Video-based smartphone app (‘VIDEA bewegt’) for physical activity support in German adults: a study protocol for a single-armed observational study

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ABSTRACT

Introduction Insufficient physical activity is one of the most important risk factors for non-communicable diseases. Physical activity should therefore be intensively promoted in all age groups. Several trials suggest that it can be effectively increased through smartphone interventions. However, few of the smartphone-interventions available on the market have been scientifically evaluated. Therefore, the described study aims to assess the short-term and long-term effects of the smartphone intervention ‘VIDEA bewegt’ to increase physical activity. The trial is designed as a single-armed observational trial to assess effects under real-life conditions.

Methods and analysis The intervention consists of the smartphone-application ‘VIDEA bewegt’, which is a video-based preventative programme to improve physical activity in everyday life. The application contains several features and components including educational videos, documentation of activity and motivational exercises. A sample size of at least 106 participants is aimed for. The primary objective of this study is to determine the effect of the application on physical activity in German adults. Secondary objectives are to evaluate the self-efficacy, health-related quality of life and usability of ‘VIDEA bewegt’.

Data collection is based on online questionnaires, as well as system-internal recorded data. Changes of outcomes from baseline to programme completion and follow-up will be calculated.

Ethics and dissemination The Ethics Committee of the Technical University Dresden approved the study on 25 May 2019 (EK 272062019). All data are processed anonymously and stored on servers only accessible by authorised personnel. The results of the study and the results of the usability test are aimed to be published in a scientific journal.

Trial registration number German Clinical Trials Registry (DRKS00017392).

INTRODUCTION

Background Insufficient physical activity is considered to be one of the most important risk factors for non-communicable diseases along with tobacco, alcohol consumption and unhealthy diet.1 Being defined as less than 150 min of moderate activity, 75 min of vigorous activity or less than 600 metabolic equivalent (MET) minutes per week,2 its prevalence for both sexes is 27.5% worldwide and 42.2% in Germany.3 As activity progressively decreases with age, it is likely to become a growing problem in our ageing population.4 In order to prevent chronic diseases and to minimise the risk of premature death, physical activity should therefore be intensively promoted in all age groups.5–7

The widespread availability of smartphones has increased the interest in exploring the potential of digitisation in preventative healthcare.8 Digital interventions can be made available without great cost and effort9 and could help to provide healthcare for those with limited access to it.10 The potential of such interventions is demonstrated by the fact that more than 80% of German citizens aged 14 or older, and already more than 40% of those aged 65 or older, were using a smartphone regularly in 2017.11 Focusing on physical activity, several scientific trials suggest that it can be effectively increased through smartphone interventions, which provide information and performance-related feedback.
monitor behaviour change, enable social networking and goal setting and include motivational messages or reminders.12–15 Although the number of health apps on the market is already high (in 2016 >80 000 apps in categories ‘health and fitness’ and ‘medicine’16), and their potential in the health sector is promising,17–19 only few of these apps use evidence-based content, practices and features.19 Developing digital interventions that work equally well for people with different expectations, opportunities and cultural backgrounds is challenging.20

A major limitation of past studies is the fact that they use apps which are specifically developed for the purpose of the study and not commercially available.21–22 As a consequence, only few of the apps available on the market have been scientifically evaluated,23–24 although an evaluation could benefit their clinical use.25 Various methodological challenges of evaluating digital devices using randomised controlled trials are known. These challenges include randomisation, timing of assessment, acceptance by patients and physicians, blind as well as defining control groups and relevant endpoints.26 Apart from that, many digital health interventions combine different intervention components (tracking of health data, motivating messages based on individual goals, performance-related feedback, etc) and can therefore differ in their effects on individuals—despite identical utilisation. This modular structure of digital applications and the associated flexibility and customisability are great strengths of those interventions,27 but make it difficult to conduct randomised controlled trials, as recommended by several studies.28–29

There is a need for mid-term and long-term data on the effectiveness and safety of such dynamic smartphone-applications.28–29 In addition, the time course of intervention effects and the strategies needed to achieve sustainable change have not yet been fully investigated.12 An iterative and user-centred development is regarded as key to identify needs and preferences of the relevant target group,30–33 and overcome barriers of prototypes in early development stages.34

Objective

The overall objective of this study is to assess the effect of the video-based smartphone app ‘VIDEA bewegt’ on physical activity in German adults.

Secondary objectives are to evaluate self-efficacy, health-related quality of life and usability of ‘VIDEA bewegt’, especially with regards to participants’ satisfaction with the app.

Hypotheses

► The users of ‘VIDEA bewegt’ who participate in the study increase their average daily step count and achieve a higher number of MET minutes per week, a significantly higher health-related quality of life, a significantly higher motivational, maintenance and recovery self-efficacy after the first 4 weeks and after completion of the 8-week course, compared with the beginning of data collection.
► The above-mentioned changes will last through the follow-up period (2/4/6/12 months after the completion of the programme). The strength of the effects is influenced by individual background (eg, educational background, socioeconomic status) and by the participants’ use of the app.

Trial design

The evaluation of the app ‘VIDEA bewegt’ is designed as a single-arm observational study, assessing the app’s effectiveness under real-life conditions. Additionally, think aloud interviews are conducted beforehand within the framework of a usability test to integrate user-oriented design.

METHODS: PARTICIPANTS, INTERVENTIONS AND OUTCOMES

Study dates

The study was developed from October 2018 to April 2019. The ethics proposal and the prestudy with usability and pretest were carried out from January 2019 to May 2019. Data collection started in July 2019. Depending on sufficient numbers of participants, it is expected to be continued until December 2019 for a first evaluation of the effects during programme use. The follow-up is planned to be continued until the end of 2020.

Study setting

‘VIDEA bewegt’ is an app enabling users to access educational content via their smartphone anywhere and at any time. ‘VIDEA bewegt’ was officially certified as a physical activity course and is covered by some health insurances. It was made available on the German market for Android and iOS in March 2019. More information can be found on the German website.35

A clinical visit or in-person contact with a physician or diabetes specialist is not required. However, it is possible to consult experts in preventative healthcare and sports science via an integrated chat function. Problems can also be discussed with other users in a forum.

As the developer of ‘VIDEA bewegt’, the TUMAINI Institut für Präventionsmanagement GmbH is responsible for the overall functioning of the app.

Eligibility criteria

All users of ‘VIDEA bewegt’ are potential participants of the study and will be invited to take part in the evaluation after registration in the app. Inclusion criteria for participation in the study are the active use of ‘VIDEA bewegt’ and informed consent to participate in the study (figure 1). In order to complete the app-registration, users must be of legal age (≥18 years old), agree to the terms of the app, confirm that physical contraindications such as musculoskeletal conditions, cardiovascular diseases and cancer do not apply, and speak and write German fluently. Inclusion and exclusion are not restricted by

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It consists of eight 45 min stages, each representing one course week including four videos (online supplementary figure 3 – stage structure). Each week, a new stage will be unlocked in ascending order. Therefore, the minimum time needed to complete the programme is 7 weeks and 1 day. The maximum period for completing the programme is 1 year.

App users are free to decide when and how intensively they want to use the programme. The app has a modular structure and contains the following features and components:

- Educational videos.
- Synchronising steps.
- Calculating and displaying activity minutes.
- Motivational exercises under ‘My Focus’.
- Motivational messages.
- Answering quiz questions.
- Chat/forum.

**Educational videos**

In these videos, an expert in preventative healthcare and an expert in sports science provide both theoretical medical content and practical exercises to participate in. The theoretical videos have a length of 2–4 min and the exercise videos each last approximately 20 min. In total there are 32 videos and 17 additional videos. In the first video of each stage, users receive theoretical information regarding physical activity and its effects on body and mind. For example, the potential negative effects of a physically inactive lifestyle are explained together with strategies for their prevention. The second video presents exercise instructions for endurance and strength exercises. The aim of these videos is for users to get to know the exercises so that they can perform them on their own. The third video refers to the theoretical change model of Greaves et al. to build up motivation to increase activity through techniques of motivational psychology applied practically in the category ‘My Focus’. The fourth video contains practical recommendations on how exercise can be integrated into everyday life with the help of different types of exercise. Add-on videos will be made available on topics like barefoot walking, sauna visiting and healthy diet.

**Synchronising steps**

In order to track activity, the app offers the opportunity to synchronise step numbers on a daily basis (online supplementary figure 4 – Activity). Users can either enter the step numbers into the app themselves, if they are using an external device, or synchronise their steps automatically. For automatic synchronisation, ‘VIDEA bewegt’ is connected with a pedometer app on the phone.

