Title: The Effectiveness of Participant Blinding of Non-Penetrating Sham/Placebo Acupuncture in Clinical Trials: A Systematic Review with Meta-Analysis

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Abstract

**Background:** Acupuncture clinical trial is important to evaluate the efficacy of acupuncture. However, it is challenging to achieve effective blinding due to the nature of acupuncture. A standardised placebo control method of acupuncture has yet to be established. The study focuses on the non-penetrating sham acupuncture because it eliminates the placebo effect and generates lesser physiological responses. The study aims to evaluate and compare the participant blinding effectiveness of non-penetrating sham acupuncture devices, and analyse the factors which may influence the participant blinding.

**Methods:** The study followed the PRISMA guidelines. An electronic search was conducted on PubMed, Ovid and CNKI up until 1st of October 2020 to include English and Chinese randomised controlled trials which evaluated the awareness on the type of acupuncture (real or sham) in any population who received acupuncture. Data screening, data extraction and quality assessment were done independently by two researchers and discrepancies were sorted out via discussion with a co-researcher. Data analysis was performed using RevMan 5.4.1.

**Results:** 34 full-text articles had been included in the systematic review and meta-analysis. The quality of the studies ranged from moderate to good. Generally, non-penetrating sham acupuncture devices were effective in blinding participants in clinical trials. The foam device demonstrated a better blinding effect, followed by Streitberger, Park and Takakura devices. Sham needles with no skin contact could not blind the participants successfully. Naive, experienced, healthy and diseased participants all could be blinded using non-penetrating sham acupuncture devices but naive and healthy participants could be blinded comparatively easily. Acupoints from different regions could achieve blinding, however, the acupoints on the back could blind the participants more easily compared to the other areas.

**Conclusion:** Non-penetrating sham acupuncture devices are valid placebo control for acupuncture clinical trials. The foam device has a better blinding effect, followed by Streitberger, Park and Takakura devices. Recruiting naive healthy participants and choosing acupoints from the back can achieve better blinding effects in the participants.

**Background:**

Acupuncture has been popular in Asia, serving as one of the major treatment methods for thousands of years. In the sixteenth century, it had been introduced to Europe and America. After that, it gradually spread all around the world and gained popularity due to its therapeutic effects, including analgesic, anaesthetic, mind-calming and body regulating effects. To investigate its clinical efficacy, acupuncture research was initiated in the eighteenth century and conducted continuously since then.

A well-blinded randomised controlled trial (RCT) is the gold standard of clinical trials because it minimises the risk of bias and maximises the validity of the results. In an ideal acupuncture clinical trial, subjects should be randomly assigned to a treatment or control group, and both the participants and acupuncturists should be blinded. It is challenging to achieve effective blinding in acupuncture clinical trials due to acupuncture’s nature, whereby the participants can feel the needling sensation and deduce the grouping, whereas the acupuncturists are not blinded most of the time because they need to know where and how to needle the participants, and can also feel the penetrating sensation from their fingers.

There are several types of control groups that are commonly used, such as no treatment, standard/conventional treatment, sham/placebo acupuncture and minimal acupuncture (shallow needle insertion). Non-treatment or standard/conventional treatment as the control cannot eliminate placebo effects. Moreover, skin penetrating sham acupuncture as the control may trigger physiological responses. For instance, it can deactivate limbic structures and, hence, reduce pain levels in patients who suffer from pain. Thus, to assess the true efficacy of acupuncture, selecting a good non-penetrating sham/placebo method is necessary.

Currently, a standardised sham/placebo acupuncture method has yet to be established. Various types of non-penetrating sham devices have been developed to raise the quality of clinical trials and the validity of the results, including Streitberger device, Park device, Takakura device and foam device. Many studies tried to evaluate the blinding effectiveness of individual sham devices but the sample size was small for most of the studies. There was insufficient evidence to demonstrate the superiority of a specific type of sham device in achieving participants’ blinding. Therefore, there is a need to perform a systematic review with meta-analysis to evaluate the blinding effectiveness of these sham acupuncture devices.

Zhang et al conducted a systematic review on the credibility of blinding healthy participants and/or acupuncturists using placebo acupuncture devices. The author also studied the penetrating sensation and DeQi sensation of the devices. The study found that the real and sham Streitberger and Park devices were significantly different in penetrating sensation when applied at sensitive acupuncture points, which may lead to failure in participants’ blinding. However, there was no meta-analysis performed to compare the blinding effectiveness of different sham acupuncture devices. The study only included the results in phase one which could lead to an incomplete assessment of the outcome, especially for cross-over studies.

Gong et al carried out a systematic review with meta-analysis on the blinding effectiveness of non-penetrating sham needles. The study included five articles and found that participants were not able to differentiate between real and sham acupuncture. The author pointed out that it may be necessary to analyse the influence of age, sex, ethnic admixture and other factors. The number of included studies was too small to perform a meaningful comparison among devices.

