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# An eHealth intervention for patients with a low socioeconomic position during their waiting period preceding cardiac rehabilitation: a randomized feasibility study

Jasper S. Faber <sup>1,\*</sup>, Jos J. Kraal <sup>1</sup>, Nienke ter Hoeve <sup>2,3</sup>, Isra Al-Dhahir <sup>4</sup>,  
Linda D. Breeman <sup>4</sup>, Niels H. Chavannes <sup>5,6</sup>, Andrea W. M. Evers <sup>1,4,7</sup>,  
Hans B. J. Bussmann<sup>2</sup>, Valentijn T. Visch <sup>1</sup>, and Rita J. G. van den Berg-Emons <sup>2,3</sup>

<sup>1</sup>Department of Human-Centered Design, Faculty of Industrial Design Engineering, Delft University of Technology, Landbergstraat 15, Delft 2628 CE, The Netherlands; <sup>2</sup>Department of Rehabilitation Medicine, Erasmus MC, Rotterdam, The Netherlands; <sup>3</sup>Capri Cardiac Rehabilitation, Rotterdam, The Netherlands; <sup>4</sup>Faculty of Social and Behavioural Sciences, Leiden University, Leiden, The Netherlands; <sup>5</sup>Department of Public Health and Primary Care, Leiden University Medical Centre, Leiden, The Netherlands; <sup>6</sup>National eHealth Living Lab, Leiden University Medical Centre, Leiden, The Netherlands; and <sup>7</sup>Medical Delta, Leiden University, Delft University of Technology, Erasmus University, Delft, The Netherlands

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## Aims

Cardiac rehabilitation (CR) shows lower effectiveness and higher dropouts among people with a low socioeconomic position (SEP) compared to those with a high SEP. This study evaluated an eHealth intervention aimed at supporting patients with a low SEP during their waiting period preceding CR.

## Methods and results

Participants with a low SEP in their waiting period before CR were randomized into an intervention group, receiving guidance videos, patient narratives, and practical tips, or into a control group. We evaluated adherence (usage metrics), acceptance (modified Usefulness, Satisfaction, and Ease of use questionnaire), and changes in feelings of certainty and guidance between the waiting period's start and end. Semi-structured interviews provided complementary insights. The study involved 41 participants [median interquartile range (IQR) age 62 (14) years; 33 males], with 21 participants allocated to the intervention group, using the eHealth intervention for a median (IQR) duration of 16 (10) days, using it on a median (IQR) of 100% (25) of these days, and viewing 88% of the available messages. Key adherence themes were daily routine compatibility and curiosity. Acceptance rates were 86% for usability, 67% for satisfaction, and 43% for usefulness. No significant effects on certainty and guidance were observed, but qualitative data suggested that the intervention helped to inform and set expectations.

## Conclusion

The study found the eHealth intervention feasible for cardiac patients with a low SEP, with good adherence, usability, and satisfaction. However, it showed no effect on feelings of certainty and guidance. Through further optimization of its content, the intervention holds promise to improve emotional resilience during the waiting period.

## Registration

This trial is registered as follows: 'Evaluation of a Preparatory eHealth Intervention to Support Cardiac Patients During Their Waiting Period (PReCARE)' at ClinicalTrials.gov (NCT05698121, <https://clinicaltrials.gov/study/NCT05698121>).

## Lay summary

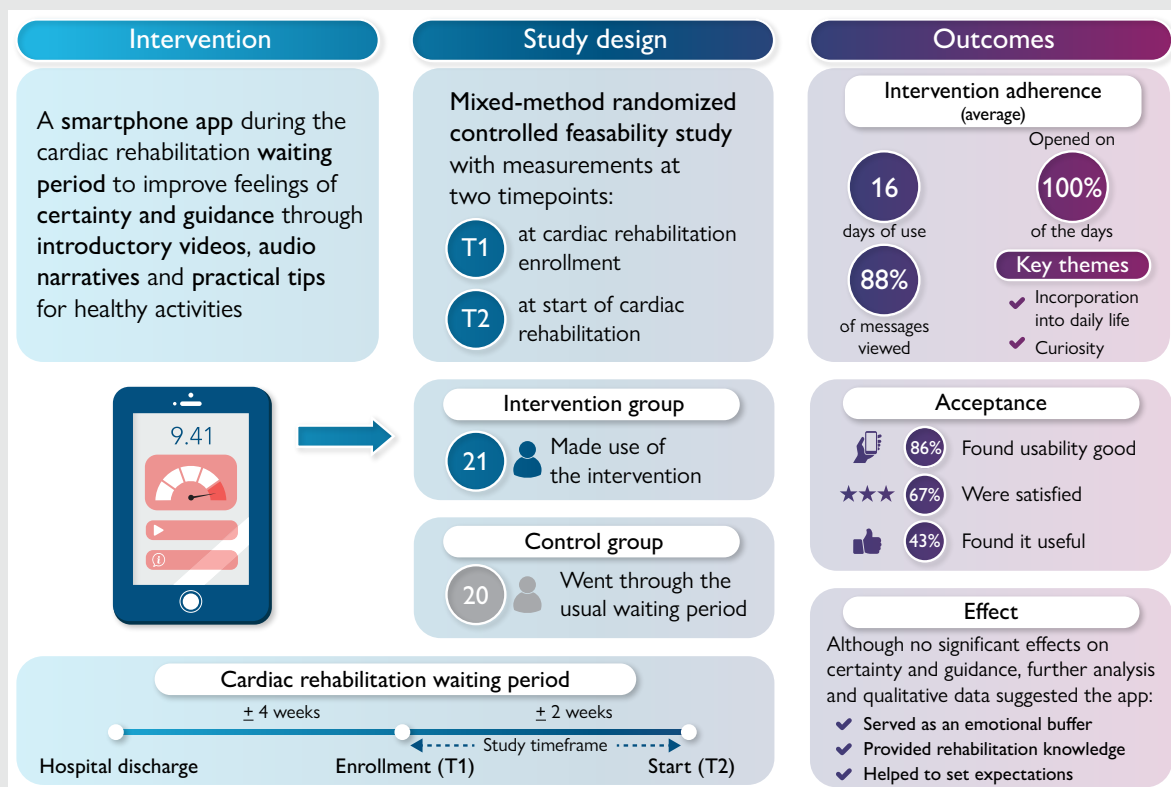
This study evaluated the feasibility of an eHealth intervention designed for cardiac patients with a low socioeconomic position (SEP) during the waiting period before cardiac rehabilitation (CR). The intervention prepares patients with a low SEP for their upcoming CR and consists of a smartphone application providing guidance videos, patient narratives, and practical tips.

\* Corresponding author. Tel: +31640850881, Email: [j.s.faber@tudelft.nl](mailto:j.s.faber@tudelft.nl)

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## Graphical Abstract



## Keywords

Cardiac rehabilitation • Socioeconomic factors • Telemedicine • Time to treatment

## Introduction

Cardiac rehabilitation (CR) is a multicomponent lifestyle intervention that includes information and coaching on healthy behaviour and supervised exercise training.<sup>1,2</sup> Cardiac rehabilitation is crucial for cardiac patients to prevent secondary health problems and decrease mortality rates. It has been shown to improve patient outcomes like physical fitness and health-related quality of life.<sup>3</sup> However, the effectiveness of CR is not uniformly experienced. Specifically, individuals with a low socioeconomic position (SEP) often show lower participation rates in these programmes and drop out more frequently.<sup>4-6</sup> This disparity can be attributed to various barriers to participation,<sup>7</sup> such as stressful life situations,<sup>8</sup> environmental accessibility issues,<sup>9</sup> inadequate social support,<sup>10</sup> stigma and distrust in healthcare,<sup>11</sup> and low health literacy.<sup>12</sup> Due to these disparities, CR is not fully benefitting patients with a low SEP, underscoring the need for solutions to make CR more inclusive and accessible.

