# The efficacy of the Rességuier method in the treatment of fibromyalgia: a randomised controlled trial

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
15/12/2008	No longer recruiting	☐ Protocol
Registration date 23/12/2008	Overall study status Completed	Statistical analysis plan
		Results
<b>Last Edited</b> 23/12/2008	<b>Condition category</b> Musculoskeletal Diseases	Individual participant data
		<ul><li>Record updated in last year</li></ul>

### Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Dr Susanna Maddali Bongi

### Contact details

Associazione per lo studio della Sclerosi Sistemica e delle Malattie Fibrosanti (ASSMaF) Viale Pieraccini 18

Firenze

Italy

50139

# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

 ${\bf Clinical Trials. gov\ number}$ 

Secondary identifying numbers

N/A

# Study information

### Scientific Title

Rehabilitation of fibromyalgia: efficacy of the Rességuier method - a randomised controlled trial

### **Study objectives**

Rehabilitation holds an important role in the treatment of fibromyalgia syndrome (FMS). Despite numerous studies, an agreement on a specific rehabilitation line, adaptable to the needs of a heterogeneous group of patients, has not been found yet. The Rességuier method is a rehabilitation technique never experimented in fibromyalgia.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Not provided at the time of registration

### Study design

Randomised controlled single centre 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

Fibromyalgia

### **Interventions**

FMS patients were recruited and randomly assigned to the Group A and Group B. The patients of Group A were individually treated with the Rességuier method for 8 weeks, 1 session a week. The patients of Group B were observed for 8 weeks.

### Intervention Type

Other

### Phase

Not Applicable

### Primary outcome measure

- 1. Reduction of pain (Numerical Rating Scale [NRS]: 0 10)
- 2. Improvement of disability and quality of life, assessed by the 36-item Short Form (SF-36) Health Survey and Fibromyalgia Impact Questionnaire (FIQ)

### Secondary outcome measures

- 1. Quality of movement (NRS: 0 10)
- 2. Ability to relax mind and body (NRS: 0 10)

### Overall study start date

10/01/2007

### Completion date

10/12/2007

# **Eligibility**

### Key inclusion criteria

- 1. Diagnosis of fibromyalgia according to the American College of Rheumatology (ACR) criteria
- 2. No age limits

### Participant type(s)

**Patient** 

### Age group

Other

### Sex

Both

# Target number of participants

36

### Key exclusion criteria

Does not meet the inclusion criteria

### Date of first enrolment

10/01/2007

### Date of final enrolment

10/12/2007

# Locations

### Countries of recruitment

Italy

### Study participating centre

### Associazione per lo studio della Sclerosi Sistemica e delle Malattie Fibrosanti (ASSMaF)

Firenze Italy 50139

# Sponsor information

### Organisation

University of Florence (Italy)

### Sponsor details

Department of Medicine Viale GB Morgagni 85 Florence Italy 50134

### Sponsor type

University/education

### Website

http://www.unifi.it

### **ROR**

https://ror.org/04jr1s763

# Funder(s)

## Funder type

University/education

### **Funder Name**

University of Florence (Italy) - Department of Medicine

# **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