In short, a limited number of studies had systematically reviewed and analysed the blinding effectiveness of sham acupuncture devices. A comparison among sham acupuncture devices was not performed to show the superiority of a specific sham device in blinding participants. Factors which may influence the blinding results, for example, acupuncture experience and health status of participants, and needling locations, were not statistically analysed in the existing
studies. As a result, this study aims to assess and compare the blinding effectiveness of non-penetrating sham acupuncture devices as well as analyse the factors which may influence the blinding effectiveness.

**Methods**

This systematic review and meta-analysis followed the Preferred Reporting Items for Systematic Reviews and Meta-Analysis Statement (PRISMA) guidelines.

**Search Strategy**

An electronic search was carried out in health-related databases such as PubMed, Ovid and CNKI for relevant studies. The literature search was conducted using search terms such as sham acupuncture, placebo acupuncture, sham needle, placebo needle, Park needle, Park device, Streitberger needle, Streitberger device, Takakura needle and Takakura device. Similar search terms in Chinese were used to search in CNKI, for example, , , , and . The search was limited to original articles published in English and Chinese language until 1st October 2020. Additional studies were identified manually from reference lists of potentially eligible articles. Title and abstract of those studies were screened to determine its relevance.

**Inclusion Criteria**

Selection of primary studies for this review was derived from the following pre-specified criteria (in PICOS format).

- **Type of participants (P)** Any population who received acupuncture treatment regardless of age, sex, race, region, underlying disease and acupuncture experience
- **Intervention (I)** Non-penetrating sham/placebo acupuncture
- **Comparison (C)** Real penetrating acupuncture
- **Outcomes (O)** Awareness of participants on the type of acupuncture
- **Type of studies (S)** Randomised Controlled Trials (RCTs)

**Exclusion Criteria**

Studies were excluded if the studies were animal studies, review articles, case reports, editorials, letters and comments. Studies published other than English and Chinese were excluded. Studies which did not meet any of the inclusion criteria were also excluded.

**Data Extraction**

Data were extracted by using a review spreadsheet, containing information such as author, publication year, study design (blinding and sample size), participant details (age, gender, health status and acupuncture experience), acupuncture methods (the type of sham acupuncture device and needling location) and results. Corresponding authors were contacted if the data was unclear. The data extraction was conducted separately by two researchers. Any discrepancies were sorted out via discussion with a co-researcher.

**Quality Assessment**

Quality and risk of bias of the eligible studies were assessed independently by two researchers at outcome level by using Cochrane risk of bias tool. Quality assessments included random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, selective reporting and other items. Studies were graded as low, unclear or high risk of bias. Discrepancies were resolved via discussion among the researchers. Publication bias across the studies was presented as funnel plot.

**Data Analysis**

Meta-analysis was performed using RevMan 5.4. The effectiveness of blinding in sham/placebo acupuncture compared to real acupuncture was estimated with the odds ratios (OR) and its 95% confidence intervals. OR > 1 means blinding is more likely to occur in the intervention arm (sham group) than in the comparator arm (real group). Trials which the patients had no events in both intervention and comparator arms were excluded from the meta-analysis. Heterogeneities were assessed using the chi-squared ($\chi^2$) test and the inconsistency index ($I^2$) statistic. A two-tailed P value of less than 0.05 was considered as statistically significant. Subgroup analyses were performed on the factors which could influence the blinding effectiveness of participants.

**Results**

**Study Selection**

A total number of 1189 studies were identified from the databases and 18 studies from the reference list of related studies (Fig. 1). After screening, 34 studies had been included in the systematic review and meta-analysis. 28 studies had been excluded with reasons during the screening of full-text articles (Table 1).
Table 1
Excluded Studies with Reasons

| Excluded Studies | Reasons |
|------------------|---------|
| Chae 2006        | The study published in Korean |
| Dilli 2013       | Not a randomised controlled trial |
| Fink 2004        | No reported data on the awareness of the device used |
| Fink 2005        | No reported data on the awareness of the device used |
| Foster 2007      | No reported data on the awareness of the device used |
| Francia 2018     | Focused on the blinding of acupuncturist only |
| Grillo 2018      | No reported data on the awareness of the device used |
| Hu 2020          | Not a randomised controlled trial |
| Jiang 2014       | Inaccessible full-text article and no reply from the author |
| Lee 2006         | Inaccessible full-text article and no reply from the author |
| Leem 2016        | No reported data on the awareness of the device used |
| Lim 2006         | The study published in Korean |
| Park 2000        | The study published in Japanese |
| Park 2002        | Inaccessible full-text article and no reply from the author |
| Park 2008        | The study published in Korean |
| Park 2009        | Not a randomised controlled trial |
| Park 2010        | Inaccessible full-text article and no reply from the author |
| Streitberger 2003| No reported data on the awareness of the device used |
| Streitberger 2004| No reported data on the awareness of the device used |
| Takakura 2010    | Focused on the blinding of acupuncturist only |
| Takakura 2013    | Inaccessible full-text article and no reply from the author |
| Takakura 2014    | No reported data on the awareness of the device used |
| Tsukayama 2005   | Inaccessible full-text article and no reply from the author |
| Vickers 2005     | Insufficient data on the awareness of the device used |
| White 2000       | Insufficient data on the awareness of the device used |
| Wong 2018        | Insufficient data on the awareness of the device used |
| Yan 2016         | No reported data on the awareness of the device used |
| Zaslawski 1997   | Inaccessible full-text article and no reply from the author |

