Orygen Youth Health Research Centre

RE-FRAME-IT: The development of a randomised controlled trial testing an e-health intervention on suicide risk among school students



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Background

Suicidality is common among young people [1-5] however there are few well designed studies testing interventions specifically among at-risk youth [6]. The development of e-health interventions have potential, yet despite some success treating depression and anxiety [7,8] their potential to reduce suicidality in youth remains unexamined.

Aims/hypotheses

Hypotheses are that the intervention will:

- 1. Reduce SI and / or DSH among participants
- 2. Reduce symptoms of depression, anxiety and hopelessness
- 3. Increase help-seeking
- 4. Improve coping and problem solving skills
- 5. Participants will find the receipt of this intervention acceptable, and
- 6. The intervention will not cause undue distress.

Methods

Design / setting: A randomised controlled trial conducted by Orygen Youth Health with government high schools in North West Melbourne.

Participants: 165 secondary school students in year 8-12 who have presented to the school welfare team for support with recent SI or DSH.

Student presents to school counsellor and screened for suicidal ideation or behaviour within the last month



Student invited to participate in RE-FRAME IT study

Student & parent give consent to participate Student chooses not to participate

Researcher contacts student to arrange a baseline assessment

School counsellor gives support to student as per school policy

Student completes a 1 hour baseline assessment. They will also identify 3 therapeutic goals and complete a safety plan.

Independent statistician randomizes student to either the **intervention** or **control group**

The **intervention** group receive an email directing them to the RE-FRAME IT website and receive the 8-wk intervention plus 2x weekly SMS messages AND standard care.

The **control group** receive standard care plus a weekly SMS message enquiring about well-being.

Researcher completes a post-intervention assessment (8 weeks)

12 week follow up



Assessment schedule: Participants will be assessed at baseline and post intervention (8 weeks) using the following:

- Demographic information and treatment history will be collected using a specifically designed questionnaire
- The Suicidal Ideation Questionnaire Junior [16].
- The Self-Harm Behavior Questionnaire [17]
- The General and Actual Help Seeking Scales [18]
- The Children's Depression Rating Scale [19]
- The Reynolds Adolescent Depression Scale [20],
- The Multidimensional Anxiety Scale for Children [21, 22]
- The screening questions for Borderline Personality Disorder from the Structured Clinical Interview for DSM-IV Axis II Personality Disorders [23].
- The Beck Hopelessness Scale [24]
- The Coping Inventory for Stressful Situations [25]. These constructs are being measured in order to determine whether or not they impact upon levels of suicidality
- The Profile of Moods States [26] and a series of specific questions asking young people if they found the content unduly distressing.

The intervention: a personalized web-site which will deliver 8 modules of psychological therapy, employing a cognitive behavioural approach, over an 8-week period. The modules will cover the following:

- Week 1: Introduction / engagement and agenda setting
- Week 2: Emotional recognition and distress tolerance
- Week 3: Identification of automatic thinking
- Week 4: Behavioural activation: help-seeking
- Week 5: Behavioural activation: activity scheduling (including relaxation techniques)
- Week 6: Problem solving: specific focus on self harming or SI
- Week 7: Detecting & challenging problematic thinking and cognitive restructuring
- Week 8: Wrap up and review

Format: Each participant will have access to their own personalized web-page via a secure log-in process. The website will be hosted by a real person talking to the viewer. They will introduce the site and provide the therapeutic content. There will also be a series of characters (young adult actors) who will provide a video diary (vlog) each week describing their problems with schoolwork or their friendships or parental relationships. There are 2 male characters and 2 female characters, one each in years 9 to 12.

Each participant will also have a message board to communicate with the research team. There will also be an activity or a game (e.g. a mood diary) that will reinforce the learning objectives each week. This will be sent back to us via the site and it will be reviewed by a member of the research team who will moderate the website and provide individualized feedback to each participant.

SMS messages will also be sent twice a week for 8 weeks, the first just as the new module is uploaded and the second following their completion of the module. The control group will receive SMS messages enquiring about well-being.

Safety: The website will be moderated by a clinical psychologist and participants will be assessed weekly for risk.

Timeline

2010	Study development
2011 Jan – July	Piloting
Aug 2011 – Aug 2013	Recruitment (2 years) Deliver intervention (8 weeks) Baseline and post-intervention assessments
Sept 2013 – March 2014	Data verification, analysis and write up

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