

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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Contact details

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Additional identifiers

Protocol serial number

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Not Specified

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(s)

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.

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/06/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

Study participating centre

UCH

London

United Kingdom

WC1E 6AU

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

University College London Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration