Mobile health-based gamification intervention to increase physical activity participation among patients with coronary heart disease: study protocol of a randomised controlled trial

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ABSTRACT

Introduction Despite proven benefits, physical activity participation remains low in patients with coronary heart disease (CHD). Scientific evidence suggests that mobile health (mHealth)-based gamification interventions could increase physical activity levels. However, several systematic reviews demonstrated that most gamification intervention designs do not appropriately leverage theories from health behaviour models, and empirical evidence on the efficacy of such interventions among patients with CHD is still emerging. This study embeds the principles of behavioural economics into a gamification intervention based on a smartphone app (WeChat applet) to explore whether a mHealth-based gamification intervention can improve participation in physical activity and other related physical and psychological outcomes in patients with CHD.

Methods We propose a single-blinded three-arm randomised controlled trial with 108 patients with CHD, who will be randomly divided into three groups (Control group: WeChat applet+step goal setting; Individual group: WeChat applet+step goal setting+gamification; Team group: WeChat applet+step goal setting+gamification+collaboration). The interventions will last for 12 weeks and follow-up for 12 weeks. All patients will receive only WeChat applet-based step goal setting in the follow-up period. The primary outcome is physical activity participation, which includes a change in daily steps and self-reported physical activity from the baseline to 12 and 24 weeks, and the proportion of patient-days that step goals achieved in 12 and 24 weeks. The secondary outcomes include biomedical and lifestyle-related risk factors, intrinsic motivation, enjoyment, competence, autonomy and relatedness, social support and mental health and patients’ satisfaction, perceptions and intervention experience.

BACKGROUND

Coronary heart disease (CHD) is the leading cause of mortality in China. Statistically, around 11 million people were affected with CHD in 2017.1,2 Exercise-based cardiac rehabilitation and secondary prevention (CR/SP) plays a crucial role in preventing the recurrence of CHD3 and has been listed as a class I recommendation for CHD treatment by the American Heart Association, the American Society of Cardiology and the European Society of Cardiology.4-7 The relevant guidelines recommend that patients with CHD should perform at least 500 Metabolic Equivalent (MET)-min/week physical activity every week.8 Although CR/SP have proven benefits, it is often challenging for patients to attain lifestyle changes needed for SP, especially with increasing physical activity...
levels. For example, owing to the poor accessibility of CR programmes, >80% of patients did not participate in the CR programmes recommended by the guidelines. Moreover, patients with CHD typically fail to attain their daily physical activity goals.

Mobile health (mHealth), defined by American Heart Association’s scientific statement, ‘the use of mobile computing and communication technologies (eg, mobile phones, wearable devices) for health services and information’, has become an essential medium to deliver behavioural change interventions and demonstrated promising ability to improve physical activity levels. For example, the Consumer Navigation of Electronic Cardiovascular Tools (CONNECT) trial examined the impact of digital health interventions on health behaviours and established the correlation between intervention and increased attainment of physical activity targets. In China, WeChat is a top-rated multipurpose social media app, with >1.151 billion active users. WeChat applets are lightweight apps that form part of the WeChat ecosystem, which could be used independently and do not need installation. Compared with mobile apps, WeChat applets are easier to be accepted and applied by people in China. The third quarter of 2019 recorded >300 million active WeChat applet users every day, thereby making it well suited to disseminate mHealth interventions in China.

Gamification is the use of game design elements (such as points, leaderboards, progress bars and badges) in non-game contexts (such as management, education, marketing and healthcare) to increase motivation and engagement. There is growing interest in the application of gamification in mHealth with the view of promoting healthy behavioural changes, especially in promoting physical activity levels. Previous studies indicated that gamification was used in 64% of the top 50 most popular smartphone apps. However, several systematic reviews reported that most gamification intervention designs did not appropriately leverage theories from health behaviour models. Moreover, as the concept of gamification is relatively new, empirical evidence on the efficacy of gamification physical activity behavioural change interventions among patients with CHD is still emerging.