Study Characteristics

The characteristics of all 34 included studies had been summarised in Table 2. The publication year of recruited studies ranged from 1996 to 2019. All the studies performed participant blinding, but only some of them tried to blind acupuncturist or outcome assessor or statistician. The number of participants recruited in these studies ranged from 10 to 321. The total number of participants involved was 2538.
| Study ID       | Study Design                          | Participants | Needling        | Awareness of the Acupuncture |
|---------------|---------------------------------------|--------------|----------------|------------------------------|
| Chae 2010^44  | Participants                          | 14           | Healthy Experienced | Park LI4 11/14^              |
| Chen 2005^45  | Participants                          | 60           | Healthy Naive | Streitberger LI4 54/60^      |
| Deng 2007^46  | Participants                          | 72           | Not reported | Streitberger GV14, GB20, BL13, PC7, HT6, KI7, ST36, SP6, ear ShenMen & ear sympathetic point 18/42 |
| Dos Santos Maciel 2016^47 | Participants & outcome assessor | 321          | Healthy Naive | Park ST25 16/23 3/2          |
| Enblom 2008^48 | Participants                          | 80           | Not reported | Streitberger PC6 13/40 13/   |
| Enblom 2011^49 | Participants                          | 215          | Cancer Naive | Park PC6 50/63 8/6          |
| Fink 2001^50  | Participants & statistician            | 68           | Episodic/chronic tension-type headache Not reported | Foam GB20, LI4, LR3 & TE5 32/32 4/3 |
| Goddard 2005^51 | Participants & outcome assessor      | 49           | Healthy Naive | Foam LI4 19/24 3/2          |
| Kim 2015^52   | Participants & acupuncturist          | 63           | Healthy Mixed | Kim LI4 35/67^ 27/ |
| Kreiner 2010^10 | Participants & acupuncturist         | 32           | Healthy Naive | Foam LI4 27/32^ 24/ |
| Lao 1999^53   | Participants & outcome assessor      | 39           | Tooth extraction Mixed Not classifiable | LI4, ST6, ST7 & TE17 11/19 4/2 |
| Lee 2010^54   | Participants                          | 79           | Not reported | Park LI4 25/79 18/ |
| Liang 2013^55  | Participants & statistician           | 60           | Healthy Experienced | Park BL23 44/60^ 12/ |
| Liu 2014^56   | Participants, outcome assessor & statistician | 60          | Healthy Mixed | Foam LI4 59/60^ 0/6 |

