Online parent training for reducing parenting stress during the coronavirus pandemic

Submission date	Recruitment status	Prospectively registered
15/05/2020	No longer recruiting	[_] Protocol
Registration date	Overall study status	[] Statistical analysis plan
23/05/2020 Last Edited	Completed Condition category	[_] Results
		Individual participant data
21/05/2020	Other	[] Record updated in last year

Plain English summary of protocol

Background and study aims

In January 2020, the World Health Organization (WHO) declared the outbreak of a new coronavirus disease, COVID-19, to be a Public Health Emergency of International Concern (WHO, 2020). In that time of crisis which generates stress throughout the population, supporting mental and psychosocial well-being is warranted. Internet-based programs have been identified as an alternative means of intervention delivery during the pandemic.

The study aims to evaluate the engagement in, satisfaction with, and efficacy of an internetbased parent-training program for reducing parenting stress during the Covid-19 pandemic.

Who can participate?

Parents of at least one child below the age of 18 years old, with internet access, own mobile phone or computer.

What does the study involve?

Participants will be randomly allocated to either complete the online training program without having any contact with other participants or the clinician who follows their completion, or in addition to the program an online forum will be added to the structure of the training, where participants can share their experiences and contact each other, and ask the clinician, or in addition to this, the clinician will give personal feedback to assignments submitted by the participants.

What are the possible benefits and risks of participating?

Possible benefits of participating will be that participants will receive information about potential mental health challenges of the pandemic and effective coping strategies that help to avoid the adverse psychological effects of the pandemic in parents and children. They can practice these skills during the training. An additional benefit is that if one of the participants is identified as needing further professional help, the moderator will contact him or her and help him or her get the appropriate help/treatment.

Potential risks: Thinking about the pandemic and raising awareness of their own and their children's stress reaction, worries and coping capacities may be anxiety provoking. In that case the participant can contact the moderator, and the moderator will help him or her get the appropriate help/treatment.

Where is the study run from? 1. Eötvös Loránd University (Hungary) 2. Heim Pál National Pediatric Institute (Hungary)

When is the study starting and how long is it expected to run for? April 2020 to December 2020

Who is funding the study? Eötvös Loránd University (Hungary)

Who is the main contact? Dr Monika Miklósi, miklosi.monika@ppk.elte.hu

Contact information

Type(s) Scientific

Contact name Dr Monika Miklósi

ORCID ID https://orcid.org/0000-0001-8316-0410

Contact details Eötvös Loránd University Izbella 46 Budapest Hungary 1064 +36 20 3948183 miklosi.monika@ppk.elte.hu

Additional identifiers

EudraCT/CTIS number Nil known

IRAS number

ClinicalTrials.gov number Nil known

Secondary identifying numbers 2020/185

Study information

Scientific Title

Online parent training for reducing parenting stress during the COVID-19 pandemic: a randomized controlled trial

Study objectives

The online training program will result in a significant reduction in parents' level of stress and a significant increase in well-being and parental self-efficacy

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 12/05/2020, Faculty of Education and Psychology, Research Ethics Committee of the Eötvös Loránd University (Izbella 46, Budapest 1064, Hungary; + (36-1) 461-2600 / 5614; molnar. mark@ppk.elte.hu), ref: 2020/185

Study design

Interventional randomized controlled study

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Internet/virtual

Study type(s) Prevention

Participant information sheet

In Hungarian: https://drive.google.com/open?id=12NHiQh8iFOInp_8fnQnfNA3lLk-2yNem

Health condition(s) or problem(s) studied

Parental stress during COVID-19 pandemic

Interventions

The study aims to examine the effect of an online parent training in reducing parental stress during Covid-19 pandemic and to determine the optimal design of the program by means of engagement in, satisfaction with, and efficacy of the program.

The training program consists of two modules, one focusing on parents' stress, and the second focusing on children's stress and parenting practices that are appropriate during the pandemic time. Each module consists of 5 topics with short psychoeducative videos and written materials, while quizzes, worksheets, and feedback forms are included for increasing engagement. Parents can flexibly go through the topics during two weeks, requiring a 15-30-minutes daily online activity.

To explore the optimal structure of the training, participants will be randomized into three conditions.

In group 1., parents complete the online training program without having any contact with other

participants or the clinician who follows their completion. The clinician contacts a parent only in case of emergency.

In group 2., an online forum will be added to the structure of the training, where participants can share their experiences and contact each other, and ask the clinician. The clinician will be involved in the forum and answer the questions.

In group 3., in addition to this, the clinician will give personal feedback to assignments submitted by the participants.

Block randomization with randomly selected block sizes of 6, 12 is used. The randomization scheme is generated by using the Web site Randomization.com (http://www.randomization.com

Intervention Type

Behavioural

Primary outcome measure

In the parent, according to self-report, measured in four time-points: at baseline, post-treatment, 1- and 3-months follow-up:

- 1. Perceived stress (Perceived Stress Scale, four-item version, PSS4)
- 2. Psychological well-being (WHO Well-being Index, WBI)
- 3. Parenting stress (Parental Stress Scale, shortened)
- 4. Parental competence (Parental Sense of Competence Scale, PSOC)

Secondary outcome measures

According to self-report, measured in four time-points: at baseline, post-treatment, 1- and 3months follow-up:

1. Parenting behaviour (Multidimensional Assessment of Parenting Behavior, MAPS) 2. Children's quality of life (Inventar zur Erfassung der Lebensqualitaet für Kindern und Jugendlichen, ILK)

Overall study start date

06/04/2020

Completion date

15/12/2020

Eligibility

Key inclusion criteria

1. Being a parent of at least one child below the age of 18 years old

2. Internet access, own mobile phone or computer

Though both parents can take part in the training, only data from one parent per family will be included in analyses

Participant type(s) Healthy volunteer

Age group Adult

Lower age limit

18 Years

Sex Both

Target number of participants 300

Key exclusion criteria Does not meet inclusion criteria

Date of first enrolment 18/05/2020

Date of final enrolment 30/11/2020

Locations

Countries of recruitment Hungary

Study participating centre Eötvös Loránd University Psychological Institute Izabella 46 Budapest Hungary 1064

Study participating centre Heim Pál National Pediatric Institute Faludi 5 Budapest Hungary 1138

Sponsor information

Organisation Heim Pál National Pediatric Institute LÉTRA Foundation

Sponsor details

Garas 9 Budapest Hungary 1026 +36 1 349-1514 mentalamb@heimpalkorhaz.hu

Sponsor type Hospital/treatment centre

Website http://www.letraalapitvany.hu/

Funder(s)

Funder type University/education

Funder Name Eötvös Loránd University

Results and Publications

Publication and dissemination plan Planned publication in a high-impact peer-reviewed journal.

Intention to publish date 15/01/2021

Individual participant data (IPD) sharing plan All data generated or analysed during this study will be included in the subsequent results publication.

IPD sharing plan summary Other