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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
19/08/2005

Recruitment status
No longer recruiting

Registration date
07/09/2005

Overall study status
Completed

Last Edited
06/01/2011

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

11.01.05

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 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	01/11/2010		Yes	No