^: mean; M: median; (): range; R: real group; S: sham group; a: reported gender ratio did not tally with the sample size; b: baseline characteristic for gender was missing (no response); c: drop-out gender was not reported; ^: based on the total responses from each participant.
| Study ID     | Study Design                                            | Participants | Needling | Awareness of the of Acupuncture |
|-------------|--------------------------------------------------------|--------------|----------|----------------------------------|
|             | Study Design                                                                                       |              |          |                                  |
|             | Blinding                                              | Sample Size  | Mean/Median Age (Range) in Years | Gender (M:F) | Health Status | Acupuncture Experience | Types of Sham Device | Location | Real | She |
| Park 2002   | Participants and outcome assessor                    | 58           | (38–87)   | 30:28  | Acute stroke | Naive          | Park  | LI4     | 57/60^   | 3/6   |
| Smith 2006  | Participants                                           | 228          | 35.9 ± 4.7 in R; 36.1 ± 4.8 in S | 0:228 | Infertility | Mixed          | Streitberger | SP4, ST29, CV3, BL32, SP8, SP6 & others based on the diagnosis | 40/202^   | 20/  |
| Smith 2011  | Participants                                           | 92           | 19.5 ± 2.9 in R; 18.9 ± 3.2 in S | 0:92  | Primary dysmenorrhoea | Mixed          | Streitberger | SP4, ST29, CV3, BL32, SP8, SP6 & others based on the diagnosis | 17/41     | 16/   |
| Streitberger | Participants                                           | 60           | 27.55     | 31:29  | Healthy     | Naive          | Streitberger | LI4     | 54/60^   | 13/    |
| Takakura 2007 | Participants & acupuncturists                        | 60           | 29.7 ± 7.5 | 35:25  | Healthy     | Experienced    | Takakura   | TE5     | 48/60^   | 35/    |
| Takakura 2008 | Participants & acupuncturist                          | 114          | 30.3 ± 7.9 | 73:41  | Healthy     | Experienced    | Takakura   | TE5     | 78/114^  | 50/    |
| Takakura 2011 | Participants, acupuncturist, outcome assessor         | 80           | 27.1 ± 6.9 | 48:32  | Healthy     | Experienced    | Takakura   | TE meridian on the posterior forearm | 65/80^   | 38/   |
| Takakura 2013 | Participants & acupuncturist                          | 109          | 28.6 ± 7.5 | 64:45  | Healthy     | Experienced    | Takakura   | TE5     | 85/109^  | 65/    |
| Takakura 2014 | Participants, acupuncturist & assistants              | 120          | 29.7 ± 9.3 | 60:60  | Functional neck/shoulder stiffness | Experienced    | Takakura   | SI14, SI15, GB21 & BL42 | 27/40     | 22/   |
| Tan 2009    | Participants                                           | 20           | 22 (18–48) | 6:14   | Healthy     | Naive          | Park       | PC meridian on the anterior forearm | 52/80^   | 44/   |
| Tan 2011    | Participants                                           | 20           | 24 (21–28) | 7:13   | Healthy     | Naive          | Park       | TE11, TE12, TE13 & TE14 | 24/40^   | 26/   |
| Tan 2019    | Participants                                           | 40           | 23 (21–40) | 13:27  | Healthy     | Not reported   | Park       | ST32 to ST39 | 108/160^ | 121   |
| To 2015     | Participants & acupuncturist                           | 5            | (23–54)   | Not reported | Healthy | Naive          | Park       | LI4, LI10, LI11, LI14, LI15 & TE14 | 7/16^     | 8/1   |
|             |                                                         | 11           | (22–74)   | Not reported | Healthy | Experienced    |            | 11/31^   | 17/    |
|             |                                                         | 19           | (22–74)   | Not reported | Should         | Not reported  |            | Above points + LV3, SI3, GB21, SI12 & ST38 | 8/8      | 0/6   |
| Tough 2009  | Participants                                           | 41           | Not reported | Not reported | Whiplash injury | Mixed       | Not classifiable | According to tender muscle points | 10/19     | 1/1   |
| Tsukayama 2006 | Participants                                        | 21           | 26 (19–68) | 15:6   | Healthy     | Experienced    | Park       | LI4     | 21/21^   | 12/    |
|             |                                                         | 20           | 24 (19–37) | 15:5   | Healthy     | Experienced    | Park       | BL23    | 14/20^   | 8/2    |

x̅: mean; M: median; (): range; R: real group; S: sham group; a: reported gender ratio did not tally with the sample size; b: baseline characteristic for gender was missing (no response); c: drop-out gender was not reported; ^: based on the total responses from each participant.
Meta-analysis was performed using RevMan 5.4.1. Data Analysis discussion among the researchers. Publication bias across the studies was presented as funnel plot. Incomplete outcome data, selective reporting and other items. Studies were graded as low, unclear or high risk of bias. Discrepancies were resolved via assessments included random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, and, hence, did not have constant points throughout the study. The effectiveness of blinding in sham/placebo acupuncture compared to real acupuncture was accepted “not sure” as one of the answers from the participants.

Quality Assessment

Quality and risk of bias of the eligible studies were assessed independently by two researchers at outcome level by using Cochrane risk of bias tool14. Quality assessments included random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, selective reporting and other items. Studies were graded as low, unclear or high risk of bias. Discrepancies were resolved via discussion among the researchers. Publication bias across the studies was presented as funnel plot.

Data Analysis

Meta-analysis was performed using RevMan 5.4.1. The effectiveness of blinding in sham/placebo acupuncture compared to real acupuncture was estimated with the odds ratios (OR) and its 95% confidence intervals. OR > 1 means blinding is more likely to occur in the intervention arm (sham group) than

