

Post-deployment Battlemind training for UK Armed Forces personnel

Submission date 28/01/2009	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 05/03/2009	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 11/07/2013	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Neil Greenberg

Contact details
Academic Centre for Defence Mental Health
King's College London
Weston Education Centre
Cutcombe Road
London
United Kingdom
SE5 9RJ

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
CSA/08/039

Study information

Scientific Title

Post-deployment Battlemind training for UK Armed Forces personnel: a cluster-randomised controlled trial

Study objectives

Primary hypothesis:

UK Armed Forces Personnel who receive Battlemind training, when compared with personnel who receive a standard post-deployment stress and homecoming brief, will report:

1. Better mental health outcomes
2. Fewer risk taking behaviours
3. Greater perceived team support
4. Fewer problems in combat-to-home transition
5. Less stigmatising beliefs about seeking help for mental health problems

Secondary hypotheses:

1. Those participants who have had greater exposure to stressors during deployment will have poorer mental health at baseline and will benefit more from the Battlemind intervention than those with fewer traumatic deployment experiences
2. Personnel receiving Battlemind will have a more positive perception of the training received than those receiving the standard post-deployment stress brief

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Ministry of Defence Research Ethics Committee approval received 16th December 2008 (ref: 0863/218)
2. King's College Hospital Research Ethics Committee approval received 18th December 2008 (ref: 08/H0808/179)

Study design

Single centre cluster randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a participant information sheet

Health condition(s) or problem(s) studied

Mental Health

Interventions

The intervention group will receive Battlemind training, a US developed post-deployment mental health training package aimed at assisting personnel to adjust to life post-deployment. Battlemind aims to build on people's combat strengths and skills by showing how these strengths can be adapted for the home environment.

The control group will receive the standard post-deployment briefs about stress and homecoming that are currently delivered to personnel returning home from operational tour.

Both intervention and control briefs will be delivered during the 'decompression' period that personnel undergo on their return from deployment. Both briefs will be delivered in a single session lasting approximately 45 minutes.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Severity of post-traumatic stress symptoms as indicated by the score on the Post-Traumatic Stress Disorder Checklist (PCL). PCL will be assessed prior to receiving the post-deployment brief and again 4 months later.

Secondary outcome measures

1. Psychological morbidity - 12-item General Health Questionnaire (GHQ-12), assessed prior to the brief and again 4 months later
2. Stigma - Help-Seeking Stigma Questionnaire assessed prior to the brief and again 4 months later

The following will be assessed at 4 months:

3. Depression - 9-items from the Patient Health Questionnaire. As of 17/02/2010 the following outcome was also added to this record: Anxiety - 7-items from the Patient Health Questionnaire.
4. Alcohol consumption - Alcohol Use Disorders Identification Test (AUDIT)
5. Driving behaviour
6. Sleep quality
7. Quality of combat-to-home transition
8. Relationship satisfaction
9. Anger
10. Team support
11. Occupational and social functioning (added to this record on 17/02/2010)

Overall study start date

01/03/2009

Completion date

31/01/2011

Eligibility

Key inclusion criteria

1. UK Armed Forces personnel returning from operational tour
2. Aged 18 years and over; both men and women will be eligible to participate

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

2000

Key exclusion criteria

1. Individual augmentees who do not return as part of a formed unit
2. Units that are primarily composed of reserve personnel or headquarters staff

Date of first enrolment

01/03/2009

Date of final enrolment

31/01/2011

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Academic Centre for Defence Mental Health

London

United Kingdom

SE5 9RJ

Sponsor information

Organisation

King's College London (UK)

Sponsor details

Strand

London

England

United Kingdom

WC2R 2LS

Sponsor type

University/education

Website

<http://www.kcl.ac.uk/>

ROR

<https://ror.org/0220mzb33>

Funder(s)**Funder type**

Research organisation

Funder Name

Haldane-Spearman Consortium (UK) (ref: TIN no. HSC_07_01_02_002)

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	01/06/2012		Yes	No