Gamification interventions are rarely based on a sound theoretical framework. Behavioural economics principles combine conventional economic principles with psychology to elucidate how individuals behave and make decisions. Behavioural economics principles can be embedded with a gamification intervention via mobile devices to aid people to attain their physical activity goals. For example, based on the loss aversion, which implies that the loss framework is more effective in stimulating behavioural change than the gain framework, Patel et al designed an intervention wherein participants lost points if they did not accomplish their step goals. Several previous studies have used behavioural economics principles to help patients lose weight, quit smoking and adhere to medications. However, limited data are available on applying these concepts to improve physical activity participation in patients with CHD.

This study will use behavioural economics principles to develop a gamification WeChat applet named ‘TahneeWeh’ to resolve the research gap mentioned above. This study aims to investigate the effects of the mHealth-based gamification intervention on participation in physical activity and evaluate the effects on biomedical and lifestyle-related risk factors, intrinsic motivation, enjoyment, competence, autonomy, and relatedness, social support, and mental health. In addition, a semi-structured interview will be conducted after the intervention to comprehend patients’ satisfaction, perceptions and their experience on the intervention.

**METHODS**

**Study design**

This is a single-blind, three-arm randomised controlled trial to evaluate the effects of the mHealth-based gamification intervention on participation in physical activity, biomedical and lifestyle-related risk factors, intrinsic motivation, enjoyment, competence, autonomy, and relatedness, social support, and mental health. Patients with CHD will be recruited in a CR centre of a tertiary-grade A class hospital in Changchun (China) through posters and email of discharged patients. A total of 108 participants will be randomly divided into three groups (Control group: WeChat applet+step goal setting; Individual group: WeChat applet+step goal setting+gamification; Team group: WeChat applet+step goal setting+gamification+collaboration). Patients in the control group will only receive daily step goal setting. The Individual and Team groups will receive gamified behaviour intervention based on behavioural economics principles. The Team group will also receive social incentives based on the Individual group. The intervention will last for 12 weeks and follow-up for 12 weeks. All patients will just receive WeChat applet-based step goal setting in the follow-up period. The study duration will be between 1 July 2021 and 30 November 2022. Figure 1 shows a flow chart of the study design. The protocol conforms to the Standard Protocol Items: Recommendations for Interventional Trials reporting guidelines, and the intervention is described per the Consolidated Standards of Reporting Trials of Electronic and Mobile HEalth Applications and onLine TeleHealth (CONSORT-EHEAL) checklist.

**Eligibility and recruitment**

Patients fulfilling the inclusion and exclusion criteria will be invited to participate in the trial. The inclusion criteria are as follows: (1) aged 18–70 years; (2) patients diagnosed with CHD (including acute myocardial infarction and unstable angina) and received percutaneous coronary intervention (PCI) treatment during admission; (3) patients evaluated by cardiologists and rehabilitation therapists if they are suitable for participating in our programme; (4) patients willing to provide written consent.
informed consent; (5) patients with a smartphone and an active WeChat account; and (6) patients with proficiency in Chinese.

The exclusion criteria include the following: (1) contraindications for exercise rehabilitation (e.g., untreated ventricular tachycardia, severe heart failure, uncontrolled hypertension or hypotension, notable exercise restriction); (2) patients unable to use WeChat applet after instruction; (3) no internet access in the place of residence; (4) patients requiring a walking aid to move and (5) patients participating in other clinical trials.

Experienced clinical nurses, rehabilitation therapists and researchers will be responsible for recruiting participants. Patients undergoing PCI will be referred to the CR centre, which is adjacent to the wards to receive physical activity counselling and obtain the follow-up booklet before discharge. The follow-up booklet will remind patients to return for review in 4 weeks, 12 weeks, 24 weeks and 12 months after discharge, which is very helpful to avoid patients’ lost to follow-up. In the rehabilitation centre, researchers will screen the eligible patients and then inform the patients about the details of the study. If they agree to participate in the study, they will have to sign the written consent form and complete the questionnaires using a traditional pen and paper method. We will mark the specific date of 12 and 24 weeks of his returning to complete the outcome measurement on his follow-up booklet and tell him if he is not available on that day; he could contact us to change the date accordingly. After that, we will teach patients to register and log in to our WeChat applet ‘TahneeWeb’; this is required to get their step data for the past 2 weeks, which will be recorded by smartphone accelerometers (as done in many prior studies) and has been proven accurate for tracking step counts. Furthermore, a baseline step count will be estimated using the mean step count of the previous 2 weeks.

**Sample size calculation**

The main outcome indicator, daily step count, is selected as the calculation standard. Based on a previous study, we will ensure, at least, 90% power to detect an 800-step difference between each intervention arm and control, an SD of 2000 steps, and a two-sided α of 0.05. In addition, we will use one-way analysis of variance F-tests in PASS V.15.0.5 software and calculate that a total of 84 participants across three arms would be recruited. By allowing for an estimated 20% drop-out rate, a sample size of 108 will be used in this study.

**Randomisation, blinding and concealed allocation**

Patients will be randomised to a study arm using block sizes of 6, stratified by the participant baseline step count (<5000, 5001–7500, or >7500 steps/day). The data collector will be unaware of patient assignments at the baseline, 12 weeks and 24 weeks of the study. Researchers could see the assignments in the backstage of the WeChat applet, and the interfaces of the WeChat applet for patients in different groups are different.

**Control**

All patients will receive step goal setting and could see their progress on the WeChat applet during the 12-week intervention and 12-week follow-up. Personalised daily
Patients tend to be more driven for aspirational behaviour.

Second, every Monday, patients will receive 140 points during the first 6 weeks and then remain fixed during the last 6 weeks, as described elsewhere. To ensure the increase in physical activity is not harmful to participants, the rehabilitation therapists in our research group will assess the condition of each patient and make appropriate adjustments to the step goals. Participants could contact us at any time to make an adjustment if it is due to physical conditions. Moreover, patients could see their daily progress toward their goals using a circular dial on the WeChat applet. Of note, patients in the control group will receive no other interventions. If a patient does not log in to the WeChat applet for over a week, a text message reminder will be sent.

### Interventions

Patients in the Individual and Team groups will receive the gamification intervention based on behavioural economics principles via the WeChat applet. Six behavioural economics principles (precommitment, fresh start effect, goal gradients, loss aversion, anticipated regret and social norms) will be embedded within the gamification intervention. The gamification intervention in the Individual group will apply to four game elements—feedback, points, levels and rewards. In the Team group, collaboration is added besides the four game elements mentioned above. Table 1 provides a summary of game elements, gamification intervention components, and behavioural economics principles.

### Individual group

First, patients will electronically sign a precommitment pledge to try their best to achieve their step goal. Precommitment is known to motivate behavioural change. Second, every Monday, patients will receive 140 points (20 for each day), which leverage the fresh start effect. Patients tend to be more driven for aspirational behaviour around temporal landmarks like the beginning of the week. Third, if patients reach the target step count, no points will be deducted; if not, 20 points will be deducted. This leverages loss aversion, demonstrating that less framing is more effective at motivating behavioural change than gain framing. Fourth, a total of five levels will be set, from low to high—bronze, silver, gold, platinum and diamonds. At the beginning of the trial, the patient is set to the gold level. If the patient has a total score of less than 80 points in a week, the level will drop, and if the total score is greater than or equal to 80 points, the level will rise. Fifth, patients in the two intervention groups will receive feedback weekly based on their progress.

### Team group

The team group will also receive social incentives based on the Individual group. Patients are assigned to a team of three people, who do not know each other before the intervention. Every Monday, the patients will receive 140 points (20 for each day, 10 for themselves, 10 for their team). If the patient achieves the step goals and the other two people in his/her team also achieve the step goals, no points will be deducted. If the patient achieves the step goals but other two people in his/her team do not, 10 points for her team will be deducted; if neither the patient nor the other two people in her team do not achieve the step goal, 20 points will be deducted.

### Outcome measures and data collection

Patients recruited in the trial will be asked to complete the questionnaires and outcome measurements in the CR clinic when they return to hospital for a review. The WeChat applet will automatically remind patients...
to complete the outcome measurements in 12 and 24 weeks. If the patient does not return on time, researchers will telephonically inquire about the reasons. We will also report the numbers and reasons of patients lost to follow-up.