Our previous research highlights an opportunity to address the barriers faced by cardiac patients, especially those with a low SEP, during the waiting period between hospital discharge and the start of CR.<sup>13</sup> This waiting period lasts, on average, 6 weeks.<sup>14</sup> It is marked by emotional vulnerability and uncertainty, as patients often leave the hospital with unmet informational needs about their condition and self-care.<sup>15-18</sup> The absence of adequate guidance during the waiting period, exacerbated by the initial shock of diagnosis or surgery, leads to a passive patient attitude<sup>19-21</sup> and a disjointed transition between healthcare

settings.<sup>22</sup> Patients with a low SEP feel this lack of guidance more strongly. Their additional challenges increase their vulnerability and uncertainty during the waiting period.<sup>23</sup> As a result, this group is less likely to adopt the necessary 'readiness' to successfully engage with CR. This leads to lower participation and higher dropout rates during CR.<sup>24,25</sup>

eHealth interventions are a promising strategy to overcome barriers that arise during the waiting period. They can better prepare cardiac patients with a low SEP for CR. For example, these interventions can fill the existing information and guidance gap by leveraging online information platforms<sup>15</sup> and goal-monitoring tools.<sup>26</sup> Due to rising healthcare costs, addressing these needs through face-to-face sessions during the waiting period may not be feasible.<sup>27-29</sup> In theory, eHealth interventions offer a cost-effective alternative to face-to-face sessions.<sup>28-32</sup> However, in practice, people with a low SEP often do not adhere to these interventions due to low technology access, low digital literacy, and other life priorities.<sup>33,34</sup> The success of these interventions depends on tailoring them to the specific needs, abilities, and preferences of this group.<sup>35</sup>

We recently developed the Inclusive eHealth Guide (IeG) to support the design of tailored eHealth interventions according to the specific needs of individuals with a low SEP.<sup>36</sup> The guide combines existing knowledge on barriers and facilitators in eHealth development for individuals with a low SEP.<sup>37</sup> It considers, among others, the target group's context, needs, preferences, and skills.<sup>38</sup> We applied the IeG in a participatory design process of an eHealth intervention specifically for and with cardiac patients with a low SEP. The intervention addresses their needs during the waiting period before CR.<sup>13</sup>

This study aimed to evaluate the feasibility of this eHealth intervention tailored towards CR patients with a low SEP in the domains of adherence and acceptance. Additionally, it explored the effects of the eHealth intervention on feelings of certainty and guidance, factors associated with changes in these constructs, and dropout rates during subsequent CR.

## Methods

### Study design

The feasibility study was executed between February 2023 and September 2023 at Capri Cardiac Rehabilitation, a CR centre with sites in Rotterdam and The Hague (The Netherlands). The participants were randomized to an intervention group and control group, and outcomes were assessed at the start and end of the waiting period before CR started.

### Recruitment

Eligible participants were adults aged 18 or above living in a low SEP neighbourhood, referred to CR by their cardiologist, able to understand Dutch (with assistance), and had a mobile device with internet access. Postal codes of potential participants were sent to the principal investigator (J.S.F.) to assess neighbourhood SEP, based on the neighbourhood residents' average income and education levels. We used a list of 40 neighbourhoods identified by the Dutch government for their socioeconomic challenges as a benchmark.<sup>39</sup> A representative from the CR centre first contacted potential participants for consent. Interested patients were then contacted by the investigator (J.S.F.), who explained the study. If they agreed to participate, they received an information letter and had an appointment scheduled for the initial assessment.

### Measures

#### Adherence

Adherence to the intervention was measured using the metrics (i) *use period length*: the number of days between the first and last day the intervention was used; (ii) *percentage of active days*: percentage of days the intervention was used; (iii) *daily use time*: average time spent on the intervention per active day within the use period; and (iv) the *total number of viewed messages*.

