

An evaluation of resilience interventions for emergency workers

Submission date 05/02/2016	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 08/02/2016	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 16/10/2020	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

The aim of this study is to evaluate resilience training for emergency workers. The group-based intervention has been developed by the national mental health charity, Mind. We will compare Mind's group-based resilience intervention for emergency personnel with the same intervention delivered with one online top-up training session.

Who can participate?

Student paramedics who do not have post-traumatic stress disorder or depression

What does the study involve?

Participants are randomly allocated to receive either Mind's resilience intervention or Mind's resilience intervention plus a new internet-based top-up session. The two groups' resilience and use of psychological coping strategies are then compared.

What are the possible benefits and risks of participating?

Participation in the study could lead to participants' resilience being improved. Participation will also guide improvements to future resilience interventions. There are no risks associated with taking part.

Where is the study run from?

University of Brighton and Oxford-Brookes University (UK)

When is the study starting and how long is it expected to run for?

January 2016 to October 2016

Who is funding the study?

Mind, the mental health charity (UK)

Who is the main contact?

Dr Jennifer Wild

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Contact information

Type(s)

Scientific

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Public

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Additional identifiers

Protocol serial number

01/02/16/V1

Study information

Scientific Title

An evaluation of resilience interventions for emergency workers: a randomised controlled trial

Study objectives

1. The format of group and top-up internet-based resilience training is effective.
2. Participants receiving the group plus internet-based resilience training will demonstrate less use of rumination as a strategy in response to an experimental task at follow-up.

Ethics approval required

Old ethics approval format

Ethics approval(s)

University of Oxford Central University Research Ethics Committee, 07/12/2015, ref: MS-IDREC-C1-2015-059

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Resilience and psychological coping strategies

Interventions

Participants will be randomly allocated to one of the following:

1. Six sessions of standard group-based resilience training
2. Six sessions of standard group-based resilience training plus a one hour internet-based top-up training

The total duration of follow-up is 6 months.

Intervention Type

Behavioural

Primary outcome(s)

Measured at baseline, post-intervention, and at follow-up (6 months):

1. Resilience (Connor-Davidson Resilience Scale [CD-RISC])
2. Rumination (Perseverative Thinking Questionnaire [PTQ] and Rumination Response Scale [RRS])

Key secondary outcome(s)

Measured at baseline, post-intervention, and at follow-up (6 months):

1. Days off work
2. Psychological coping strategies (Responses to Intrusions Questionnaire [RIQ])

Completion date

31/10/2016

Eligibility**Key inclusion criteria**

Student paramedics who do not have posttraumatic stress disorder or depression

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Student paramedics who are suffering from PTSD or major depression

Date of first enrolment

15/01/2016

Date of final enrolment

12/02/2016

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

University of Brighton

United Kingdom

BN1 9PH

Study participating centre

Oxford-Brookes University

United Kingdom

OX3 0BP

Sponsor information

Organisation

University of Oxford (UK)

ROR

Funder(s)

Funder type
Charity

Funder Name
Mind, national mental health charity (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes