

Emotion regulation as the stepping stone for youth mental health: testing the efficacy of the prevention program 'Boostcamp'

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Registration date 13/06/2017	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 23/07/2018	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 transition from childhood into adolescence is a time of development, which involves physical, mental, emotional challenges. These developments often coincide with issues such as stronger reactions and vulnerability to developing mental health problems. Studies have shown that at least 15-20% of adolescents suffer from mental health issues. Boost Camp is a new universal prevention program which aims to stimulate the use of emotion regulation strategies, improve adolescent mental health, lower risk for developing mental health problems during the stressful life period of school transition, and increasing mental health knowledge while decreasing stigmatization (negative associations). The aim of this study is to find out whether delivering this program to adolescents can help improve their mental health and emotion regulation.

Who can participate?

Adolescents in their first year of secondary school.

What does the study involve?

Participating schools are randomly allocated to one of two groups. Those in the first group continue as normal for the duration of the study. Those in the second group take part in the Boost Camp program. This involves learning and practicing emotional awareness, acceptance, relaxation, distraction, cognitive reappraisal and problem solving through exercises, both as individuals and in groups. Participants fill out questionnaires about mental health and emotion regulation four times through the first year of high school. This happens during school time and under the supervision of the classroom teacher and someone of the department.

What are the possible benefits and risks of participating?

Participating in the study can be beneficial for the participating schools, because they can receive an overall report of the results about the mental health status of their students. The classes who participate gain a class photo-shoot as a reward for their commitment. There are no significant risks involved with participating. Although participation requires time and investment from management board, teachers and their students.

Where is the study run from?
Six schools located in Flanders (Belgium)

When is the study starting and how long is it expected to run for?
April 2016 to May 2018

Who is funding the study?
Research Foundation Flanders (Belgium)

Who is the main contact?
1. Miss Brenda Volkaert (public)
2. Miss Laura White (scientific)

Contact information

Type(s)
Public

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Additional identifiers

Protocol serial number
G0E9217N

Study information

Scientific Title

Evaluating efficacy of a school-based transdiagnostic adolescent mental health prevention program, targeting emotion regulation: a randomised controlled trial

Study objectives

Compared to adolescents in the control group, adolescents in the Boost Camp group will show better mental health and more adaptive emotion regulation.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Cluster randomised controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Emotion Regulation (ER)

Interventions

When schools agree to participate, they are randomly assigned to the control or the intervention condition. In this way, randomisation is clustered since the participating classes of one and the same school are always in the same condition.

Intervention group: Students from schools in the intervention group receive an school-based adolescent training, named 'Boostcamp'. This is a group training for adolescents who are making the transition from primary to secondary school and focuses on improving ER and social skills. The program consists of three different parts:

The program consists of three different parts:

1. An information moment in which the program is explained and written informed consent will be collected
2. A two-day 'immersion' training program
3. Two booster sessions over the subsequent three months of the first semester.

The training is organized at the beginning of secondary school. During two full days, youngsters learn about adaptive ER strategies, social skills, and how to use them in challenging emotional and social situations. The trainers or employees or students of our department, with at least a bachelor's degree in psychology and are coached and guided by the two main investigators who are also trained clinical psychologists.

Control group: Students from schools in the control condition are asked to fill out all the self report questionnaires just like students in the experimental condition and on the same time. However, the control condition do not participate in the two-way training program or the boostersessions.

There are two follow-up measurements: one at the end of the first semester (December 2017) and one at the end of the second semester (May 2018). Both assessments include the same questionnaires as at the baseline.

The students in the experimental condition receive two 'boostersessions', in which they repeat the learned theory and skills of the Boostcamp and get opportunities to practice with lifelong situations under the supervision of trainer. These Boostersessions take place before and between the two follow-up moments (at the end of December and at the end of March).

Intervention Type

Behavioural

Primary outcome(s)

1. Adolescent mental health is assessed using the Youth Self Report (YSR), Positive and Negative Affect Schedule for Children (PANAS-C), Children's Depression Inventory (CDI), Compententie Belevingsschaal voor Adolescenten (CBSA), and KIDSCREEN-10 HRQoL for Children and Adolescents (KIDSCREEN-10) at baseline, 8 weeks and 6 months
2. Emotion regulation is assessed using the FEEL-KJ questionnaire, the emotion regulation skills questionnaire (ERSQ) and he subscale 'Emotional Awareness' of the Difficulties in Emotion Regulation Scale (DERS) at baseline, 8 weeks and 6 months. The positive and negative affect (PANAS-C), depression (CDI), emotion regulation skills (ERSQ) and Global Self Worth (CBSA) are assessed immediatly after the two days training (in the last hour of the training).

Key secondary outcome(s)

1. Mental health knowledge and stigma is measured using qualitative questions at baseline, 8 weeks and 6 months
2. School absenteeism and academic achievement are measured by the school through the whole school year as usual and will be delivered to the investigator after the last questionnaires

Completion date

31/05/2018

Eligibility

Key inclusion criteria

1. Aged 10 - 14 years
2. In the first year of secondary school
3. In a participating class

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Child

Lower age limit

10 years

Upper age limit

14 years

Sex

All

Key exclusion criteria

Not meeting inclusion criteria.

Date of first enrolment

01/02/2017

Date of final enrolment

01/09/2017

Locations

Countries of recruitment

Belgium

Study participating centre

Leiepoort campus Sint-Hendrik

Guido Gezellelaan 105

Deinze

Belgium

9800

Study participating centre

VTI

Leon Declercqstraat 1

Deinze

Belgium

9800

Study participating centre

Koninklijk Atheneum Erasmus Deinze

Volhardingslaan 11

Deinze

Belgium

9800

Study participating centre**Ten Doorn**

Zuidmoerstraat 125

Eeklo

Belgium

9900

Study participating centre**Mariagaard**

9230, Oosterzelesteenweg 80

Wetteren

Belgium

9230

Sponsor information

Organisation

Ghent University

ROR

<https://ror.org/00cv9y106>

Funder(s)

Funder type

Government

Funder Name

Fonds Wetenschappelijk Onderzoek

Alternative Name(s)

Research Foundation Flanders, Flemish Research Foundation, The FWO, Het FWO, FWO

Funding Body Type

Government organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Belgium

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated and/or analysed during the current study during this study will be included in the subsequent results publication.

IPD sharing plan summary

Other

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	protocol	21/07/2018		Yes	No