#### Acceptance

Acceptance was measured using a modified Usefulness, Satisfaction, and Ease of use (USE) questionnaire. The original USE questionnaire consists of 30 items on a 7-point Likert scale focusing on usefulness, satisfaction, ease of use, and ease of learning.<sup>40</sup> In alignment with the specific needs and challenges faced by our target population of individuals with a low SEP, we recognized that lengthy questionnaires often lead to disengagement among this group.<sup>41</sup> Therefore, we adapted the original questionnaire to a more manageable version with only nine of the original items (three per category) on a 5-point Likert scale, focusing on usefulness, satisfaction, and ease of use (see [Supplementary material online, Appendix S1](#)). The questions retained were chosen for their relevance to the unique context and goals of the current intervention.

#### Certainty and guidance and influencing factors

We developed the Certainty and Guidance Questionnaire (CGQ), consisting of seven items measured using a 5-point Likert scale, for use in this study (see [Supplementary material online, Appendix S1](#)). High scores indicate good certainty and guidance. The questionnaire focuses on patient needs identified in a previous study.<sup>13</sup> These needs include feeling certain during the waiting period, confidence to be physically active, managing expectations about the contents of CR, good management of emotions, the feeling of hope for future recovery, understanding the current health status, and feeling guided before the start of CR. The questions are derived from existing scales, including the Motivation for Traumatic Brain Injury Rehabilitation Questionnaire for motivation,<sup>42</sup> the Patient Evaluation of Emotional Comfort Experienced for experienced emotional comfort,<sup>43</sup> and the Credibility and Expectancy Questionnaire for expectancy and credibility<sup>44</sup> to strengthen the validity of our measurements. Finally, to better understand

the factors influencing changes in feelings of certainty and guidance, we explored associations between age, education level, length of waiting period, baseline level of CGQ, and the change in CGQ scores in both the intervention and control groups.

### Qualitative insights

In line with our mixed methods approach, we complemented the quantitative data for adherence, acceptance, and feelings of certainty and guidance with qualitative insights collected with semi-structured interviews. We asked questions relating to reasons for adherence (e.g. Why did or did you not succeed in using the intervention daily?), acceptance (e.g. What was your experience with using the intervention?), and effects on feelings of certainty and guidance (e.g. How has the intervention been able to help you the most during the waiting period?) (see [Supplementary material online, Appendix S2](#), for the full interview guide).

