integrated Traditional and Western Medicine, Chinese Journal of Integrative Medicine, China Journal of Chinese Materia Medica) of TCM with a top ranking impact factor (IF) (ranked as No. 2, 3, 4 and 7) among the 38 journals in the category of Chinese traditional medicine and Chinese materia medica based on the 2007 data for Sci-tech Journal Citation Reports of China (Table 1). We selected these four high impact journals as they are indexed in MEDLINE and each abstract to be reported in English.

Identification of RCTs
Two reviewers (L Wang and YL Li) hand searched the four journals independently. We included all abstracts published between 2006 and 2007 that mentioned "randomization", "randomized allocation" or "randomized" in the abstract or full text. We excluded the abstracts reported as "quasi-randomized controlled trials", "laboratory study", "randomized sampling study", "meta-analysis", or "systematic review". All the eligible abstracts were retrieved from the full text database CNKI (China National Knowledge Infrastructure/Chinese Academic Journals full text Database).

Assessment of abstracts
We assessed the reporting quality of included Chinese and English abstracts based on criteria outlined in the CONSORT for abstracts [4]. We refined several of the items included in the CONSORT for Abstracts checklist to assess certain items in more detail and to add some unique characteristics relevant to TCM. For example, we expanded "interventions" to "interventions intended for experimental group" and "interventions intended for control group", and expanded the item "blinding" to "specified exactly who was blinded" and "simply described the trials as single blind or double blind". We assessed each item as being "reported" or "not reported" according to whether the author had reported all the contents listed in the refined items or not.

Two reviewers (L. Wang and Y.L. Li) assessed each abstract independently after discussing and comprehending the items thoroughly with J Li and MM Zhang. We sought further clarification from S Hopewell if there was any problem with understanding of the CONSORT for Abstracts checklist criteria. A pilot study was performed on 10 RCT abstracts selected at random (both the Chinese and English version of the same abstract) and inter-rater agreement was calculated using the kappa statistic. Discrepancies were resolved by discussion. Once we had ensured consistency in the interpretation of the data extraction form we carried out double data extraction on all remaining abstracts. We determined the overall number and proportion (%) of RCT abstracts that reported each of the CONSORT for Abstracts item. We also assessed the consistency in reporting between the Chinese and English version of the same abstract using the kappa statistics. Data were analyzed using Microsoft Excel 2003 and SAS version 9.1.

Table 1: Number of abstracts identified and included from four TCM a of Chinese medical journals (2006 to 2007)

| Journals                                      | IF b   | Identified RCT c | Included abstracts |
|-----------------------------------------------|--------|------------------|--------------------|
| Chinese Acupuncture and Moxibustion           | 0.437  | 165              | 154                |
| Chinese Journal of Integrated Traditional and Western Medicine | 0.837  | 261              | 158                |
| Chinese Journal of Integrative Medicine       | 0.598  | 21               | 19                 |
| China Journal of Chinese Materia Medica       | 0.634  | 48               | 14                 |
| Total                                         | 495    | 345              |                    |

a TCM: traditional Chinese medicine
b Impact factor (IF) comes from «2007 data for sci-tech Journal Citation Reports of China»
c RCT: randomized controlled trial
Results
We identified 495 RCTs from hand searching the full-text and abstracts of the four leading TCM journals from 2006 to 2007, of which 345 (70%) included both the Chinese and English version of the same abstracts and all were structured abstracts as it is a formal requirement for the four journals. The remaining 150 RCTs were published as short reports with no abstract. Seven RCT reports mentioned “randomized” in the abstract but not in the full text, while 18 mentioned “randomized” in the full text but not in the abstract. Eighty-five papers mentioned “randomized” in the abstract but were judged as “quasi-randomization” after reading the full text. Table 1 provides information on the impact factor for the four journals, the number of RCTs identified and the number of reports with both the Chinese and English version of the same abstracts for each journal. Agreement between the two reviewers was good with a kappa score of greater than 0.75 for each item.

Table 2 presents the criteria used to judge whether the abstract reported the item, and the number and proportion (%) of items reported for both the Chinese and English version of the same abstracts.

Reporting quality of Chinese RCT abstracts

General items
Only 8% (28/345) abstracts could be judged as RCTs from their titles, other abstracts were most described as "clinical trial", "comparison research", "effect comparison", or "effect observation". Only one (1/345) abstract did not report the author’s addresses, seven abstracts reported the trial design as "parallel comparison RCTs", no abstract reported details of trial registration and only 27% (93/345) reported the source of funding.