**Calculating and displaying activity minutes**

When registering, users select an average activity level of <30 min, 30–60 min and >60 min/day to set a daily activity goal (online supplementary figure 4). Physical activities such as cycling, swimming, housework can

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**Interventions**

‘VIDEA bewegt’ is a preventative programme to improve physical activity in everyday life and strength endurance. It was developed in cooperation with physicians of the university hospital Dresden. The programme uses videos to present content and exercises and aims to motivate users to increase physical activity on a regular basis (see online supplementary figures 1–5).

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**Patient and public involvement**

Potential participants were included in the study development in order to make methods understandable and functional, to better understand observed effects and to gather errors and problems to serve future projects. One step in this process involved pretesting the questionnaires in order to assess their logic and understandability. Another important step was the usability test which included interviews with selected individuals. Possibilities to improve the app could be demonstrated.

In the context of actual app usage and study participation, participants are involved in system improvement by recording app usage problems in questionnaires.

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**Evaluation “VIDEA bewegt”**

Einen Studie der Universitätsklinikum Carl Gustav Carus Dresden

Herrlich willkommen!

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Der Umgang mit Ihren Daten wird gemäß der Datenschutz-Grundverordnung (DSGVO) und dem deutschen Bundesdatenschutzgesetz (BDSG) gewährleistet. Ihre Daten werden nur für den Zweck der Studie gespeichert und sind nicht für Dritte unterwegs.

Sollten Sie Fragen haben, können Sie bei uns telefonisch oder per E-Mail erreicht werden.

In der vorliegenden Studie werden folgende Arzneimittel eingesetzt:

- Paracetamol
- Ibuprofen

Bitte beachten Sie, dass die Arzneimittel nur in Tablettenform angeboten werden.

Sollte die Medikamenteneinnahme nicht möglich sein, können Sie uns im Vorfeld anrufen, um eine Ausnahme zu beantragen.

Die Studie ist in Übereinstimmung mit der Deklaration von Helsinki 2013 durchgeführt worden.

Bitte lesen Sie daher die zweiseitigen Informationen in Absprache mit Ihrem Arzt/Arztin gegeben.

Zusätzlich zu den medizinischen Beratungsleistungen werden Ihnen auch praktische Anleitungen zur Gesundheits- und Bewegungserhöhung gegeben.

Die Informationsmaterialien sind im Folgenden aufgeführt:

- Broschüre: ‘VIDEA bewegt’
- Infotafel: ‘VIDEA bewegt’
- Website: www.videa-bewegt.de

Die Teilnehmer/-innen erhalten ein persönlichestagsbuch, in dem sie ihre täglichen Aktivitäten aufschreiben können. Die Daten werden anonymisiert und nur für die Auswertung der Studie verwendet.

Die Teilnehmer/-innen erhalten eine individuelle Gesundheitsberatung von einem Arzt/Arztin oder einer Ärztin/Arztin.

In der Studie werden folgende Maßnahmen zur Motivation der Teilnehmer/-innen eingesetzt:

- Freiwillige Teilnahme
- Peer- und Selbsthilfe
- Gruppenarbeit
- Individualberatung

Die Daten der Teilnehmer/-innen sind nach ihrer Genehmigung der Datenschutzbehörde gespeichert.

Die Studie wird von der Ethikkommission der Technischen Universität Dresden (EKTU) und der Ethikkommission der Universitätsklinikum Carl Gustav Carus Dresden (EKU) überwacht.

Die Ethikkommission der EKTU hat die Studie genehmigt.

Die Teilnehmer/-innen sind aufgefordert, innerhalb der angegebenen Studiendauer aktiv zu bleiben.

Die Teilnehmer/-innen erhalten eine entsprechende Prämie für ihre Teilnahme an der Studie.

Die Prämie beträgt 20 Euro pro Teilnehmer/-in.

Die Prämie wird innerhalb von drei Monaten nach Abschluss der Studie ausgezahlt.

Die Prämie wird nur innerhalb der angegebenen Studiendauer ausgezahlt.

Die Prämie wird nur an Teilnehmer/-innen gezahlt, die zu Beginn der Studie eine Einwilligungsschreiben haben ausgefüllt.

Die Teilnehmer/-innen haben das Recht, jederzeit aus der Studie auszutreten.

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Table 1: Overview of the measures used

| Measure               | Topic                      | Description                                                                                                                                                                                                 | Item number |
|-----------------------|----------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------|
| GPAQ                  | Physical activity          | The Global Physical Activity Questionnaire (GPAQ) measures the intensity and duration of physical activity at work, during transport and during leisure time. The data are quantified by a conversion into metabolic equivalent (MET) minutes per week. One MET represents the energy consumption at rest. The GPAQ defines 4 METs for moderate activity and 8 METs for vigorous activity. | 16          |
| AppEx (self-constructed) | General information       | Author-constructed questions about sociodemography, individual backgrounds, self-assessment of endurance and muscle strength.                                                                                             | 12          |
| SF-8                  | Health related quality of life | The SF-8 (Short Form Health Survey 8) covers the eight dimensions of SF-36 on the basis of eight questions: physical functioning, role-physical, bodily pain, general health, vitality, social functioning, role-emotional and mental health. | 8           |
| MotSE, adapted to     | Motivational self-efficacy | Self-efficacy related to the motivation to increase physical activity is estimated on the basis of three items.                                                                                                | 3           |
| MainSE, adapted to    | Maintenance self-efficacy  | Self-efficacy, related to the maintenance of increased everyday activity, is estimated on the basis of three items.                                                                                           | 3           |
| RecSE, adapted to     | Recovery self-efficacy     | Self-efficacy, related to the return to activity after a setback, is estimated on the basis of three items.                                                                                                   | 3           |
| AppEx-Mod (self-constructed) | General information         | Shortened version of the App-External Questionnaire (AppEx): questions about weight, height, endurance and muscle strength, participation in further courses.                                                          | 5           |
| Peva-FB UEQ           | Process evaluation         | A series of author-constructed questions to assess satisfaction with individual components of the programme and user behaviour. Additional integration of UEQ-S (User Experience Questionnaire Short), which assesses the user-friendliness of a product using eight opposing pairs of characteristics. | 5           |
| NH (self-constructed) | Sustainability             | Author-constructed questions to assess the use of the exercises and components of the programme after it has been completed.                                                                              | 5           |

be documented by selecting from a list and entering a number of minutes. In addition, the collected step numbers and training sessions contribute to the activity goal.

Motivational exercises
Based on contents of the third video of each stage, a new section in the app category ‘My Focus’ will be unlocked each course week, enabling users to express their thoughts in text fields for self-reflection and motivation. Sections of ‘My Focus’ include the user’s personal motive, possible action plans, power sources and strengths, social support, rewards, stopping negative thoughts, if-then plans and dealing with setbacks.

Motivational messages
Motivational messages appear each time the app is opened. These messages contain a review of the previous video, the reminder for a specific motivational exercise under ‘My Focus’ and a preview of the upcoming content. As such, the motivational messages are individualised using the aspects of the motivational exercises, which were initially selected by the patients themselves.

Answering quiz questions
At the end of each stage, two to three multiple-choice questions will be displayed on the content of the videos, primarily to review recently received information. Feedback is provided on given answers, yet accuracy does not influence the further programme.

Chat/forum
The ‘Exchange’ app category offers two features to users with questions or problems (online supplementary figure 5). Experts can be contacted directly in a chat. Furthermore, a forum is included to discuss questions on the topics ‘Questions and discussions about the course’, ‘Smalltalk’, ‘Technology and support’ with other users.

Measurement of outcomes
Primary outcome: physical activity
► Step numbers/minutes of activity.
► Mean MET minutes per week.

Secondary outcomes
► Self-efficacy (motivational, maintenance, recovery self-efficacy).
Health-related quality of life: Mental Health Component Summary score and Physical Health Component Summary score.

User behaviour and obstacles to the use of the programme.

See table 1 for detailed information.

**Participant timeline**

The evaluation will begin with people registering in the ‘VIDEA bewegt’ app without knowing anything about this study (figure 2). Once registration is complete, all users of the app will be invited to participate in the study via a pop-up window. Further information as well as the link to the first questionnaire will be sent by e-mail. Data are collected using a total of seven online questionnaires (figure 3). After completion of stage four and stage eight, selected system-internal data, will also be extracted from ‘VIDEA bewegt’.

**Sample size**

In similar projects, which were conducted as randomised controlled trials, the total number of participants ranged from 40 to 361.9 15 37 38

In the SMART MOVE study, for example, 90 participants were divided into two groups.15 After an a priori power analysis using G*Power39 with an assumed small to medium effect size for change of self-efficacy as well as for change of physical activity40–42 a necessary sample size of at least 27 was calculated for the t-tests. Starting from 20% study or intervention drop-out (conservative assumption), a total of n=33 participants will have to be recruited. In order to assess the influence of potential confounders such as age or period of use, a multiple regression analysis is planned. Therefore, a larger sample size of at least 106 persons is aimed for.43 44

**Recruitment**

Users of ‘VIDEA bewegt’ will be invited to the study after registration via a pop-up-window. Since two health insurance companies were involved in the development of the app, people insured with these health insurance companies can use the app free of charge. For members of other health insurance companies, a fee is due. In order to recruit further study participants, a fee waiver will be offered on the condition of participation.

Study participation will be compensated with €15. Consent to participate in the study will be given at the beginning of the first questionnaire (figure 1). Participants will receive emails containing a link to access each questionnaire at the relevant stage. If no reply was received within 2 days, a reminder e-mail will be sent.

**METHODS: DATA COLLECTION, MANAGEMENT AND ANALYSIS**

**Usability test**

A usability test was carried out before starting data collection. For this usability test, the think aloud method was used, which is common when developing user centred designs.45 Think aloud describes an interview in which, for example, an app is tested, and the test persons are asked to express their thoughts and impressions, particularly concerning obstacles they encounter when using the app, verbally. This approach provides an insight into strengths and weaknesses of a system.46 Think-aloud methods are a robust tool to identify about three quarters of the usability problems with the help of only four to five test persons.47
Ten test persons were recruited to work through the structure of the app while aiming to achieve various specified goals (eg, input their daily step count). They were asked to describe their thoughts, impressions and problems verbally. The audio track was recorded along with the screen of the smartphone. The recorded interview was transcribed and analysed according to Mayring.48 These insights in the usability of ‘VIDEA bewegt’ were used to further improve the programme’s usability. Thereby, the effectiveness of ‘VIDEA bewegt’ will be less likely to be confounded by usability problems.

Pretest of questionnaires

The questionnaires used in the evaluation contain both validated and non-validated questions constructed by the authors. The questionnaires were analysed with the help of expert opinions regarding their structure and selectivity. The non-validated questions were tested in a think aloud test and optimised on the basis of these results.

Apart from that, a pretest of the complete questionnaires was conducted by sending a link to a test questionnaire to 20 people. They were asked to record the time needed to answer the questionnaire and to comment on spelling, grammar, comprehensibility, presentation, structure and logic of the questions.

Evaluation

The actual study is based on two essential data sets. (1) Data gathered via the online questionnaires is stored in SURVEY (based on Limesurvey) and (2) system-internal data using step numbers and activity minutes (see table 2).

The questionnaires use internationally established scores for physical activity, health-related quality of life and self-efficacy (see table 1 and online supplementary file).

Plans to promote participant retention and complete follow-up

Subjects who withdraw their study participation will receive a final questionnaire for process evaluation. Data from respondents who withdraw or stop completing questionnaires will be included as long as questionnaires were completed at least at two different stages.

| Table 2 System-internal data |
|-----------------------------|
| **Email address** |
| **Day of registration** |
| **Day of completion** |
| **Health insurance company** |
| **User behaviour** |
| **Date/time at which a video was viewed** |
| **Active use of expert chat** |
| **Active use of forum** |
| **Documented data** |
| **Activity level** |
| **<30 min** |
| **30–60 min** |
| **>60 min** |
| **Pedometer—manual or automatic synchronisation?** |
| **Number of steps per day** |
| **Number of activity minutes reached per day** |
| **Activities selected on a given day** |

Data management

All collected primary data will be processed under strict confidentiality and will not be made accessible to third parties. Questionnaire data are collected via SURVEY (https://bildungsportal.sachsen.de/umfragen/). This service is based on Limesurvey and is provided by BPS Bildungsportal Sachsen GmbH. When using it, the following privacy policy applies (https://bildungsportal.sachsen.de/umfragen/datenschutz.html). The app data of ‘VIDEA bewegt’ is stored on the servers of TUMAINI. Members of the research team will have access to both data sets. In order to be able to contact app users, their email addresses must first be transferred from ‘VIDEA bewegt’ to a participant table in SURVEY. At both points of data analysis (T2 and T6) app-internal data will be transferred. This allows for the connection and anonymisation of both data sets. Data collection, coding and database cleaning is performed by at least two researchers (authors). A plausibility check is performed by descriptive univariate analysis. A university email address is provided to participants in case of questions or problems. A data sharing plan was added to the trial registration.

