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A randomized controlled trial of venlafaxine versus placebo for depression amongst persons with lymphoma or leukaemia

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 30/11/2015	**Condition category** Mental and Behavioural Disorders	

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
N0079096832

Study information

Scientific Title
A randomized controlled trial of venlafaxine versus placebo for depression amongst persons with lymphoma or leukaemia

Study objectives
Is venlafaxine an effective antidepressant for depression amongst persons with lymphoma or leukaemia?

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)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Depression amongst persons with lymphoma or leukaemia

Interventions
A randomized controlled trial of venlafaxine versus placebo

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Venlafaxine

Primary outcome measure
Hamilton Rating Scale for Depression

Secondary outcome measures
Not provided at time of registration

Overall study start date
07/01/2001

Completion date
31/12/2003

Eligibility

Key inclusion criteria
Persons with lymphoma or leukaemia with depression as assessed using the Hamilton Rating Scale for Depression.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
07/01/2001

Date of final enrolment
31/12/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

University of Exeter
Exeter
United Kingdom
EX2 5AF

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

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

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Devon Partnership 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