# Dialectical behaviour therapy in patients with borderline personality disorder who self-harm: a pragmatic exploratory trial

Submission date	<b>Recruitment status</b> No longer recruiting	Prospectively registered		
30/07/2008		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
25/09/2008	Completed	[X] Results		
<b>Last Edited</b> 23/07/2014	Condition category  Mental and Behavioural Disorders	Individual participant data		

# Plain English summary of protocol

Not provided at time of registration

# Contact information

## Type(s)

Scientific

#### Contact name

Prof Stefan Priebe

#### Contact details

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# Additional identifiers

# Protocol serial number

PB-PG-0906-10540

# Study information

#### Scientific Title

#### Acronym

**DIALECT** 

#### **Study objectives**

Self-harming patients receiving a one-year course of dialectical behaviour therapy (DBT) will self-harm less over the 12 months than patients receiving one year of treatment as usual.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

The Camden and Islington Community Local Research Ethics Committee, 18/02/2008, ref: 07/H0722/98

#### Study design

Pilot randomised controlled pragmatic exploratory trial

#### Primary study design

Interventional

#### Study type(s)

Treatment

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

Borderline personality disorder, self-harm

#### **Interventions**

DBT and care co-ordination versus waiting list control group with standard NHS care, over 12 months. DBT consists of 3 hours of therapy a week: this comprises 1 hour of individual therapy and 2 hours of group skills classes.

#### Intervention Type

Other

#### Phase

Not Applicable

#### Primary outcome(s)

Days with self-harm during the 12-month period.

#### Key secondary outcome(s))

- 1. Pre-post changes in self-harming during the 12-month period
- 2. Number of accident and emergency (A&E) attendances during the 12-month period
- 3. Inpatient admissions during the 12-month period
- 4. Use of other services in primary and secondary care during the 12-month period
- 5. Service costs during the 12-month period
- 6. Use of medication during the 12-month period

- 7. Pre-post changes in self-rated and observer-rated symptom level and quality of life at the end of the 12 month period
- 8. Quality of the therapeutic relationship at the end of the 12-month period
- 9. Treatment satisfaction at the end of the 12-month period

## Completion date

01/07/2011

# **Eligibility**

#### Key inclusion criteria

- 1. Frequent self-harm (more than 5 days with self-harm over 12 months)
- 2. Aged 16 years and older, either sex
- 3. Sufficient command of English
- 4. At least one personality disorder

#### Participant type(s)

**Patient** 

### Healthy volunteers allowed

No

#### Age group

Adult

#### Lower age limit

16 years

#### Sex

All

#### Key exclusion criteria

Learning disabilities

#### Date of first enrolment

01/02/2008

#### Date of final enrolment

01/07/2011

# Locations

#### Countries of recruitment

United Kingdom

England

## Study participating centre

# Unit for Community and Social Psychiatry

London United Kingdom E13 8SP

# Sponsor information

## Organisation

East London NHS Foundation Trust (UK)

#### **ROR**

https://ror.org/01q0vs094

# Funder(s)

#### Funder type

Government

#### **Funder Name**

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) Programme

# **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	results	01/01/2012	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes