The effect of flicker stimulation of peripheral visual field on short-term recall.

Submission date	Recruitment status No longer recruiting	Prospectively registered		
30/09/2004		☐ Protocol		
Registration date 30/09/2004	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 14/07/2009	Condition category Signs and Symptoms	[] Individual participant data		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Jonathan Williams

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0176135497

Study information

Scientific Title

Study objectives

To test if flicker can enhance short-term recall in older people. To test if any flicker effects on short term recall are frequently-specific.

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)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Signs and Symptoms: Memory

Interventions

Randomised trial. The intervention is learning and recall of words. A flicker machine will occur on light emitting diodes, the randomisation will be the flicker frequencies. It is thought that specific frequencies (near 10Hz) will enhance recognition and lower and higher frequencies will not.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The endpoints of the study will be recognition of words that participants saw during the learning phase. We will test if particular frequencies of flicker that occurred in the learning phase enhance recognition.

Secondary outcome measures

Not provided at time of registration

Overall study start date

19/12/2003

Completion date

28/02/2005

Eligibility

Key inclusion criteria

50 healthy people, 50 patients

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

100

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

19/12/2003

Date of final enrolment

28/02/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre OPTIMA

Oxford United Kingdom OX2 6HE

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Oxford Radcliffe 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	05/03/2006		Yes	No