| Study ID | Study Design | Participants | Needling | Awareness of the Type of Acupuncture |
|----------|--------------|--------------|----------|--------------------------------------|
| Vase 2015 | Participants & acupuncturists | 67 | x̅: 25.8 ± 5.0 | ST44, LI4, ST7, ST6 & TE17 | 22/32 |
| White 1996 | Participants & outcome assessor | 10 | x̅: 57.2 ± 13.6 R; 57.4 ± 19.9 S | L14 with local points like GB14, GB20, GB21, EX-HN3 YinTang & EX-HN5 TaiYang | 4/4 |
| White 2003 | Participants | 37 | x̅: 65.8 ± 8.3 R & 64.4 ± 12.7 (37–79) | GB30, GB31, GB34, BL34, BL39, BL40, BL60, ST31, ST35, ST36, ST41, SP9 & EX-LE5 XiYan | 25/37 |
| White 2007 | Participants | 20 | x̅: 50.8 ± 14.9 (35–80) | LI4 | 16/20 |
| Xie 2013 | Participants | 60 | x̅: 23.07 ± 0.753 R & 23.13 ± 1.056 in S (22–25) | BL23 | 45/60 |

All recruited subjects were adults. The mean age ranged from 18.9 to 65.8 years old. In some of the studies, diseased subjects were recruited such as cancer, tooth extraction, stroke, infertility, dysmenorrheoe, infertility, shoulder impingement syndrome, whiplash injury and joint pain/osteoarthritis, whereas the rest of them were either healthy or not reported. Most studies reported acupuncture experience of the subjects, for example, naive (never experienced acupuncture before), experienced (at least experienced acupuncture once before) or multiple times of evaluation. Most studies recorded their results in Yes-No format, whereas the 15 of them accepted “not sure” as one of the answers from the participants.

A variety of non-penetrating sham acupuncture devices was used in the studies. They were classified and named according to the name of the authors (Park, Streitberger, Takakura and Kim) or materials (foam and cocktail stick). The studies had chosen acupoints or non-acupoints located on the head, abdomen, back, upper limbs, lower limbs and ears. The coding and naming of acupoints were converted if they did not follow a Proposed Standard International Acupuncture Nomenclature by the World Health Organization (WHO). Some selected acupoints based on syndrome differentiation of Chinese medicine or tender muscle point and, hence, did not have constant points throughout the study.

The total number of responses from the participants due to cross-over study design, using a mixture of real and sham acupuncture devices or multiple times of evaluation. These responses were extracted and analysed because the blinding effectiveness of a particular sham device ideally should be maintained even after several times of evaluation. Most studies recorded their results in Yes-No format, whereas the 15 of them accepted “not sure” as one of the answers from the participants.
in the comparator arm (real group). Trials which the patients had no events in both intervention and comparator arms were excluded from the meta-analysis. Heterogeneities were assessed using the chi-squared ($\chi^2$) test and the inconsistency index ($I^2$) statistic. A two-tailed $P$ value of less than 0.05 was considered as statistically significant. Subgroup analyses were performed on the factors which could influence the blinding effectiveness of participants.

**Blinding Effectiveness of Sham Acupuncture Devices**

The OR of overall blinding effectiveness of non-penetrating sham acupuncture devices (Fig. 5) was 5.11 [3.36, 7.76] with a $P$-value ($< 0.00001$), which indicated that the blinding was more likely to occur in the sham group and the result was statistically significant. However, the studies were not homogenous and the heterogeneity was high in the meta-analysis including the subgroup analyses, hence, a random effect model was selected. After switching to the random effect model, the heterogeneity was still high ($I^2 = 88\%$) in the overall blinding effectiveness analysis, so subgroup analyses were performed to evaluate the factors which may influence the blinding effectiveness of sham acupuncture devices. For example, acupuncture experience of the participants, the health status of the participants and the location of acupoints.

A comparison between sham acupuncture devices was performed through evaluating and comparing the individual blinding effectiveness of sham acupuncture devices. Kim and cocktail stick devices were excluded because they only possessed one study each. As shown in Fig. 6, the foam device was the most successful device in blinding the participants because it has the highest OR (44.78) [10.03, 199.92], followed by Streitberger (4.69) [1.87, 11.78], Park (3.16) [1.61, 6.21], Takakura (2.66) [1.98, 3.55] and no-touch Takakura (1.47) [0.82, 2.62]. The $P$-value was statistically significant ($< 0.05$) in all devices except for the no-touch Takakura device. The heterogeneity was high in Park, Streitberger and foam but low in Takakura and no-touch Takakura. The heterogeneity across the subgroups was high ($I^2 = 79\%$).

**Blinding Effectiveness in Different Types of Participants**

Naive participants demonstrated superiority in achieving blinding during acupuncture clinical trials with an OR 5.73 [2.76, 11.89] compared to experienced participants 3.22 [1.94, 5.37] (Fig. 7). Both naive and experienced participants were successful in blinding because they were both statistically significant ($P < 0.00001$). The heterogeneity was high in both groups individually but did not differ much between the two subgroups ($I^2 = 37.7\%$).