Statistical methods

Sociodemographic data and user behaviour will be analysed descriptively.

The effect on outcomes is calculated using a two-sided t-test for dependent samples at T1 to T6. The t-test requires metric, normally distributed variables.49 The Kolmogorov-Smirnov test will be used to test for normal distribution. In case of normal distribution, the connected t-test will be used, if there is no normal distribution the Wilcoxon sign rank test. In each case, the CI (95%) is tested with p=0.05 (see table 3). The possible alpha-error inflation of multiple mean value comparisons is controlled by Bonferroni correction. In addition, a power analysis using G*Power39 is performed, the CI for significance estimation is determined and the Bayesian correction is applied.50

There is no agreed definition of a general clinical relevance margin for increase of daily step count, as these improvement rates differ for certain patient characteristics (like sex, diseases and the baseline level of physical activity).51 In earlier trials, the clinical importance of change in step count (as one of the primary outcomes in our study) was considered as 600 and 110052 53 and above54 55 in different chronic diseases.

In the light of our relatively broad inclusion criteria, effects on primary and secondary outcomes will be interpreted with caution and in strong accordance to the enrolled patient profiles.

Methods for any additional analyses (e.g. subgroup and adjusted analyses)

A multiple regression analysis (possibly multilevel models) is planned. Predictors are age, gender, body mass index, baseline activity, participation in further lifestyle interventions.
Table 3: Statistical methods

| Outcome                                                                 | Hypothesis                                                                 | Statistical analysis                                      |
|-------------------------------------------------------------------------|---------------------------------------------------------------------------|-----------------------------------------------------------|
| **Primary outcome**                                                     |                                                                           |                                                           |
| Average number of steps/activity-minutes per day                         | Increase within first 4 weeks and until programme completion compared with baseline | t-test for combined samples (or Wilcoxon test)            |
| **Secondary outcomes**                                                  |                                                                           |                                                           |
| Average MET-minutes per week                                            | Increase within first 4 weeks and until programme completion compared with baseline, maintained over 2/4/6/12 months | t-test (or Wilcoxon-test)                                 |
| Self-efficacy (motivational, maintaining, restoring self-efficacy)      | Increase within first 4 weeks and until programme completion compared with baseline, maintained over 2/4/6/12 months | t-test (or Wilcoxon-test)                                 |
| Self-assessment endurance, muscle strength                              | Increase within first 4 weeks and until programme completion compared with baseline, maintained over 2/4/6/12 months | t-test (or Wilcoxon-test)                                 |
| Health-related quality of life (SF-8)                                    | Increase within first 4 weeks and until programme completion compared with baseline, maintained over 2/4/6/12 months | t-test (or Wilcoxon-test)                                 |
| User behaviour (Peva-FB after completion of the programme, or in case of dropout, additional continuously recorded internal system data (see table 1)) |                                                                           | Descriptive analysis with frequencies, mean values, SD    |
| **Subgroup analysis**                                                   |                                                                           |                                                           |
| (eg, age, gender, BMI, baseline activity, participation in additional lifestyle interventions) |                                                                           | Multiple linear regression (multilevel model, if applicable) |

BMI, body mass index; MET, metabolic equivalent; Peva, process evaluation; SF-8, Short Form Health Survey 8.

Auditing
A pretest of the questionnaires took place before starting data collection.
Additionally, system-internal data collection is checked using test accounts.

