ISRCTN77981324 https://doi.org/10.1186/ISRCTN77981324

Randomised controlled trial of a telephone support 'help card' versus usual care for self harm (SH) patients: effects on repetition

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 14/04/2015	**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
N0079102564

Study information

Scientific Title
Randomised controlled trial of a telephone support 'help card' versus usual care for self harm (SH) patients: effects on repetition

Study objectives
Does presenting a 'help card' with details of relevant telephone support/advice to self harm patients alter the rate of self harm repetition?

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)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Mental and Behavioural Disorders: Self harm

Interventions
Card versus treatment as usual

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Repetition of deliberate self harm (identified) by 6 months.

Secondary outcome measures
Not provided at time of registration

Overall study start date

01/08/2000

Completion date
01/01/2006

Eligibility

Key inclusion criteria
Patients presenting with deliberate self harm to a district general hospital and subsequently seen by members of a specialist self harm service are randomised to receive a card versus treatment as usual. Criteria are adulthood and valid consent.

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/2000

Date of final enrolment
01/01/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Wonford House Hospital
Exeter
United Kingdom
EX2 5AF

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
Devon Partnership NHS Trust (UK)

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