As for the health status of participants, both healthy and diseased participants were statistically significantly successful in blinding ($P < 0.00001$) using sham acupuncture devices (Fig. 8). Healthy participants were blinded slightly better than diseased participants with an OR 6.28 [3.62, 10.91] compared to 5.79 [2.71, 12.35]. The heterogeneity was high within the subgroups but low when comparing the two subgroups ($I^2 = 0\%$).

**Blinding Effectiveness in Different Locations of Acupoints**

The acupoints of the included studies had been classified according to its regions, for example, head and neck, chest, abdomen, back, upper limbs and lower limbs. Studies which utilised acupoints from multiple parts of the body were excluded. Besides, there was only one study which had used acupoints purely from the head and neck region, therefore it was excluded from the meta-analysis. Among the remaining regions, the abdomen demonstrated a better blinding effect with an OR 11.33 [0.71, 180.77], followed by the back (9.10) [3.14, 26.36], the upper limbs (4.31) [2.50, 7.43] and the lower limbs (3.69) [0.82, 16.62] (Fig. 9). The back and the upper limbs exhibited a statistically significant result ($P < 0.00001$), whereas the results of the abdomen and the lower limbs were not statistically significantly different ($P = 0.09$). However, heterogeneity was high in all of the individual subgroups but low across the subgroups ($I^2 = 0\%$).

**Discussion**

**Blinding Effectiveness of Sham Acupuncture Devices**

**Overall Blinding Effectiveness of Sham Acupuncture Devices**

When conducting acupuncture clinical trials, subjects from both real acupuncture group and sham acupuncture group should believe that they receive real acupuncture to mimic the actual scenario which takes place during acupuncture treatment. As shown in Fig. 5, the existing non-penetrating sham acupuncture devices such as Park, Streitberger, Takakura, foam and cocktail stick displayed a result which favoured the sham group and was also statistically significant. In other words, more participants in the sham acupuncture group who did not identify the sham acupuncture treatment correctly compared to the real group and hence, they were blinded. As a result, non-penetrating sham acupuncture devices can act as an effective placebo control method to be applied in acupuncture clinical trials, especially in replacing other less effective types of control methods such as using no treatment, standard/conventional treatment or skin-penetrating sham acupuncture as the control.

**Comparison between Sham Acupuncture Devices**

Among all the non-penetrating sham acupuncture devices included in the data analysis, the foam device demonstrated the best ability in achieving the blinding of participants. There were five studies that used the foam device in their control group. The foam device is usually self-prepared by the researchers, so there are some variations in design. It is made out of a certain thickness of foam with double-sided adhesive tape at the bottom (Fig. 10). The foam pad can act as a supportive material to hold the needle in place even in the placebo group, whereas the adhesive tape can stick the device on the skin. The real foam device utilises a real acupuncture needle with a sharp tip which can penetrate the skin of participants; the placebo foam device uses a shorter blunted acupuncture needle which cannot penetrate the skin. Ultimately, the appearance after needle insertion will remain the same and hence, achieve blinding of participants in terms of vision. After needle insertion, the real device will penetrate the skin with a certain depth, whereas the placebo device will only touch the skin to blind the participants by mimicking the feeling of penetration. In the studies of Fink and Goddard, the placebo needles were gently twisted to enhance the feeling of penetration. Besides having a good blinding effect, the foam device is also less pricey and easily accessible compared to the
other devices, so it can be a good option to be used in the control group of acupuncture clinical trials. However, owing to the preparation of the foam device is usually self-made, hygiene in preparing the device will become the main concern. All the equipment must be sterilised adequately before being applied to the participants.

The other sham acupuncture devices such as Streitberger, Park and Takakura (Fig. 11) are also effective in blinding the participants. Their sham devices look identical with the real devices. Similar to the foam device, their sham devices possess shorter blunted-tip needles which touch on the skin to mimic penetrating sensation. Hence, all of them can be applied in clinical trials. The characteristics of the foam, Streitberger, Park and Takakura devices have been summarised in Table 3. On the other hand, the sham device of no-touch Takakura was not statistically significantly superior to the real device (P = 0.19), so it is not recommended to be used in clinical trials. No-touch sham Takakura device has no contact with the skin of participants. It may be useful in achieving visual blinding but not tactile blinding.

In short, each sham acupuncture design has its own advantages and limitations. Researchers should take that into consideration when designing the experimental and control methods. Due to high heterogeneity of the studies across the subgroups (Fig. 6), the OR value may be influenced by other factors and hence, the results of blinding effectiveness of sham devices can only serve as a reference.

**Limitations of the Existing Sham Acupuncture Devices**

The limitations of the existing sham acupuncture devices can be discussed in several aspects, including the suitability in performing electro-acupuncture, needling location, needling angle and blinding effect in acupuncturists.