### Procedures

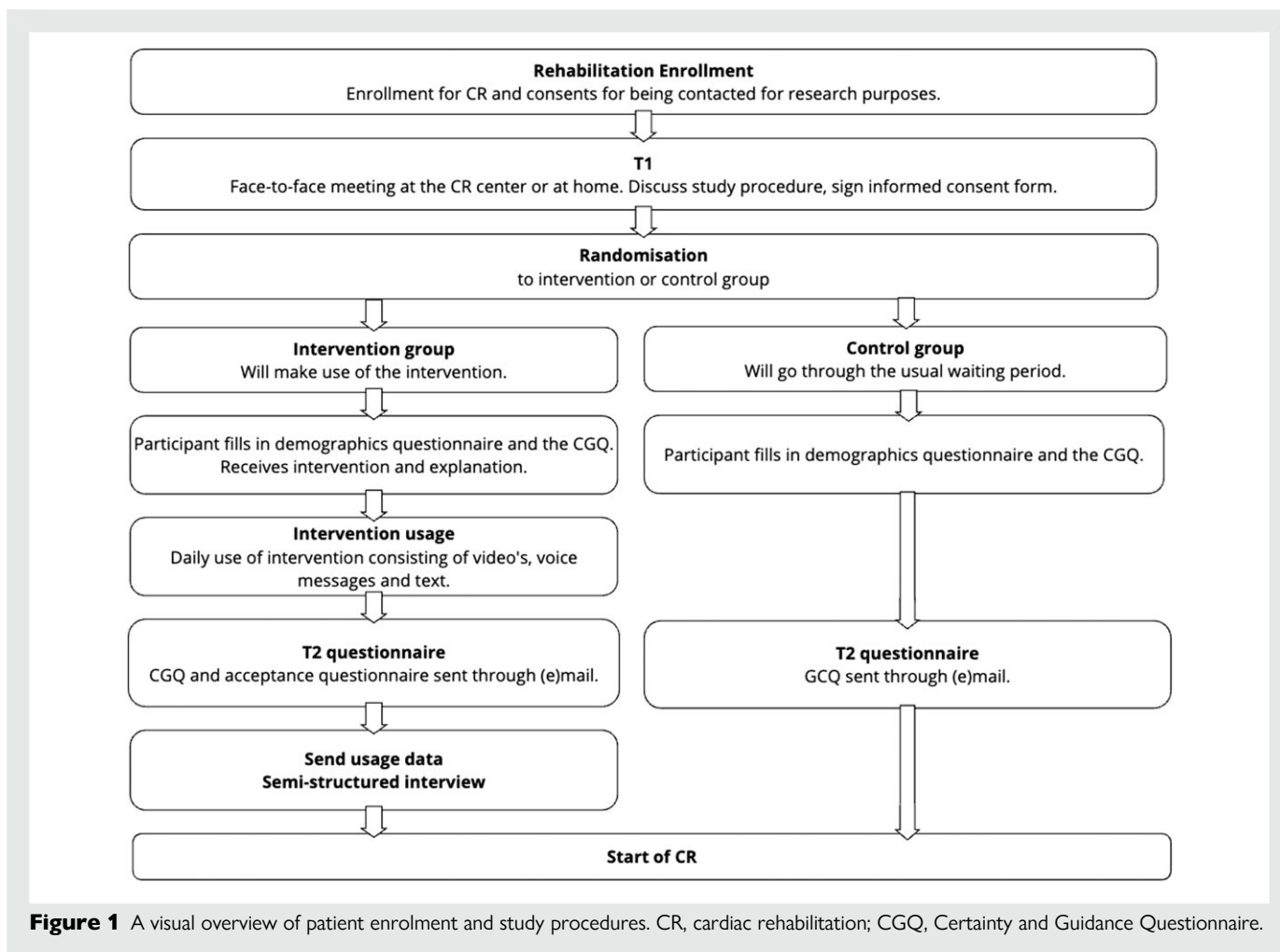
We performed assessments in both study groups during two contact moments: initially upon enrolment in CR ( $T_1$ ) and again just before the CR programme began, usually 2–8 weeks after  $T_1$  ( $T_2$ ) (see [Figure 1](#)). At  $T_1$ , participants were briefed on the study, signed consent forms, and completed demographic and CGQ questionnaires. The intervention group received additional information about the smartphone app and help installing it. At  $T_2$ , both groups completed a second CGQ questionnaire, with the intervention group also submitting usage data, filling out an acceptance questionnaire, and participating in a phone interview. Participants received a 20-euro gift voucher for their participation.

### Intervention

The CapriXpress application is a tailored digital intervention developed to support patients with a low SEP during their waiting period between discharge from the hospital and the start of their CR. This intervention was co-designed in a participatory design study with the target group.<sup>13</sup> The CapriXpress application addresses the need for certainty and guidance for people with a low SEP during their waiting period. To enhance patient adherence and acceptance of the intervention, we ensured that the intervention is grounded in established theoretical frameworks, namely the taxonomy of behaviour change techniques (BCTs)<sup>45</sup> and the persuasive systems design (PSD) model<sup>46</sup> (see [Table 1](#)). Combining the two frameworks is valuable as it combines the rigour of scientifically validated methods for behaviour change with engaging, user-focused aspects of persuasive technology design.<sup>47</sup> Additionally, the intervention integrates recommendations derived from our previously developed *leG*,<sup>36</sup> which served as a foundational resource (see [Table 1](#)).

