An exploratory randomised trial of brief intervention for alcohol misuse among people who self harm and misuse alcohol: impact on repetition of self harm

Submission date	Recruitment status No longer recruiting	Prospectively registered	
19/08/2005		☐ Protocol	
Registration date 07/09/2005	Overall study status Completed	Statistical analysis plan	
		[X] Results	
Last Edited 06/01/2011	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

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Acronym

SHARP (Self Harm Alcohol Reduction Programme)

Study objectives

Among people who present to emergency hospital services following deliberate self harm and are found to be misusing alcohol, the proportion who repeat self harm over the following six months will be lower among those offered a self-help leaflet and an appointment for brief intervention from an Alcohol Health Worker than among those who receive a self-help leaflet on alcohol and health.

Ethics approval required

Old ethics approval format

Ethics approval(s)

West London REC 2 on 24/05/2010 10/H0706/29

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

Alcohol misuse

Interventions

Experimental treatment: A leaflet on alcohol and health together with an appointment card asking the patient to attend an appointment with an Alcohol Health Worker (AHW). The AHW will provide a single session of assessment, brief counselling based on the FRAMES approach (Bien et al., 1993), and referral on to other services.

Control treatment (CT): A leaflet on alcohol and health

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Repetition of deliberate self harm over the following 6 months (as measured by self report together with an examination of hospital records).

Secondary outcome measures

Level of alcohol consumption, mental health and satisfaction with care at six months

Overall study start date

01/09/2005

Completion date

31/08/2007

Eligibility

Key inclusion criteria

All those aged over 18 years or over, who are found to be misusing alcohol (according to the Paddington Alcohol Test), and have an address in the local area (within the M25).

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

270

Key exclusion criteria

All those unwilling to provide informed consent to participate in the trial, all those unable to complete baseline assessment because of an inability to communicate in English, and all those who are already in contact with alcohol misuse services will be excluded from the study.

Date of first enrolment

01/09/2005

Date of final enrolment

31/08/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Psychological Medicine
London
United Kingdom
W6 8LN

Sponsor information

Organisation

Imperial College London (UK)

Sponsor details

G02, Sir Alexander Fleming Building Imperial College London London England United Kingdom SW7 2AZ

Sponsor type

University/education

ROR

https://ror.org/041kmwe10

Funder(s)

Funder type

Charity

Funder Name

St Marys Charitable Trust (AE08)

Results and Publications

Publication and dissemination planNot provided at time of registration

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	01/11/2010		Yes	No