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Can hypnosis decrease nausea and vomiting in a defined 3-day period in patients diagnosed with lymphoma or myeloma who are undergoing autologous bone marrow/stem cell transplantation?

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 12/09/2003	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 19/02/2014	**Condition category** Signs and Symptoms	[ ] Individual participant data [ ] Record updated in last year

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
N0263120043

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)
Hospital

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Nausea and vomiting

Interventions
1. Standard hypnotic technique
2. Self-hypnosis plus standard hypnotic technique

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

It is planned to use multiple t-tests to assess for significant differences in nausea and distress and vomiting levels between the hypnosis and relaxation groups. More power can be achieved by grouping all the data and it should be possible to look specifically for effects within each type of cancer group.

Secondary outcome measures
Not provided at time of registration

Overall study start date
06/01/2003

Completion date
01/06/2007

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
60 patients from Haematology

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
06/01/2003

Date of final enrolment
01/06/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

UCH
London
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Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
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Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
University College London Hospitals NHS Trust (UK)

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