Hypnosis and self-hypnosis, administered and taught by nurses, for the reduction of chronic pain: a controlled clinical trial

Submission date	Recruitment status	Prospectively registered		
06/09/2005	No longer recruiting Overall study status	[X] Protocol		
Registration date		Statistical analysis plan		
04/10/2005	Completed Condition category	Results		
Last Edited		Individual participant data		
10/09/2009	Signs and Symptoms	Record updated in last year		

Plain English summary of protocolNot provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Bernard Burnand

Contact details

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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

Hypnosis and auto-hypnosis do not reduce pain in patients with chronic pain

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

Various conditions engendering chronic pain

Interventions

Hypnosis and auto-hypnosis administered and taught by trained nurses

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Level of pain (VAS)

Secondary outcome measures

- 1. Physical activity (tridimensional accelerometer)
- 2. Health related quality of life (SF-36)

Overall study start date

01/05/1993

Completion date

31/12/1994

Eligibility

Key inclusion criteria

- 1. To be referred (including occasional self-referral) for the management of chronic pain
- 2. To live within manageable distances from the hospital
- 3. To be aged 18 or above
- 4. To understand and speak French adequately
- 5. To be able to see, hear and communicate (i.e. no alteration of consciousness)
- 6. To have no diagnosis of mental illness (e.g. psychosis)
- 7. To give informed consent

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

 $2 \times 65 = 130$

Key exclusion criteria

Patients were excluded from the study whenever they wished and/or when no conventional treatment could be proposed.

Date of first enrolment

01/05/1993

Date of final enrolment

31/12/1994

Locations

Countries of recruitment

Switzerland

Study participating centre

IUMSP

Lausanne Switzerland CH-1010

Sponsor information

Organisation

Hospital of Morges (Switzerland)

Sponsor details

Morges Switzerland CH-1110

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/026qe1m81

Funder(s)

Funder type

Government

Funder Name

Swiss National Science Foundation (Switzerland) (SNF 4034-35883)

Alternative Name(s)

Schweizerischer Nationalfonds, Swiss National Science Foundation, Fonds National Suisse de la Recherche Scientifique, Fondo Nazionale Svizzero per la Ricerca Scientifica, Fonds National Suisse, Fondo Nazionale Svizzero, Schweizerische Nationalfonds, SNF, SNSF, FNS

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Switzerland

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?
Protocol article	protocol	01/08/1994		Yes	No