The content of the CapriXpress application is divided into concise, manageable units, with a limited number of messages presented per day ([Figure 2, 1.1](#)). The interface is designed to be playful, aesthetically pleasing, and simple to understand ([Figure 2, 1.2 and 1.3](#)). A 'travel bag' feature stores completed messages, giving patients a sense of achievement. When the bag is filled to a certain level, it receives an aesthetic upgrade as a reward ([Figure 2, 1.4 and 1.5](#)). The information is primarily conveyed through multimedia formats and is articulated in easily understandable language, adopting a positive tone ([Figure 2, 1.6](#)). A push notification is sent if the participant has not engaged with the intervention for two consecutive days. The 'help' section provides contact information for research or application-related questions ([Figure 2, 1.9](#)).

Several intervention features have been implemented to address the target group's needs during the waiting period. The application uses a calendar-based train journey metaphor to symbolize the patient's progression towards the start of CR ([Figure 2, 2.1](#)). This progression occurs automatically over time, aiming to provide a sense of certainty during the waiting period. The app delivers a total of 43 multimedia messages, defaulting to three per day, with the frequency adjusting based on the patient's specified CR start date. Patients can choose from three message types: introductory videos from healthcare providers like a physiotherapist, dietitian, and psychologist to inform and connect with the CR team ([Figure 2, 2.3](#)); audio narratives from former patients to offer emotional support and hope ([Figure 2, 2.4](#)); and actionable advice promoting healthy activities and improving understanding of their condition and the rehabilitation process ([Figure 2, 2.5](#)).



## Data analysis

We analysed our quantitative data in RStudio (Version 2023.06.0, Posit Software, PBC). We utilized medians, interquartile ranges (IQRs), and non-parametric statistical tests to ensure robustness and minimize assumptions about data distribution, given our limited sample size. The level of significance was set at  $P \leq 0.05$ .

We transformed the raw intervention usage data into specific metrics to evaluate adherence. We calculated the use period length from the first day of use ( $T_1$ ) to the last completed message. The percentage of active days was determined by dividing the number of days the intervention was used by the total use period length. Daily use time was derived by summing the duration of all visits and dividing it by the number of active days. We also totalled the number of viewed messages. Adherence was considered satisfactory if participants used the intervention on at least half the days and viewed more than half the available messages.

To analyse the acceptance from the adapted USE questionnaire, we classified the Likert scores as negative (1 or 2), neutral (3), or positive (4 or 5) and calculated the percentages of participants in each category. We then calculated individual scores for usability, usefulness, and satisfaction and determined the median, IQR, minimum, and maximum scores for these metrics across intervention group participants. Overall acceptance was similarly assessed using these statistics. For this prototype, a score was deemed good if over 60% of the participants rated it positively.

To assess the intervention's effect on certainty and guidance during the waiting period, we calculated the median Likert scores for the CGQ items for each participant and determined group medians for both the intervention and control groups. Wilcoxon rank-sum and Mann-Whitney  $U$  tests were used to assess within- and between-group differences, respectively. Rank correlation tests examined the relationship between changes in

CGQ scores and factors such as age, education level, initial CGQ scores, and waiting period length. Fisher's exact test was used to analyse differences in dropout rates.

For the qualitative data, we performed a thematic analysis<sup>49</sup> using ATLAS.ti (Version 9.1.3, ATLAS.ti Scientific Software Development GmbH). Interviews were transcribed verbatim, followed by coding individual quotations and corresponding interpretations. These codes were then grouped into overarching themes related to the outcome measures, such as adherence, acceptance, and impact on feelings of certainty and guidance.

## Ethics and data management

This study adhered to the Declaration of Helsinki principles and was approved by the Medical Ethics Committee of Erasmus MC (MEC-2022-0483) and registered in clinicaltrials.gov (NCT05698121). Written informed consent was obtained from all study participants.