Table 2 shows the summary of the outcome measures for the study. The primary outcome is physical activity participation, which includes a change in daily steps and self-reported physical activity from the baseline to 12 and 24 weeks, and the proportion of patient-days that step goals achieved in 12 and 24 weeks. The daily step counts will be measured and recorded by smartphone accelerometers, proven accurate for tracking step counts.\(^{39}\) Self-reported physical activity level will be measured by the International Physical Activity Questionnaire (IPAQ).\(^ {47}\)

The secondary outcomes include biomedical risk factors, including smoking, intrinsic motivation, enjoyment, competence, autonomy, and relatedness, social support, anxiety symptom, and depressive symptoms. The IPAQ will be filled out online at the baseline, 12 weeks and 24 weeks through the WeChat applet, while the other measurements will be taken in the hospital (at the baseline, 12 weeks and 24 weeks). In addition, usability will be tested at the end of the intervention (at 12 weeks) in the two intervention groups using the System Usability Scale.\(^ {48}\) Furthermore, we will conduct a semi-structured interview to understand patients’ satisfaction, perceptions and experiences in the two intervention groups.

All adverse events will be reported to the ethics committee as required during the 24 weeks study period. Adverse events are defined as medical occurrences resulting in hospitalisation, disability or death.
Statistical analysis and data management
All continuous variables will be reported as mean and SD, and categorical variables will be described as frequencies and percentages. Within-group changes in daily step counts, the proportion of patient-days that step goals attained, self-reported physical activity, biomedical and lifestyle-related risk factors, intrinsic motivation, enjoyment, competence, autonomy, and relatedness, social support, and mental health will be compared using a paired t-test or Wilcoxon rank-sum test depending on the data distribution. Besides, a one-way analysis of variance will be used to compare the intergroup differences between the baseline and postintervention among the outcomes. Main analysis and secondary analysis will be

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BMI, body mass index; DBP, diastolic blood pressure; RHR, resting heart rate; SBP, systolic blood pressure.
conducted of all the outcomes. In the main analysis, the analysis of outcomes will be conducted per the intention-to-treat principle. In addition, multiple imputations for data will be used that are missing and with step values <1000 because evidence indicates that these values are unlikely to represent the capture of actual activity. In the secondary analysis, data analysis will be conducted without multiple imputations, both with and without step values <1000. Furthermore, adjusted analyses include sex, age, BMI, severity of disease and baseline variables. Moreover, we will compare the baseline differences between patients lost to follow-up and patients who adhere to follow-up. All statistical analyses will be two sided, and p<0.05 will be considered statistically significant. We will use SPSS V.20.0 for data analysis.

In this study, well-trained clinical researchers will record all patients’ data using standardised case report forms (CRFs). The original data will be recorded timely and accurately, and a copy of the report will be kept in the laboratory. All CRFs will be stored in a locked file cabinet to prevent data leakage. All laboratory data will be identified with a code number to ensure the confidentiality of subjects’ data. The clinical research data management platform of the School of Nursing of Jilin University will be accountable for data monitoring. The chief investigator can directly access the dataset, and the data scattered to the project team members cannot identify any participant identity information.

Patient and public involvement

Patient and public involvement (PPI) played a vital role in this study. Before designing the WeChat applet, the authors conducted a survey among patients with CHD and found that patients lacked physical activity and were willing to be supervised and motivated via their smartphones to promote participation in physical activity. During the development of WeChat applet, patients with CHD will be invited to participate in our discussion, allowing the authors to consider the thoughts and needs of patients in developing the WeChat applet. In the pilot study, patients will be invited to give reasonable recommendations for study design, questionnaire selection and outcome measurements while considering the burden of intervention. The results of this study will be disseminated to PPI representatives and study participants who wish to be notified.

Validity and reliability

This study will use a rigorous research design (randomised controlled design) and a block random method to assign groups. The grouping results will be numbered and placed in a sealed envelope. Participants and data collectors will be blinded to the assignments. All questionnaires will be completed by the researcher’s guidance or ghostwriting. The questionnaires will be distributed and collected on the spot to avoid data bias caused by different researchers. Two researchers will enter all the data to avoid objective typing errors.