DISCUSSION
The main aim of the described study is to assess the effectiveness of the video-based smartphone app ‘VIDEA bewegt’ for increasing physical activity in German adults under real-life conditions. To the best of our knowledge, ‘VIDEA bewegt’ is one of the first apps on the German market to offer a primary preventative course to increase physical activity being developed under the guidance of physicians and being certified and covered by health insurances. Although the potential of digital prevention programmes is very promising,9 17 18 very few apps available on the market have yet been scientifically evaluated.23 In order to increase the clinical benefit of digital interventions, scientific analysis is urgently needed.31 35

Data are collected using various methods. The app system provides synchronised step numbers and activity minutes while established questionnaires allow for further data collection.

Since the acceptance of technical innovations substantially depends on ease of use and perceived usefulness,36 a usability test was carried out as the basis of this study. Previous studies have shown the importance of usability testing for successful app development.37–39 The think-aloud method has proven to be a useful way to gain insight into the thought processes and problems of test persons, especially in digital health interventions.31 58 In order to apply the results of the usability test, a process evaluation is included in the study, in which usage data of the app, as well as obstacles during app use, will be evaluated.

The development of the study protocol followed important recommendations formulated by Eysenbach and the Consolidated Standards of Reporting Trials-EHEALTH Group.60 As such, we describe the development of the video-based app and how to access it and provide detailed information on the intervention, including the mode of delivery, features/functionalities/components and prompts/reminders used.

In summary, the described study aims to contribute to exploring the potential of digital interventions for effectively increasing physical activity.

Limitations and strengths
The described study design contains a series of limitations.
First of all, there is a risk of selection bias. Since participation in the evaluation is voluntary, it is reasonable to assume that people who are already highly motivated to increase their level of physical activity are going to show more interest in participating than users who are struggling with their awareness and motivation. Additionally, it is planned to offer financial incentive to participate in the study by waiving the otherwise applicable course fee, which may bias the results. As both healthy subjects, as well as individuals at risk for and those already living with chronic disease are addressed, inclusion criteria are broad. This may result in an inhomogeneous group of study participants, which is also influenced by characteristics like age, frequency/intensity of smartphone use or level of health literacy.

Due to the multilayered structure of the intervention with several components, the actual intervention will differ for each individual (risk of performance bias). This study can only collect limited data on how intensively the individual components of the app have been used and how they contributed to the overall effect.

The app is regularly updated by the responsible company. The evaluation team does not have any influence on these updates but will be informed via email in case an update has been conducted. However, these updates mostly intend to eliminate technical problems and do not lead to major changes of the overall intervention.

Most of the data collected is based on self-assessment, there is no possibility of verification. The study is not conducted as a randomised controlled trial, there will be no initial trial visit and no in-person contact during the intervention. For this reason, no blinding of the participants and the study personnel is intended. The design of the study resembles a pragmatic study. Due to the pragmatic approach measuring precise baseline data are difficult. Since the users will need to register in the app before participating in the study, the participants’ level of knowledge may differ by the time the first questionnaire is answered.

These limitations are countered by the strength of the described study to evaluate the effectiveness of the app under the most realistic conditions possible. The project includes several different approaches, such as a usability test, questionnaires, system data and a follow-up to assess sustainability. The aim is to create a comprehensive impression of the app, its weaknesses, strengths, and potential.

ETHICS AND DISSEMINATION
Ethics approval
After an ethics application was submitted to the Ethics Committee of the Technical University of Dresden on 12 April, the Ethics Committee approved the study on 25 May 2019 (EK 272062019).

Protocol amendments
If changes to the procedure are necessary, a report will be made to the ethics committee of the Technical University of Dresden as well as to the study participants.

Consent to participate
The informed consent to participate is obtained written by clicking a button in the first online questionnaire in accordance with the General Data Protection Regulation (GDPR) as a prerequisite for participation (figure 1).

Dissemination policy
The results of the study are planned to be published in a scientific journal.

Contributors TF, GR and PS conceived and designed the study protocol, wrote the manuscript, will collect the data and perform the data analysis. PT regularly provided feedback on the overall study protocol, was involved in the development of the usability test, and participated in the writing of the manuscript. PEHS advised and provided feedback on the overall protocol and reviewed the manuscript. All authors reviewed and approved the final version of the manuscript before submission.

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Competing interests The principal investigator PEHS was involved in the development and implementation of the app ‘VIDEA bewegt’ as a medical expert. He is responsible for the medical and theoretical background and is shown in the app’s videos. He received no payment for his participation in the app.

Patient consent for publication Not required.

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