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Title: A feasibility study of auricular therapy and self-administered acupressure for insomnia following cancer treatment

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Abstract

Introduction: Many cancer patients experience sleeping difficulties which can persist several years after the completion of cancer treatment. Previous research suggests that acupuncture, and variants of acupuncture (acupressure, auricular therapy) may be effective treatment options for sleep disturbance. However, current evidence is limited for cancer patients.

Methods: Feasibility study with 3 arms. Seven cancer patients with insomnia randomised to receive either auricular therapy (attaching semen vaccariae seeds to ear acupoints) (n=4), self-acupressure (n=1) or no treatment (n=2). Participants assigned to receive auricular therapy or self-acupressure stimulated the acupoints each night an hour before retiring to bed. The duration of participant involvement was 5 weeks. Subjective sleep quality was measured at baseline and post-treatment using the Pittsburgh Sleep Quality Index (PSQI). The impact of treatment on concerns of importance to the participants themselves was measured using the Measure Yourself Concerns and Wellbeing (MYCaW). Each participant also completed a treatment log book.
Results: All participants completed their treatment. All auricular therapy and self-acupressure participants recorded clinically significant improvements in global PSQI scores. In the auricular therapy arm mean global PSQI reduced from 12.5 at baseline to 8 following completion of treatment. In the self-acupressure arm PSQI reduced from 15 to 11. While in the no treatment arm the mean PSQI score was 14.5 at both baseline and follow up.

Conclusions: Despite the limited sample size, both auricular therapy and self-acupressure may represent potentially effective treatments for cancer patients with insomnia. The positive findings suggest further research is warranted into both treatment modalities.

Keywords:
Cancer, insomnia, auricular therapy, acupressure.

Introduction:
Between 25% and 59% of cancer patients experience sleeping difficulties [1-4]. Patients ranked sleep problems as the 5th highest out of 14 distressing symptoms before their cancer treatment, and 4th highest after cancer treatment [5]. In between 23% and 44% of cancer patients, sleep problems persist several years after the initiation of adjuvant therapy for cancer, suggesting that insomnia develops a chronic course in substantial numbers of cancer patients [6]. Many of these cancer patients receive hypnotic medications to relieve their symptoms of insomnia. However, the usage of hypnotic medications is associated with a number of risks and limitations, such as daytime drowsiness and associated increased risk of falls and fractures [6]. Prolonged usage is also associated with risks of dependence or tolerance (reduced efficacy due to prolonged usage, and the need to increase the dosage to maintain therapeutic effects) [6]. Given the limitations of hypnotic medications non-pharmacologic approaches to treating insomnia should be considered.

The findings from published systematic reviews suggest that acupuncture, and variants of acupuncture (acupressure; auricular acupuncture; auricular therapy), may be effective treatment options for sleep disturbance [7-13]. However, evidence of the effectiveness of acupuncture and related interventions within cancer are more limited [14]. A non-randomised single arm feasibility study evaluated a short course of acupuncture treatment for sleep disturbance and hot flushes in 10 postmenopausal breast cancer survivors. The findings suggested a short term benefit from acupuncture on a range of sleep outcomes (total sleep time, sleep latency, and night-time awakenings) [15]. A randomized controlled trial comparing acupuncture with fluoxetine in 80 cancer patients presenting with depression and insomnia concluded that acupuncture significantly improved sleep quality [16].

However many previous studies of acupuncture and related interventions have contained numerous methodological weaknesses [7-16], and there is no conclusive evidence to suggest which modality of acupuncture may be more effective for sleep disturbance. This feasibility study aimed to address this gap in the evidence base and provide preliminary data on the effectiveness of two variants of acupuncture (auricular therapy (attaching semen vaccariae seeds to ear acupoints) and self-administered acupressure) in the relief of insomnia in a sample of patients with cancer in a methodologically rigorous feasibility study.

Methods:
The study was a randomised controlled feasibility study with 3 arms. Treatment arms consisted of (1) auricular therapy (attaching semen vaccariae seeds to ear acupoints), (2) self-administered acupressure, and (3) no additional treatment. Patients were informed about the study by treating healthcare professionals at the study site. Members of the research team provided interested patients with a patient information sheet and detailed verbal information regarding the study. Participants were required to be hospital outpatients with breast, prostate or colorectal cancer; meet the criteria for chronic insomnia syndrome (require more than thirty minutes to fall asleep and/or have more than thirty minutes of nocturnal awakenings, occurring three nights a week or more, for six months or more and affecting daytime functioning); have a score of five or more on the Pittsburgh Sleep Quality Index; be older than 16 years; and receiving acupressure or auricular therapy for the first time. Exclusion criteria included those with self-reported sleeping difficulties prior to receiving their diagnosis of cancer; those currently receiving, or having received within the last month, chemotherapy or radiotherapy; and those scheduled to receive anticancer treatments within the next three months, with the exception of adjuvant hormone therapy. Patient recruitment occurred between February 2010 and June 2012.