ETHICS AND DISSEMINATION

This study will comply with the ethical principles of the Declaration of Helsinki, and the Human Research Ethics has been approved by the School of Nursing, Jilin University (HREC 2020122401). All participants will be required to provide written informed consent. Research reports will be disseminated through scientific forums, including peer-reviewed publications and presentations at national and international conferences.

DISCUSSION

The authors aim to develop a WeChat applet in this study. Based on the WeChat applet and under the guidance of behavioural economics principles, the authors will develop a gamification intervention using five game elements, including points, levels, feedback, rewards and collaboration. This study will evaluate the role of mHealth-based gamification intervention on physical activity participation and the effects on biomedical and lifestyle-related risk factors, intrinsic motivation, enjoyment, competence, autonomy, and relatedness, social support, and mental health. Moreover, the authors will conduct a semi-structured interview after the intervention to elucidate patients’ satisfaction, perceptions and experience of the intervention.

Despite proven benefits, patients with CHD do not often attain their physical activity goals on their own. Behavioural interventions are needed to help them increase physical activity participation. With technological advancement, numerous smartphone apps have appeared, and gamification was used in most of these apps, which could increase physical activity motivation and promote behavioural change. However, most gamification intervention designs did not appropriately leverage theories from health behaviour models, and empirical evidence on their efficacy is still emerging. A previous study established that behavioural economics principles could be embedded with a gamification intervention to significantly increase physical activity among overweight and obese adults. However, thus far, there is limited evidence of interventions that use these methods to effectively improve physical activity participation among high-risk patients, such as patients with CHD.

Thus, this study will develop a gamification intervention based on a WeChat applet that has been specifically developed for this study. Personalised goal setting and promotion of patients’ satisfaction, perceptions and experience of the intervention.

The key of gamification interventions is to organically combine game elements to form a resultant force to improve physical activity, and the key of the resultant force is to comprehend the driving force or motivation behind the incentive mechanism. Research indicates links between self-determination theory and gamification.
concepts. Self-determination theory suggests that satisfying three innate psychological needs of competence, autonomy and relatedness could promote autonomous motivation and well-being. Reportedly, individuals with autonomous motivation had higher physical activity participation and better physical activity adherence than those primarily driven by external factors. Furthermore, the fact that gamification could make interventions more enjoyable aligns with self-determination theory, which assumes that a key aspect of intrinsic motivation is enjoyment. We plan to investigate the internal psychological mechanism of gamification to promote physical activity; thus, we will evaluate competence, autonomy, relatedness, enjoyment and intrinsic motivation. We assume that gamification intervention promotes the transformation of controlled motivation into autonomous motivation by satisfying competence, autonomy, relatedness, enjoyment and ultimately promote physical activity participation. Figure 4 shows the hypothesised model of physical activity behaviour regulation.

In China, access to CR is often limited. Patients with CHD often lack physical activity. Our intervention could help increase physical activity participation and bring more health benefits.

Limitations
This study has several limitations. First, we will not measure the intensity of physical activity via the smartphone accelerometer. In future, we plan to use wearable devices to evaluate the intensity of physical activity. Second, the study is limited to patients with smartphones and a WeChat account, which could lead to a selective bias. Third, the gamification interventions are comprehensive, and it would be difficult to analyse the component that worked. Fourth, it is a multilayered and complex intervention, and the projected sample size will make it challenging to say that the results will be much more than a pilot study given there will be three groups.

Acknowledgements
The authors would like to thank the valuable contribution made by the patients and public representatives during the study design and intervention development.

Contributors
LX, QT and FL conceived the original concept of the study and wrote the first draft of the protocol manuscript. LX, JL, XZ, YP, TY, XL, TY and LZ contributed to the design of the study. All authors read and approved the final manuscript.

Funding
This work is financially supported by a Construction Programme of Independent Innovation Ability of Community Health Nursing Engineering Laboratory in Jilin Province (Study code: 2020C038-8) awarded to FL.

Competing interests
None declared.

Patient consent for publication
Consent obtained directly from patient(s)

Provenance and peer review
Not commissioned; externally peer-reviewed.

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REFERENCES


