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

| Submission date 30/07/2008 | Recruitment status No longer recruiting | Prospectively registered | | |
|-------------------------------|--|--------------------------------|--|--|
| | | ☐ Protocol | | |
| Registration date | Overall study status | Statistical analysis plan | | |
| 25/09/2008 | Completed | [X] Results | | |
| Last Edited 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

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

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 measure

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

Secondary outcome measures

- 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

Overall study start date

01/02/2008

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

Age group

Adult

Lower age limit

16 Years

Sex

Both

Target number of participants

60

Key exclusion criteria

Learning disabilities

Date of first enrolment 01/02/2008

Date of final enrolment 01/07/2011

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Unit for Community and Social Psychiatry
London
United Kingdom
E13 8SP

Sponsor information

Organisation

East London NHS Foundation Trust (UK)

Sponsor details

Trust Headquarters
EastONE
22 Commercial Street
London
England
United Kingdom
E1 6LP
+44 (0)20 7655 4000
webadmin@eastlondon.nhs.uk

Sponsor type

Hospital/treatment centre

Website

http://www.eastlondon.nhs.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

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 | 01/01/2012 | | Yes | No |