The duration of participants' involvement was 5 weeks. Participants were randomly allocated to the trial groups through simple envelope method. Trial allocations were randomly placed in sealed numbered opaque envelopes by a member of university staff not involved in the research study or the care of the participating patients. Upon participants consenting to take part in the study the next numbered envelope was opened to reveal the trial allocation. As part of the feasibility study aspects of the study design, such as recruitment, acceptability of the intervention, and appropriateness of the outcome measures were evaluated. The study was conducted at the Christie NHS Foundation Trust, Manchester. Ethical approval for the feasibility study was obtained from Stafford and Trafford Research Ethics Committee [REC reference number: 09/H1004/7].

Interventions:
Participants were randomly allocated to receive either auricular therapy, self-administered acupressure, or no treatment. The modalities of acupressure and auricular therapy were chosen for this study due to considerations of ease of administration, and because they are less invasive than needle insertion. The acupuncturists instructing patients and administering treatments were all nurses who had received training from either the British Medical Acupuncture Society or the British Academy of Western Medical Acupuncture. All acupuncturists had at least 3 years of clinical experience in treating patients with cancer.

Auricular therapy: participants assigned to auricular therapy attended Christie NHS Foundation Trust on a weekly basis for five weeks. Appointment times lasted approximately ten minutes and took place at a time convenient for participants (total therapist contact time 50min). Acupoints selected for the feasibility study were based on those utilised in previous research; acupuncture textbooks; and consultations with practicing acupuncturists. The auricular acupoints stimulated included shenmen bilaterally, and any two from insomnia 1, insomnia 2, heart, liver, kidney and subcortex bilaterally based on the patient's presenting condition. Semen vaccariae seeds were fixed tightly to each acupoint with a piece of adhesive plaster by the treating acupuncturist. The seeds were left in place and replaced each week by the acupuncturist. Based on previous research and consultations with practicing acupuncturists, participants were asked to press the seeds for one minute each night an hour before retiring to bed.

Self-administered acupressure: Participants allocated to the self-administered acupressure group received a 50 min appointment with an acupuncturist, at a time convenient for the participant. The
acupuncturist provided participants with instructions on how to locate and apply pressure to specific acupoints. Acupoints selected for the feasibility study were based on points used within previous research; consulting acupuncture/acupressure textbooks; and consultations with practicing acupuncturists. Consideration was additionally given to the likely ease of location and stimulation by patients. All patients were instructed to apply pressure to HT7 bilaterally, and two from PC6, GB20, Ex8 (anmian 1), KL6, BL62 bilaterally based on the patients presenting condition. Based on previous research and consultations with practicing acupuncturists, participants were asked to stimulate each acupoint for approximately one minute, each evening an hour before retiring to bed.

No additional treatment: Participants allocated to the no treatment group received no additional treatments for the duration of the study. Participants allocated to no additional treatment were offered the choice of receiving either auricular therapy or self-acupressure at the end of their participation in the study.

Outcome measures:
Subjective sleep quality was measured using the Pittsburgh Sleep Quality Index (PSQI) [17], which contains seven indices to represent patients’ quality of sleep over the previous month: overall sleep quality; sleep latency (the amount of time it takes to fall asleep); sleep duration; sleep efficiency; sleep disturbances; day dysfunction due to sleepiness; and use of sleeping medication. Global PSQI scores range from 0 to 21, with a score of ≥5 indicating sleeping difficulty. In addition the impact of treatment on concerns of importance to the participants' themselves were measured using the Measure Yourself Concerns and Wellbeing (MYCaW) [18]. The MYCaW is an individualised questionnaire that has been developed for evaluating complementary therapies in cancer care. MYCaW contains two patient-generated problems. Patients rate two concerns or problems which they would most like help with using a seven point likert scale, with 0 being ‘not bothering me at all’ and 6 being ‘bothers me greatly’. MYCAW also contains an overall measurement of wellbeing, also rated on a seven point likert scale with 0 being ‘as good as it could be’ and 6 being ‘as bad as it could be’.