The sham devices of foam, Streitberger, Park and Takakura solely rely on the adhesive double-sided tape or pedestal to attach to the skin. The attachment is not as firmly as those in the real acupuncture, therefore, sham electro-acupuncture is difficult to perform by using these devices. Moreover, the adhesive tape may not be suitable for hairy skin or areas which are not flat. As a result, using sham acupuncture devices on the scalp, hairy regions and the skin with a great curvature such as the ears, fingers and toes can be challenging.

Apart from that, needling angle is usually limited to perpendicular for the foam, Park and Takakura devices due to the presence of guide tube, so only acupoints that allow perpendicular insertion can be selected when using these devices. Acupoints such as LU7 LieQue and EX-HN3 YinTang or acupoints on the scalp which require oblique or transverse insertion cannot be chosen in the trials. On the contrary, Streitberger device, which does not have a guide tube, can perform perpendicular and oblique insertions. Yet, having no guide tube can also make the needle unstable especially in the sham group, increasing the risk of exposure of grouping allocation.

Last but not least, most of the sham acupuncture devices did not demonstrate the ability to blind acupuncturists, except for the Takakura device. The Takakura device has added a lower stuffing within its guide tube to mimic the feeling of skin penetration when the acupuncturist pushes the sham needle into the lower stuffing. In other words, the performance bias will be high in the acupuncture clinical trials that use other types of sham acupuncture devices.

**Considerations of Designing Sham Acupuncture Controlled Clinical Trials**

**Guidelines of Sham Acupuncture Controlled Clinical Trials**

In 1995, the Guidelines for Clinical Research on Acupuncture by the WHO stated that placebo acupuncture should fulfil two criteria: it must be a less effective form of acupuncture and also mimic acupuncture in a credible manner. Zhang’s paper also stated that placebo acupuncture should have no or minimal specific treatment effects on the tested disease and the treatment and control groups should be identical to achieve blinding. To evaluate the true efficacy of acupuncture, the difference in specific effects shall be maximised but the difference in non-specific effects shall be minimised (Fig. 12).

| Devices  | Characteristics                                                                 | Price $/per needle | Advantages                                      | Limitations                                      |
|----------|----------------------------------------------------------------------------------|--------------------|-------------------------------------------------|-------------------------------------------------|
| Foam     | Needle is supported by an opaque guide tube which is attached to the foam base;  | N/A                | Inexpensive, easily accessible and self-prepared | Potential risk of hygienic issue and inconsistency of quality of the foam |
|          | adhesive tape at the bottom                                                      |                    |                                                 |                                                 |
| Streitberger | Sham: retractable needle supported by plastic ring covered with plastic sheet as the base; adhesive tape at the bottom | $6.3               | Can insert in different angles                  | Sham needle attachment is not firm               |
| Park     | Needle is supported by an opaque guide tube and a Park tube which is connected with the ring base; adhesive tape at the bottom | $2.9               | Able to perform simple needle manipulations     | Can only insert perpendicularly                 |
| Takakura | Needle is supported by an opaque guide tube which is filled up with stuffings at upper and lower ends and the tube is connected to an adhesive pedestal | N/A                | Potentially blind the acupuncturist             | Can only insert perpendicularly                 |

N/A: not available

Notes: price was cited from the other article. Current price may differ from the past time.
Determining a research question is important before selecting the type of control because each control method can be used to answer different types of research questions. As shown in Table 4, no treatment and standard treatment as control can rule out regression to the mean and study the general effectiveness of acupuncture; non-penetrating sham as control can rule out regression to the mean and psychological responses (placebo effect) and study the efficacy of acupuncture, including skin-penetrating physiological effects and acupoint specific effects; lastly, penetrating sham as control can rule out three other aspects and study specifically on the efficacy of acupoint specific effects. As far as the authors know, non-penetrating sham acupuncture is the only method that can eliminate the placebo effect and meanwhile, minimise the physiological responses being generated. So, it can be widely used in a broad range of acupuncture clinical trials that study the specific effects of acupuncture. On the other hand, penetrating sham acupuncture (e.g. shallow needling/minimal acupuncture and needling on non-acupoints) is suitable to study narrower specific effects of acupuncture that will not be generated by skin penetration. Do note that the acupuncturists of penetrating sham acupuncture are not blinded most of the time owing to different needling techniques and locations in the sham group and hence, it may lead to performance bias of personnel. In addition, it has to be ensured that skin penetration will not trigger any desired specific effects of the study, if not it will result in no significant difference in both arms.

