

# The Effectiveness of Acupuncture for Insomnia

<b>Submission date</b> 18/08/2007	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 03/09/2007	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 03/09/2007	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

096XDAA00028

## Study information

Scientific Title

**Acronym**

EAI

**Study objectives**

Traditional Chinese Medicine (TCM) Meridian Theory.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Approved by the Institutional Review Board at Taipei City Hospital (IRB/TCH) (ref: 960302)

**Study design**

Randomized controlled trial.

**Primary study design**

Interventional

**Secondary study design**

Randomised controlled trial

**Study setting(s)**

Not specified

**Study type(s)**

Treatment

**Participant information sheet****Health condition(s) or problem(s) studied**

Insomnia

**Interventions**

1. Acupuncture
2. Stilnox® (non-benzodiazepines) 5 mg/day at bedtime (hs) for 4 weeks

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome measure**

Pittsburgh Sleep Quality Index (PSQI) records (PSQI assessment will be carried out on a weekly basis).

**Secondary outcome measures**

Serious adverse events.

**Overall study start date**

03/08/2007

**Completion date**

31/05/2008

## Eligibility

**Key inclusion criteria**

Participants suffering from insomnia who have not taken any sleep medications associated with psychiatric disorders within 30 days.

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Both

**Target number of participants**

90

**Key exclusion criteria**

1. Pregnancy
2. Central nervous system disease such as seizure disorder, cerebrovascular disease, dementia etc.

**Date of first enrolment**

03/08/2007

**Date of final enrolment**

31/05/2008

## Locations

**Countries of recruitment**

Taiwan

**Study participating centre**

145 Zhengzhou Road

Taipei City

Taiwan

10341

# Sponsor information

## Organisation

Taipei City Hospital (Taiwan)

## Sponsor details

145 Zhengzhou Road

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## Sponsor type

Hospital/treatment centre

## ROR

<https://ror.org/02gzfb532>

# Funder(s)

## Funder type

Not defined

## Funder Name

Taipei City Government (Taiwan)

# Results and Publications

## Publication and dissemination plan

Not provided at time of registration

## Intention to publish date

## Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration