

Efficacy of a school-based psychosocial intervention to deal with the psychosocial impact of armed conflict on school-aged children in Indonesia

Submission date
09/02/2006

Recruitment status
No longer recruiting

Prospectively registered

Protocol

Registration date
09/06/2006

Overall study status
Completed

Statistical analysis plan

Results

Last Edited
15/08/2008

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

Prof Joop de Jong

Contact details

c/o HealthNet TPO

Tolstraat 127

1074 VJ

Amsterdam

Netherlands

1074 VJ

+31 (0)20 6200005

jdejong@healthnettpo.org

Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Study objectives

A psychosocial school-based program is capable of reducing conflict-related psychosocial symptoms, and increasing children's strength to deal with armed-conflict related psychosocial difficulties

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ongoing at ethical review board of Vrije Universiteit Amsterdam as of 09/06/06

Study design

Cluster randomisation of schools to intervention (structured school-based psychosocial program) or waitlist condition (receiving treatment after the research)

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Multiple impact of armed conflict on psychosocial wellbeing (post-traumatic stress disorder)

Interventions

Classroom-based structured, manualized psychosocial intervention, called the class-room based intervention (CBI), as designed by the Center for Trauma Psychology (Boston, Massachusetts) versus the waitlist condition.

The CBI entails a 15-session program that encompasses working with the trauma narrative, creative techniques (drama, music), and specifically designed games. The waitlist condition simply entails the provision of treatment after the research is finished.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Post-traumatic stress disorder (PTSD) and depressive symptoms

Key secondary outcome(s))

1. Anxiety
2. Aggression
3. Daily functioning
4. Social support
5. Sociometric measurements (sociogram)

6. Coping
7. Family functioning
8. Hope
9. School functioning (grades, absenteeism)

Completion date

31/05/2007

Eligibility

Key inclusion criteria

Children screened for exposure to traumatic events, post-traumatic stress symptoms or depressive anxiety symptoms, with the use of symptom checklists

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Sex

All

Key exclusion criteria

Serious psychopathology and psychiatric disorders (mutism, retardation, psychotic symptoms) or incapability to function in a group (conduct disorders, harming others), as judged by local psychosocial counsellors

Date of first enrolment

15/03/2006

Date of final enrolment

31/05/2007

Locations

Countries of recruitment

Indonesia

Netherlands

Study participating centre

c/o HealthNet TPO
Amsterdam
Netherlands
1074 VJ

Sponsor information

Organisation

HealthNet TPO (The Netherlands)

ROR

<https://ror.org/0088fq38>

Funder(s)

Funder type

Charity

Funder Name

Plan International (UK)

Results and Publications

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	13/08/2008		Yes	No