# The Efficacy of Reiki in the Treatment of Fibromyalgia

Submission date	Recruitment status  No longer recruiting	<ul><li>Prospectively registered</li></ul>		
14/08/2004		☐ Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
21/09/2004		[X] Results		
Last Edited	Condition category	Individual participant data		
25/01/2019	Musculoskeletal Diseases			

#### Plain English summary of protocol

Not provided at time of registration

## Contact information

#### Type(s)

Scientific

#### Contact name

Dr Dedra Buchwald

#### Contact details

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## Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number NCT00051428

Secondary identifying numbers

R21 AT001075

## Study information

#### Scientific Title

Reiki for the treatment of fibromyalgia: a randomized controlled trial

#### **Study objectives**

Not provided at time of registration

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Not provided at time of registration

#### Study design

Randomised controlled trial

#### Primary study design

Interventional

#### Secondary study design

Randomised controlled trial

#### Study setting(s)

Not specified

#### Study type(s)

**Not Specified** 

#### Participant information sheet

### Health condition(s) or problem(s) studied

Fibromyalgia (FM)

#### **Interventions**

One hundred Reiki-naive FM patients will participate in an 8-week trial. Patients will be randomized into one of two Reiki groups (direct-contact and distant Reiki) or one of two control groups (sham and placebo). Patients will receive either Reiki or placebo 16 times during the course of the study.

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome measure

Not provided at time of registration

#### Secondary outcome measures

Not provided at time of registration

#### Overall study start date

01/01/2003

#### Completion date

28/02/2005

## **Eligibility**

### Key inclusion criteria

Male or Female, aged 18-75 with Fibromyalgia

#### Participant type(s)

**Patient** 

#### Age group

Adult

#### Lower age limit

18 Years

#### Upper age limit

75 Years

#### Sex

Both

#### Target number of participants

100

#### Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/01/2003

#### Date of final enrolment

28/02/2005

## Locations

#### Countries of recruitment

United States of America

#### Study participating centre

#### **University of Washington**

Seattle United States of America 98101

## Sponsor information

#### Organisation

National Center for Complementary and Alternative Medicine (NCCAM) - USA

#### Sponsor details

6707 Democracy Blvd room 401 Bethesda United States of America 20892-5475

#### Sponsor type

Government

#### Website

http://nccam.nih.gov/

#### ROR

https://ror.org/00190t495

## Funder(s)

#### Funder type

Government

#### **Funder Name**

National Center for Complementary and Alternative Medicine (NCCAM) - USA

## **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

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2008	25/01/2019	Yes	No