Trial methodology
A total of 68% (234/345) of abstracts reported details of participants included in the trial; the remainder simply used the term "patient". Forty-seven percent (163/345) of abstracts reported details of the experimental intervention and 52% (179/345) the control intervention. Forty-two percent (144/345) of abstracts reported the name of the experimental intervention and 27% (92/345) the control intervention, 11% (38/345) described the experimental intervention as "Chinese medicine" and 21% (74/345) of the control intervention as "western medicine". Only 14% (48/345) of abstracts reported the study objective, 5% (17/345) the method of random sequence generation; no abstracts mentioned details of allocation concealment. A total of 11% (39/345) mentioned the trial as "single blind" or "double blind"; only one specified exactly who was blinded.

Trial results
Seventy-three percent (253/345) of abstracts reported the number of participants randomized to each group, however, only 4% (13/345) reported the number of participants included in the analysis. Fifty-seven percent (195/345) of abstracts reported the result for each group and 44% (150/345) reported a P value for the between-group comparison, however only two abstracts reported the effect size and its precision. Fifteen percent (50/345) of abstracts mentioned adverse events or side effects of the treatment.

Trial conclusion
Four abstracts clearly stated the benefits and harms of the results when interpreting the clinical application, with 55% (190/345) of abstracts reporting only the benefits.

Comparison of the Chinese and English of the same RCT abstracts
The proportion of abstracts reporting each checklist item was almost the same between the Chinese and corresponding English abstracts (Table 3). The percentage agreement in reporting (between the Chinese and English version of the same abstracts) ranged from 84% to 100% across individual checklist item. The median of kappa statistic for each item was 0.86 (0.64-1.00). The items where there was most disparity between the Chinese and English abstracts were details of the trial participants and the number of participants randomized.

Discussion
Our study identified 345 of RCTs published in four leading Chinese TCM journals with both the Chinese and English version of the same abstracts between 2006 and 2007. Based on the CONSORT for Abstract checklist criteria, the quality of reporting of trial abstracts in these journals was far from satisfactory. Except for details of the trial authors, participants, the intervention for control group, and the number of participants randomized in the trial, all other items were reported in less than half of the abstract reports. Reporting of methodological quality or key features of trial design and trial results was even worse. Less than 5% of the included abstracts fully or correctly reported relevant items including details of the trial design, clearly defining the primary outcome, the method of random sequence generation and allocation concealment, specifying details on who was blinded, the number of participant analyzed in each arm of the trial, and the estimated effect size and its precision for the primary outcome.

Our study has several strengths. We hand searched four high impact TCM medical journals to ensure that all RCTs were identified. In addition, all items in modified version of the CONSORT for Abstracts checklist were carefully discussed and understood within the research team. The identification of RCTs, eligibility decision and data extraction were conducted in duplicate. However, our study has limitations. We did not take a random sample of all RCT published in TCM medical journals and
therefore our findings may not reflect the whole picture of reporting quality of trial abstracts in the TCM field.

Our findings are consistent with or worse than previous studies that have assessed the quality of reporting of journals RCT abstracts and conference RCT abstracts. Previous work has shown that abstracts frequently under report key features of study design [1,2,5] and omit important results. In the mid-1990s, Li and Hu reported that the quality of abstracts in Chinese medical journals was poor [6,7]. The problem is particularly apparent when trying to assess details of methodological quality. A recent study evaluating 227 RCT abstracts published in four high impact general medical journals in 2006 found that only one abstract reported details of allocation concealment, 23% reported the use of intention-to-treat analysis, 14% reported loss of follow-up, and 9% reported details of blinding [8]. A further study evaluating RCT abstracts presented at the American Society of Clinical Oncology Conference and found no abstracts reported allocation concealment, 16% reported on blinding, and only 14% reported intention-to-analysis [9].

Possible explanations for our findings of poor reporting include poor reporting of RCT full-text articles in China, when an item isn't reported in the full publication, it may

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**Table 2: Criteria for assessing the quality of reporting of abstracts of RCTs in TCM**

| Items               | Criteria for judgment                                                                 | No. of abstracts reporting each item (n = 345) |
|---------------------|----------------------------------------------------------------------------------------|-----------------------------------------------|
|                     |                                                                                       | Chinese (%) | English (%)   |
| **General items**   |                                                                                       |                |               |
| Title               | Mentioned randomized or randomly allocation in the title                               | 28(8)         | 25(7)         |
| Authors             | Reported the contact details of corresponding author, e.g. postal address, email       | 344(99)       | 344(99)       |
| Trial design        | Described as "parallel group", "cluster randomized", "crossover", etc.                | 7(2)          | 5(1)          |
| Trial registration  | Reported the trial registration number and name of trial register                      | 0(0)          | 0(0)          |
| Funding             | Reported the source of funding for the trial                                          | 93(27)        | 0(0)          |
| **Methods**         |                                                                                       |                |               |
| Participants        | Reported the details of the participants (included the demography, diagnosis, TCM syndrome) | 234(68)       | 239(69)       |
| Interventions       | Reported the details of the interventions for experimental group (included denomination, usage and course of treatment) | 163(47)       | 152(44)       |
|                     | Reported the details of the interventions for control group (included denomination, usage and course of treatment) | 179(52)       | 168(49)       |
| Objective           | Described as "to compare interventions(A) and (B) in patients with condition(C)"     | 48(14)        | 44(13)        |
| Outcome             | Defined the primary outcome                                                           | 11(3)         | 10(3)         |
| Randomization       | Reported the method of random sequence generation                                     | 17(5)         | 14(4)         |
|                     | Mentioned the allocation concealment (such as central randomization by telephone)     | 0(0)          | 0(0)          |
| Blinding            | Specified exactly who was blinded                                                     | 1(0.3)        | 0(0)          |
|                     | Simply mentioned the trial as "single blind", or "double blind"                     | 39(11)        | 39(11)        |
| **Results**         |                                                                                       |                |               |
| Numbers randomized  | Reported the number of participants randomized to each group                           | 253(73)       | 253(73)       |
| Numbers analyzed    | Reported the number of participants included in analysis for each group               | 13(4)         | 12(4)         |
| Outcomes            | Reported the result for each group (if the primary outcome not defined, reported the result of one listed outcome) | 195(57)       | 190(55)       |
|                     | Reported effect size, precision for the difference between groups                     | 2(1)          | 2(1)          |
| Harms               | Reported important adverse events or side effects                                      | 50(15)        | 48(14)        |
| **Conclusions**     |                                                                                       |                |               |
|                     | Clearly stated the benefits and harms when interpreting the results                   | 4(1)          | 3(1)          |
|                     | Only stated the benefits when interpreting the results                                 | 190(55)       | 190(55)       |
also be missing from the abstract. A study evaluating the reporting quality of 142 RCT full-text articles published in five leading Chinese medical journals found that only 27% described the method to generate the randomized sequence, 4% had adequate allocation concealment, and only 17% mentioned blinding [10]; these findings are supported by others in the field of TCM [11,12].

Lack of detailed requirements for structured abstracts in journal’s “Instruction for Authors” in China is also a problem since none of the journals included in our study mentioned CONSORT for Abstracts in their Instructions to Authors. Some studies have proved that journal adoption of structured abstracts can improve the reporting of abstracts [13-17]. Although most of the Chinese medical journals have adopted four-heading structured abstracts (objectives, methods, results and conclusion), the structured format alone is insufficient to guide authors regarding their content for reporting RCTs.

Finally, the word limitation of an abstract restricts the amount of detail an author is able to provide. Most Chinese medical journals have abstract word restrictions, usually within 150-500 Chinese words, making it difficult for authors to comprehensively report trial methods and results, especially when authors are not aware the importance of research methodology and the way to correctly present the clinical relevance of research results. The word limit for the journals we assessed is 150-250 Chinese words and 150-400 English words.

**Conclusion**

In summary, we found that the quality of reporting of trial abstracts in these four leading TCM medical journals is far from satisfactory, especially with respect to details of the trial methodology. Many studies have proved that adoption of the CONSORT Statement has demonstrated benefits in improving the reporting quality of RCT of full-text articles [18], and we hope that it will do the same with the introduction of CONSORT for Abstracts in Chinese medical journals of TCM field.

**Competing interests**

This research is supported by a project of the Cochrane Collaboration entitled "Making Cochrane review more accessible to policy makers", but no funding for the publication. Dr Sally Hopewell is a lead author on the CONSORT extension for reporting abstracts of randomized trials.

**Authors’ contributions**

LW and YL participated in the design of the study, carried out the hand searching of journals, assessment of the abstracts and drafting the manuscript. JL, MZ and SH conceived the study, interpreted the data and revised the manuscript. LX and WY participated in the design of the study and performed the data analysis. GW participated in the design of the study, selection of journals and interpretation of the data. All authors read and approved the final manuscript.

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**Author Details**

1. Chinese Cochrane Centre, West China Hospital, Sichuan University, Chengdu, Sichuan, China; 2. Department of Nephrology, West China Hospital, Sichuan University, Chengdu, Sichuan, China; 3. Department of Integrated Chinese Medicine and Western Medicine, West China Hospital, Sichuan University, Chengdu, Sichuan, China; 4. NHS R&D Programme, UK Cochrane Centre, Oxford, UK

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