ISRCTN81861513 https://doi.org/10.1186/ISRCTN81861513

A randomised trial of a General Practice based intervention to reduce repeat deliberate self harm (parasuicide).

Submission date
23/01/2004

Recruitment status
No longer recruiting

Registration date
23/01/2004

Overall study status
Completed

Last Edited
22/02/2008

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] Individual participant data

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
C/HP/07/11.95/Gunnell

Study information

Scientific Title

Acronym
DASH

Study objectives
To what extent can a general practitioner (GP) based intervention affect the rate of repetition of deliberate self harm?

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
Depression, anxiety, neuroses

Interventions
1. GP intervention
2. Standard care

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
1. Deliberate self-harm repetition rates
2. Cost effectiveness of the intervention
3. Production of GP management guidelines for DSH

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/1996

Completion date
30/09/2000

Eligibility

Key inclusion criteria
1. Patients attending accident and emergency (A&E) departments in Bristol and Bath after an episode of deliberate self harm (DSH), registered with general practitioners in Avon.

1932 - patients were studied in total.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
1932

Key exclusion criteria
Cases of alcohol taken alone and illicit drug overdose, except where the casualty officer felt that the purpose of the act was self-harm or suicide.

Date of first enrolment
01/10/1996

Date of final enrolment
30/09/2000

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Bristol
Bristol
United Kingdom
BS8 2PR

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
NHS Executive South West (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	25/05/2002		Yes	No