# Psychological mechanisms in suicidal behaviour

Submission date	Recruitment status  No longer recruiting	Prospectively registered		
03/03/2005		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
22/07/2005	Completed	[X] Results		
<b>Last Edited</b> 28/11/2012	Condition category  Mental and Behavioural Disorders	[] Individual participant data		

#### Plain English summary of protocol

Background and study aims.

This study aimed to examine the acceptability and mechanisms of action of Mindfulness-based Cognitive Therapy (MBCT) delivered to patients in remission, but with a history of serious suicidal thoughts or behaviour. The aim of the study was to explore the feasibility of delivering MBCT to this population, and the extent to which treatment with MBCT could reduce vulnerability to suicidal thoughts and behaviour through an examination of its effects on psychological factors linked to of suicidal vulnerability.

#### Who can participate?

Participants aged between 18 and 65 who were currently well but reported at least one prior episode of major depression accompanied by serious suicidal thoughts were recruited from general practitioners and local psychologists/psychiatrists and from the community. All participants were required to be well (no more than one week of minimal depressive symptoms in the past 8 weeks), and to have experienced no episodes of mania for at least 6 months. Additional exclusion criteria included current psychosis, obsessive compulsive disorder or eating disorder as their main problem, current deliberate self harm on a regular basis, a neurological disorder or an inability to complete assessments due to language difficulties or cognitive impairment.

#### What does the study involve?

Participants were interviewed using the Mini International Neuropsychiatric Interview to establish psychiatric history. Following this, participants completed a number of other assessment measures assessing aspects of cognitive vulnerability to depression and suicidality. These measures included assessment of residual symptoms of depression, mood-related impairments in problem solving and future thinking, autobiographical memory deficits, tendency to suppress unwanted thoughts and self-discrepancy (perceptions of distance between how one currently sees themselves and how one would like to be). Participants were then randomly allocated to either immediate treatment with MBCT or a waitlist condition. The measures were completed again at the end of treatment or waitlist phase. Following this the waitlist group received treatment with MBCT.

What are the possible benefits and risks of participating?

Benefits included the fact that all participants were offered treatment with Mindfulness Based Cognitive Therapy, either immediately or at the end of the waitlist phase of the study. Potential

risks to participants related to the distress of reporting on prior psychiatric history and the inherent challenges of engaging in a therapeutic process designed to target vulnerability to recurrent depression and suicidality.

Where is the study run from? Department of Psychiatry, University of Oxford

When is the study starting and how long is it expected to run for? The study started in May 2005 and ended in December 2005

Who is funding the study? The Wellcome Trust

Who is the main contact?
Professor Mark Williams
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## Contact information

# Type(s)

Scientific

#### Contact name

Prof J. Mark G. Williams

#### Contact details

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# Additional identifiers

Protocol serial number PRG/06/01

# Study information

Scientific Title

#### Study objectives

This is a preliminary (explanatory) trial to assess the immediate effects of Mindfulness-based Cognitive Therapy (MBCT) on cognitive reactivity (the tendency to react to small changes in mood with a catastrophic and rapidly escalating pattern of suicidal thinking).

#### 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

#### Study type(s)

**Treatment** 

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

Suicidality

#### **Interventions**

Treatment:

Mindfulness-Based Cognitive Therapy (MBCT): manualised, eight week treatment combining stress reduction techniques and cognitive therapy and Treatment As Usual (TAU)

#### Control:

Waiting List control and Treatment As Usual (TAU)

#### Intervention Type

Other

#### Phase

Not Applicable

#### Primary outcome(s)

- 1. Depression
- 2. Frequency of thought suppression
- 3. Change in problem solving and future thinking following experimental induction of mood
- 4. Self-rated mindfulness
- 5. Discrepancy between actual and ideal self-guides
- 6. Specificity of autobiographical memory

#### Key secondary outcome(s))

No secondary outcome measures

#### Completion date

01/12/2005

# **Eligibility**

#### Key inclusion criteria

- 1. Aged 18 to 64
- 2. History of major depression with suicidal ideation and/or suicidal behaviour
- 3. Not currently depressed or suicidal (at least eight weeks symptom free)

### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

Adult

#### Lower age limit

18 years

#### Sex

All

#### Key exclusion criteria

- 1. Visually impaired
- 2. Not fluent in English
- 3. Habitual self-damaging acts
- 4. Bipolar disorder
- 5. Schizophrenia
- 6. Active substance abuse
- 7. Eating disorder
- 8. Primary Obsessive Compulsive Disorder (OCD)

#### Date of first enrolment

01/05/2005

#### Date of final enrolment

01/12/2005

# **Locations**

#### Countries of recruitment

United Kingdom

England

# Study participating centre Oxford University

Oxford United Kingdom OX3 7JX

# Sponsor information

## Organisation

University of Oxford (UK)

#### **ROR**

https://ror.org/052gg0110

# Funder(s)

# Funder type

Charity

#### Funder Name

The Wellcome Trust (UK) (grant ref: 067797)

# **Results and Publications**

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	preliminary results	01/04/2008		Yes	No
Results article	results	01/12/2008		Yes	No
Results article	results	01/06/2009		Yes	No