ISRCTN32129167 https://doi.org/10.1186/ISRCTN32129167

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

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

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
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 x 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