Reducing Suicide In Men through general practice (The SIM Study)

Submission date	Recruitment status No longer recruiting	Prospectively registered		
23/01/2007		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
04/05/2007	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
23/09/2021	Mental and Behavioural Disorders			

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

N/A

Study information

Scientific Title

Reducing Suicide In Men through general practice (The SIM Study)

Acronym

SIM

Study objectives

The objectives of this study are to examine whether General Practitioners (GPs) who have completed the Suicide In Men (SIM) training package in assessing and managing suicide risk in men, demonstrate:

- 1. An increased understanding of the science, epidemiology, interventions and statistics related to male suicide
- 2. A raised awareness of the scale of the problem and the barriers to managing it
- 3. Increased competency to improve suicide assessment and management of at-risk patients

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval is currently in progress at the University of Melbourne Human Research Ethics Committee

Study design

This study is a randomised controlled trial of the SIM training package, with GPs randomised to training and wait-listed (control) groups.

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Depression and suicide risk in male general practice patients

Interventions

The SIM education package includes:

- 1. A training workshop
- 2. Completion of a ten minute survey before and after training
- 3. Undertake two 30 minute simulated patient consultations at their practice during the study

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

GPs improvement in:

- 1. Knowledge
- 2. Attitude/awareness of suicide management
- 3. Competence in the assessment and management of male suicide risk

Secondary outcome measures

Comparison of pre-training GP knowledge and attitudes with performance in simulated consultations will be undertaken

Overall study start date

31/01/2007

Completion date

31/12/2007

Eligibility

Key inclusion criteria

GPs registered in the state of Victoria, Australia. This study does not involve patients.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

56 GPs in total, 28 in each arm of the study

Key exclusion criteria

None specified

Date of first enrolment

31/01/2007

Date of final enrolment

31/12/2007

Locations

Countries of recruitment

Australia

Study participating centre Department of General Practice

Carlton Australia 3053

Sponsor information

Organisation

The beyondblue Victorian Centre of Excellence in Depression and Related Disorders (Australia)

Sponsor details

PO Box 6100 West Hawthorn Australia 3122 +61 (0)3 9810 6100 bb@beyondblue.org.au

Sponsor type

Government

Website

http://www.beyondblue.org.au/index.aspx?link_id=6.841

ROR

https://ror.org/05mwvz623

Funder(s)

Funder type

Government

Funder Name

The beyondblue Victorian Centre of Excellence in Depression and Related Disorders (Australia)

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		10/09/2013	23/09/2021	Yes	No