Participants completed the PSQI and MYCaW with a member of the research team at baseline, and self-completed the questionnaires five weeks later at the end of the study intervention. Participants also completed a log book documenting their use and experience of conventional and unconventional treatments for sleep disturbance during the study (including auricular therapy and self-administered acupressure if assigned to one of these groups).

Population sample:
The planned sample for the study was a stratified sample of 18 hospital outpatients (6 breast cancer patients, 6 prostate cancer patients, 6 colorectal cancer patients). However, due to recruitment difficulties only 7 participants were recruited. 4 participants were randomised to receive auricular therapy, 1 self-acupressure, and 2 no additional treatment. Five participants were diagnosed with breast cancer, and 2 with colorectal cancer. The age range of the participants was 43 to 71 (mean 55), 6 were female and 1 male. See Table 1 for sociodemographic characteristics of participants.

RESULTS:
Data from the log books indicate that participants allocated to auricular therapy and self-acupressure largely administered the treatments as instructed on a nightly basis. One auricular therapy participant reported forgetting to apply the intervention on three evenings, whilst another
auricular therapy participant reported purposely not pressing the ear seeds on three separate evenings, to test if there was continuation of the improvement in sleep that had been experienced.

One participant allocated to auricular therapy contracted shingles during the study and received painkillers, antiviral medication and antidepressants. No other pharmacological interventions were taken by participants during the study. Apart from the trial interventions, the only non pharmacological methods that participants reported using to help with sleep were warm milky drinks and bubble baths. Two participants reported some ear tenderness from the auricular therapy, although this did not appear to cause distress or be long lasting. Two of the participants described individual seeds falling off, and both of them reported pressing the site where the seed had been instead until the seed was replaced. Participants receiving active treatments commented in their log books that they had positive perceptions of the experience of undertaking the therapy and the effect the treatment had on their sleep.

It had been intended that data from the PSQI and MYCaW would be inputted into Statistical Packages for Social Sciences (SPSS) and analysed using appropriate between-group tests. However, due to the small sample size achieved only the mean improvement in scores is presented. Mean global PSQI score improved by 4.5 in the auricular therapy group, by 4 in the self-acupressure group, and did not change in the no additional treatment group. For the auricular therapy group improvements in PSQI scores were seen in duration of sleep, day dysfunction due to sleepiness, sleep efficacy, and overall sleep quality. In the self-acupressure participant improvements were seen in sleep latency, sleep disturbance, and overall sleep quality.

All participants listed an improvement to their sleep as their primary concern when completing the MYCaW questionnaire. Mean MYCaW score for sleep reduced by 2.25 in the auricular therapy arm, by 2 in the self-acupressure arm, and increased by 0.5 in the no additional treatment arm. Participants’ second item of concern included anxiety, fatigue, and pain. In both the auricular therapy arm and the self-acupressure arm the mean reduction in the second item of concern to participants reduced by 1, while the score for the no additional treatment arm remained the same. See Table 2 for all mean baseline and post-treatment scores.

INSERT TABLE 2 HERE

Discussion:
To the authors knowledge the present feasibility study of auricular therapy and self-acupressure is one of the first studies to evaluate these modalities for insomnia in cancer patients. All participants allocated to auricular therapy or self-acupressure completed the intervention, with only minor issues with ear tenderness reported by two auricular therapy participants, suggesting that both interventions are acceptable to cancer patients. The PSQI and MYCAW appeared to be appropriate tools to use and were completed by participants with no concerns raised. The tools were sensitive to reported changes in sleeping patterns and the scores of the two tools corresponded, ie, changes or similar scores in the PSQI were reflected in the scores of the MYCAW. Recruitment rates for the study were lower than anticipated, and are a major limitation of the current study. The key problem was that, despite current evidence suggesting that between 23% and 44% of cancer patients develop symptoms of chronic insomnia following treatment for cancer [6], the present study found very low rates of cancer patients reporting insomnia to their treating healthcare practitioner. However, there is evidence that people often do not report their experience of insomnia to their health professionals [22], which may have further hindered recruitment in this study. A large number of those patients who did present with insomnia were also ineligible to participate, due to currently receiving, or having received in the previous month, chemotherapy.
and/or radiotherapy. It may be advisable for future studies to either expand inclusion criteria to include those receiving chemotherapy and/or radiotherapy, or target post active treatment patients through alternative sites, such as GP practices.