Table 4

| Types of Control          | Aspects                  | Regression to the Mean | Psychological Responses | Physiological Responses* | Acupoint Specific Effects |
|---------------------------|--------------------------|------------------------|-------------------------|--------------------------|--------------------------|
| No treatment              | X                        | X                      | Efficacy                | Efficacy                 | Efficacy                 |
| Standard treatment        | X                        | X                      | Efficacy                | Efficacy                 | Efficacy                 |
| Non-penetrating sham      | X                        | X                      | Efficacy                | Efficacy                 | Efficacy                 |
| Penetrating sham          | X                        | X                      | X                       | Efficacy                 |                         |

Note: the boxes with "X" in Table 4 indicate the areas that are eliminated when comparing with the real acupuncture group.

Factors which May Influence the Blinding Effectiveness of Sham Acupuncture Devices

 Besides the selection of sham acupuncture method and device, there are other factors which may also play a role in achieving successful blinding, for example, acupuncture experience and health status of the participants and needling location. As shown in Fig. 7, the naive participants are more easily blinded than the experienced participants. The experienced participants are more familiar with the acupuncture process and DeQi (needling) sensation and hence, they are more likely to guess the grouping accurately. However, both groups showed a significant difference in blinding effectiveness. Ideally, naive subjects should be recruited in acupuncture clinical trials, but experienced participants can also be considered if naive ones are not available or sufficient.

Next, both healthy and diseased participants demonstrated a significant difference in blinding effectiveness, in which the healthy ones were slightly superior to the other group (Fig. 8). Healthy subjects should be prioritised when designing acupuncture clinical trials, nonetheless, recruiting diseased participants is unavoidable when studying the efficacy of acupuncture on a particular disease.

Lastly, the locations of acupoints may also influence the effectiveness of blinding. As shown in Fig. 9, the OR values of the abdomen (11.33) [0.71, 180.77] and the back (9.1) [3.14, 26.36] were higher compared to the upper limbs (4.31) [2.50, 7.43] and the lower limbs (3.69) [0.82, 16.62]. This phenomenon is associated with the higher sensitivity of the skin and stronger needling sensations in the four limbs due to rich nervous distribution. Yet, only the back and upper limbs showed statistically significant different P-values.

Limitations of the Study

Four studies had been excluded due to language barrier and seven studies had been excluded due to inaccessibility of the full-text articles. Excluding articles other than English and Chinese may introduce language bias. Excluding inaccessible articles may also reduce the precision of combined estimates of blinding effectiveness. However, the authors were not able to overcome it due to limited resources. The included studies utilised different ways to present data, so some data had to be converted before performing the meta-analysis. Also, some studies presented their results in terms of the number of participants, whereas some were based on the total responses from the participants.

A variety of study designs also led to high heterogeneity in the results of the meta-analysis. Other potential influencing factors which might contribute to the heterogeneity such as the diameter of acupuncture needle, the depth of needle insertion, the duration of needle retention, needle manipulation techniques and the number of treatments were not analysed in the study. For instance, the participants may be aware of the grouping by observing the procedure and treatment effects after having multiple and long-time treatment.

The number of articles in some of the subgroups was small. For example, Kim device and cocktail stick device in the analysis of the types of sham acupuncture devices possessed solely one study; besides, the head and neck, abdomen and lower limbs in the analysis of the locations of acupoints possessed a small number of studies.

Conclusions
Non-penetrating sham acupuncture devices are valid placebo control for acupuncture clinical trials. It should be applied in future clinical trials because it can blind the participants and meanwhile, produces lesser physiological responses. The foam device has a better blinding effect, followed by the Streitberger, Park and Takakura devices. Sham needles with no skin contact could not successfully blind the participants. Naive, experienced, healthy and diseased participants all can be used in acupuncture clinical trials but naive and healthy participants can be blinded more easily. Choosing acupoints from the back can blind participants more easily compared to the other areas.

Future clinical trials can study and compare the blinding effectiveness of sham acupuncture devices by performing multiple sham groups with a larger sample size to evaluate all different types of sham devices under the same setting. This can eliminate heterogeneity and show real blinding efficacy of these devices. Besides, researchers can also try to develop a better type of non-penetrating sham acupuncture device with the considerations of the blinding efficacy, strengths and limitations as mentioned in the results and discussion.

**Abbreviations**

PRISMA: Preferred Reporting Items for Systematic Reviews and Meta-Analyses; RCT: Randomised Controlled Trial; OR: Odd Ratio; WHO: World Health Organization

**Declarations**

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**Authors’ Contributions**

GYH provided the idea of the study. SNA provided the study design. GYH conducted the literature search from the databases. GYH, SNA and SYS performed article screening, data extraction and quality assessment. GYH and SNA performed meta-analysis. GYH wrote the manuscript. All authors approved the final submission.

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**Availability of data and materials**

All data generated or analysed during this study are included in this published article.

**Ethics approval and consent to participate**

Not applicable.

**Consent for publication**

All authors agree for publication in BMC Complementary Medicine and Therapies.

**Competing interests**

All authors declare that there are no conflicts of interest.

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