ISRCTN24549546 https://doi.org/10.1186/ISRCTN24549546

A Pragmatic Evaluation of Manualised cognitive therapy on suicide ideation in service users with deliverate self-harm in acute psychiatric settings.

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 03/09/2012	**Condition category** Mental and Behavioural Disorders	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0573157722

Study information

Scientific Title

Study objectives
Will a brief manualised cognitive therapy intervention reduce suicide ideation, duration of level one observation and length of stay, in an adult acute in-patient setting?

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
Mental and Behavioural Disorders: Self harm

Interventions
Cognitive therapy intervention vs standard care

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
1. Beck Scale for Suicide Ideation (BSSI)
2. Duration of Level One Observation
3. Length of Stay.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2004

Completion date
30/12/2006

Eligibility

Key inclusion criteria
Patients admitted to the adult acute in-patient wards in Newcastle & North Tyneside with suicide ideation and at significant risk of Deliberate Self Harm.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
40

Key exclusion criteria
1. Diagnosis of personality disorder
2. Diagnosis of organic disorder
3. Incapacity to consent
4. Unable to comprehend English

Date of first enrolment
01/11/2004

Date of final enrolment
30/12/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Newcastle General Hospital
Newcastle upon Tyne
United Kingdom
NE4 6BE

Sponsor information

Organisation
Department of Health

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Richmond House
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Sponsor type
Government

Website
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Funder(s)

Funder type
Government

Funder Name
Newcastle, North Tyneside and Northumberland Mental Health NHS Trust (UK) NHS R&D Support Funding

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