A reduction of 3 points on global PSQI is considered a clinically significant reduction [17,19,20], and it is noteworthy that all 5 participants allocated to receive either auricular therapy or self-acupressure had a reduction of 3 or more in global PSQI scores. Although the sample size was very small, it may also be noteworthy that those who received auricular therapy reported different improvements in insomnia symptoms to the participant who received self-acupressure. While participants in both groups saw an improvement in overall sleep quality, the four participants who received auricular therapy noted improvements in the duration of sleep, day dysfunction due to sleepiness, and sleep efficacy; whereas the one self-acupressure participant noted a reduction in sleep latency and sleep disturbance, which may be an interesting hypothesis to explore in future research.

The clinical improvement seen in patients receiving auricular therapy or self-acupressure is comparable to that seen in previous studies of acupuncture and related interventions in patients with different presenting conditions. In the Cochrane systematic review by Cheuk et al [8] the findings from 5 randomised controlled trials of acupressure versus no additional treatment which had utilised the PSQI were collectively analysed. The 5 trials included a diverse range of patient populations including those with depression/anxiety, renal disease, and haemodialysis patients. Their findings showed that the global PSQI score was better in the acupressure group compared to the no treatment group (P < 0.00001), with the mean reduction of PSQI score being 3.17 greater in the acupressure group compared to the no additional treatment group. Similar improvements have been reported for trials of auricular therapy for insomnia. Wang et al [21] reported a mean greater improvement of 2.5 in global PSQI score compared to no treatment in nursing students with an initial PSQI score of 5 or more. However, all the trials contained methodological weaknesses, with Cheuk et al [8] noting that the trials contained within their review had a high risk of bias and were heterogeneous in their definition of insomnia, participant characteristics, acupoints and treatment. Despite the very limited sample size, all patients allocated to receive either auricular therapy or self-acupressure experienced clinically relevant improvements in their symptoms of insomnia. Both auricular therapy and self-acupressure are non-invasive interventions, which appear safe and acceptable to cancer patients, and may represent effective treatments for chronic insomnia syndrome. Further large scale trials of both interventions are warranted to establish the effectiveness of auricular therapy and self-acupressure for insomnia in cancer patients.

Conflict of interest:
The authors declare they have no competing interests.

Acknowledgements:
The authors wish to thank the acupuncturists at Christie NHS Foundation Trust for administering treatments during the study, the patients who kindly participated in the study, and the Cancer Experiences Collaborative (CECo) for funding the research.

References


Table 1 Sociodemographic characteristics of participants

<table>
<thead>
<tr>
<th>Participant</th>
<th>Gender</th>
<th>Age</th>
<th>Cancer diagnosis</th>
<th>Ethnic origin</th>
<th>Marital Status</th>
<th>Educational Level</th>
<th>Occupational group</th>
<th>Occupational status</th>
<th>Intervention</th>
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<td>Female</td>
<td>45</td>
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<td>Caucasian</td>
<td>Married</td>
<td>College/ diploma</td>
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<td>71</td>
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<td>Self-acupressure</td>
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<td>Married</td>
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<td>Skilled manual</td>
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<td>Data not available</td>
<td>Data not available</td>
<td>Auricular therapy</td>
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Table 2 Mean baseline and post-treatment scores

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<th>Mean (range)</th>
<th>Change</th>
<th>Mean (range)</th>
<th>Change</th>
<th>Mean (range)</th>
<th>Change</th>
</tr>
</thead>
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<td>End of treatment</td>
<td>Baseline</td>
<td>End of treatment</td>
<td>Baseline</td>
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<td></td>
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<td>End of treatment</td>
<td>Baseline</td>
<td>End of treatment</td>
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<td>Baseline</td>
<td>End of treatment</td>
<td>Baseline</td>
<td>End of treatment</td>
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<tr>
<td>Baseline</td>
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<td>8 (4-12)</td>
<td>4 (4-6)</td>
<td>5 (4-6)</td>
<td>14.5 (13-16)</td>
<td>4 (4-6)</td>
</tr>
<tr>
<td>Baseline</td>
<td>5 (4-6)</td>
<td>2.75 (1-4)</td>
<td>-2.25</td>
<td>6</td>
<td>-4</td>
<td>4.5 (3-6)</td>
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<tr>
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<td>4.5 (4-5)</td>
<td>-1</td>
<td>3</td>
<td>2</td>
<td>5 (4-6)</td>
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<tr>
<td>Baseline</td>
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<td>2.75 (1-4)</td>
<td>-0.25</td>
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<td>3</td>
<td>2.5 (2-3)</td>
</tr>
<tr>
<td>Baseline</td>
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<td>-1.67</td>
<td>4.33</td>
<td>3</td>
<td>4.17</td>
</tr>
</tbody>
</table>

*MYCAW profile scores are the mean of the concerns and wellbeing scores combined.*