# Effect of rapid transcranial magnetic stimulation (rTMS) on subjective responses to methamphetamine in healthy subjects

Submission date	<b>Recruitment status</b> No longer recruiting	Prospectively registered		
06/04/2000		∐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
06/04/2000	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
03/05/2011	Mental and Behavioural Disorders			

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Mr PJ Cowen

#### Contact details

University Department of Psychiatry University of Oxford Warneford Hospital Old Road Headington Oxford United Kingdom OX4 4XN

# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers

# Study information

#### Scientific Title

#### **Study objectives**

The purpose of this study is to test the hypothesis that repetitive Transcranial Magnetic Stimulation (rTMS) can attenuate the behavioural response to amphetamine in healthy volunteers.

### 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

### Health condition(s) or problem(s) studied

Neurosciences, psychiatry

#### Interventions

rTMS versus sham application (placebo). Both groups receive methamphetamine.

#### **Intervention Type**

Drug

#### Phase

**Not Specified** 

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

Methamphetamine

#### Primary outcome measure

Not provided at time of registration

#### Secondary outcome measures

Not provided at time of registration

## Overall study start date

01/03/1999

# Completion date

01/03/2000

# **Eligibility**

#### Key inclusion criteria

- 1. No significant current or past psychiatric disorder or substance misuse
- 2. No current physical illness

# Participant type(s)

**Patient** 

#### Age group

Adult

#### Sex

Both

# Target number of participants

24

## Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/03/1999

#### Date of final enrolment

01/03/2000

# Locations

#### Countries of recruitment

England

**United Kingdom** 

## Study participating centre

## **University Department of Psychiatry**

Oxford United Kingdom OX4 4XN

# Sponsor information

#### Organisation

Medical Research Council (MRC) (UK)

#### Sponsor details

20 Park Crescent London United Kingdom W1B 1AL +44 (0)20 7636 5422 clinical.trial@headoffice.mrc.ac.uk

#### Sponsor type

Research council

#### Website

http://www.mrc.ac.uk

# Funder(s)

#### Funder type

Research council

#### **Funder Name**

Medical Research Council (UK)

#### Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

#### Funding Body Type

Government organisation

#### Funding Body Subtype

National government

#### Location

**United Kingdom** 

# **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	01/12/2000		Yes	No