Sicher im Ernstfall (Safe in an emergency)

Submission date	Recruitment status No longer recruiting	Prospectively registered		
04/03/2019		☐ Protocol		
Registration date 07/03/2019	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited	Condition category	[] Individual participant data		
26/10/2020	Circulatory System			

Plain English summary of protocol

Background and study aims

In Germany it is recommended for school children to receive basic life support (BLS) training. Even trained people often hesitate to help in cardiac arrest. This study investigates if special training methods are effective at improving school children's preparedness and positive beliefs in their own capability to help in cardiac arrest.

Who can participate?

Students in grade seven and eight at high schools in Hamburg

What does the study involve?

Classes are randomly allocated to either the intervention group or the control group. All classes participate in practical training in BLS. The intervention group receive self-regulated training in peer groups after demonstration and deconstruction of the steps by the trainer. The control group receive Instructor-led training following the four-step approach. Self-efficacy for helping in cardiac arrest is measured immediately after the training and 9 months later.

What are the possible benefits and risks of participating? Participants receive training in first aid, and no risks are expected.

Where is the study run from?
University Medical Center Hamburg-Eppendorf (Germany)

When is the study starting and how long is it expected to run for? February 2015 to October 2017

Who is funding the study? University Medical Center Hamburg-Eppendorf (Germany)

Who is the main contact? Dr Stefanie Beck st.beck@uke.de

Contact information

Type(s)

Public

Contact name

Dr Stefanie Beck

Contact details

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

Nil known

Study information

Scientific Title

Promotes self-regulated learning in peer- groups self-efficacy for helping in cardiac arrest?

Acronym

Sicher im Ernstfall

Study objectives

Self-regulated learning in peer- groups promotes self-efficacy and long-term retention of practical Basic Life support (BLS) skills in school children.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The study was rated by the head of the local Ethics Committee of the Medical Association of Hamburg as a study with humans but not on humans. Therefore, this educational study does not refer to the statutes of the ethics committee (§ 9 des Hamburgischen Kammergesetzes für

Heilberufe) and the study was not appropriate for ethics consultation. This study was conducted with the approval of the Ministry of education of the state of Hamburg (Ministerium für Schule und Weiterbildung des Landes Hamburg).

Study design

Prospective randomised controlled interventional trial

Primary study design

Interventional

Secondary study design

Cluster randomised trial

Study setting(s)

Community

Study type(s)

Other

Participant information sheet

not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Helping in cardiac arrest

Interventions

Students were randomised class-wise into the intervention and control arm in a 1:1 allocation ratio. The randomisation was performed in advance by drawing balls of two colours blindly. A cluster was represented by students of one class.

All students participated in a practical training in BLS and an objective structured examination of skills:

Intervention: Basic Life support training with self-regulated training in peer-groups after demonstration and deconstruction of the steps by the trainer Control: Instructor-led basic life support training following the four-step approach

Intervention Type

Behavioural

Primary outcome measure

Self-efficacy of school children for helping in cardiac arrest, measured with a four-point Likert scale in three dimensions with two questions for each by adding the values of the two questions. The dimensions were helping in general, helping in cardiac arrest and diminished emotional arousal to cardiac arrest. The questionnaire, based on the general self-efficacy scale of Schwarzer and Jerusalem, was transformed into a special self-efficacy scale following the authors recommendations. Measured immediately after the training and 9 months later.

Secondary outcome measures

Pass-rates and means for items in a practical basis life support assessment. The assessment of the practical BLS skills was structured as an OSCE using MiniAnne-mannequins® (LaerdalTM).

For assessment, the raters used a structured rating checklist with nine binary items. To pass the practical assessment in total, all nine points had to be rated with yes. Measured immediately after the training and 9 months later.

Overall study start date

04/02/2015

Completion date

31/10/2017

Eligibility

Key inclusion criteria

- 1. Volunteering school children
- 2. Written informed consent of the parents

Participant type(s)

Healthy volunteer

Age group

Child

Sex

Both

Target number of participants

15 classes (clusters) with a mean number of 22 students per class for every group (15 x 22 x 2)

Total final enrolment

600

Key exclusion criteria

- 1. Not able to perform basic life support
- 2. No written informed consent from the parents

Date of first enrolment

01/07/2016

Date of final enrolment

01/09/2016

Locations

Countries of recruitment

Germany

Study participating centre

University Medical Center Hamburg-Eppendorf

Department of Anaesthesiology Martini-Str. 52 Hamburg Germany 20246

Sponsor information

Organisation

University Medical Center Hamburg-Eppendorf

Sponsor details

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Sponsor type

University/education

ROR

https://ror.org/01zgy1s35

Funder(s)

Funder type

Other

Funder Name

Investigator initiated and funded

Results and Publications

Publication and dissemination plan

Planned publication in an open access medical journal.

Intention to publish date

15/03/2019

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Dr Stefanie Beck (st.beck@uke.de). The final dataset used for analyses could be provided, data are anonymised, consent to publish anonymised result obtained by all participants.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	13/01/2020	26/10/2020	Yes	No