

Norepinephrine-evoked pain in fibromyalgia

Submission date 15/01/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 15/01/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 06/09/2007	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Fibromyalgia

Interventions

1. Subcutaneous injection of 10 micrograms of norepinephrine diluted in 0.1 ml normal saline solution
2. Subcutaneous injection of 0.1 ml normal saline

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Norepinephrine

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2000

Completion date

01/01/2001

Eligibility

Key inclusion criteria

1. 20 patients with fibromyalgia
2. 20 patients with rheumatoid arthritis
3. 20 normal controls

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2000

Date of final enrolment

01/01/2001

Locations

Countries of recruitment

Mexico

Study participating centre

Chief Rheumatology Department

Mexico DF

Mexico

14080

Sponsor information

Organisation

Ignacio Chavez National Institute of Cardiology (Instituto Nacional de Cardiologia Ignacio Chavez) (Mexico)

Sponsor details

Juan Badiano 1
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Sponsor type

Research organisation

ROR

<https://ror.org/046e90j34>

Funder(s)

Funder type

Research organisation

Funder Name

Ignacio Chavez National Institute of Cardiology (Instituto Nacional de Cardiologia Ignacio Chavez) (Mexico)

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	01/01/2002		Yes	No