## Results

### Participants

Out of the 835 patients referred to the CR centre during the recruitment period (January 2023 to June 2023), 149 patients (18%) were eligible, of which 42 patients (28%) consented to participate. Frequently reported reasons for non-participation were personal circumstances, logistical issues, lack of interest, technological barriers, and language and cognitive barriers. Twenty-one participants were assigned to the intervention group and 21 to the control group (see [Figure 3](#)). One

**Table 1** Intervention features that address adherence and acceptance, linked to principles from the Inclusive eHealth Guide, behaviour change technique, and persuasive systems design framework, and features that address patient needs as identified in our prior study

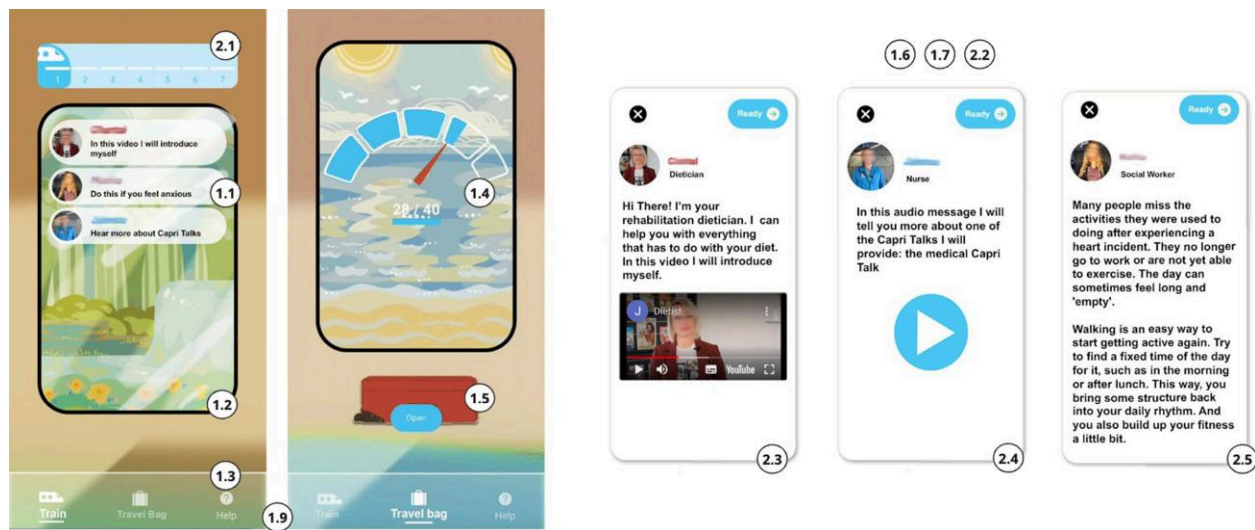
Features addressing adherence and acceptance				
Number	Feature	leG recommendation	BCT	PSD principle
1.1	Limited number of daily messages	Realistic, achievable goals, align with life situation	Graded tasks	Reduction
1.2	Playful interface	Positive approach		Liking
1.3	Simple interface	Simplicity		Reduction
1.4	Done-pile tracker	Short-term goals, apply gamification	Self-monitoring/feedback	Self-monitoring
1.5	Bag upgrade	Reward for adherence, apply gamification	Non-specific reward	Rewards
1.6	Use of multimedia and simple language	Simplify communication		Tailoring
1.7	Tone of voice	Positive approach		
1.8	Notification	Send reminders		Reminders
1.9	Support page	Offer technical support		

Features addressing patient needs		
Number	Feature	Needs
2.1	Calendar-based progression	Certainty during waiting period, pre-CR guidance
2.2	Information provided by healthcare provider	Certainty during waiting period, pre-CR guidance
2.3	Video introductions	Certainty during waiting period, CR expectancy
2.4	Spoken peer stories	Certainty during waiting period, managing emotions, future recovery
2.5	Textual advice	Physical activity confidence, health status understanding

Inclusive eHealth Guide (leG)<sup>36</sup>, behaviour change technique (BCT) taxonomy<sup>48</sup>, and persuasive system design (PSD) model,<sup>46</sup> and features that address patient needs as identified in our prior study.

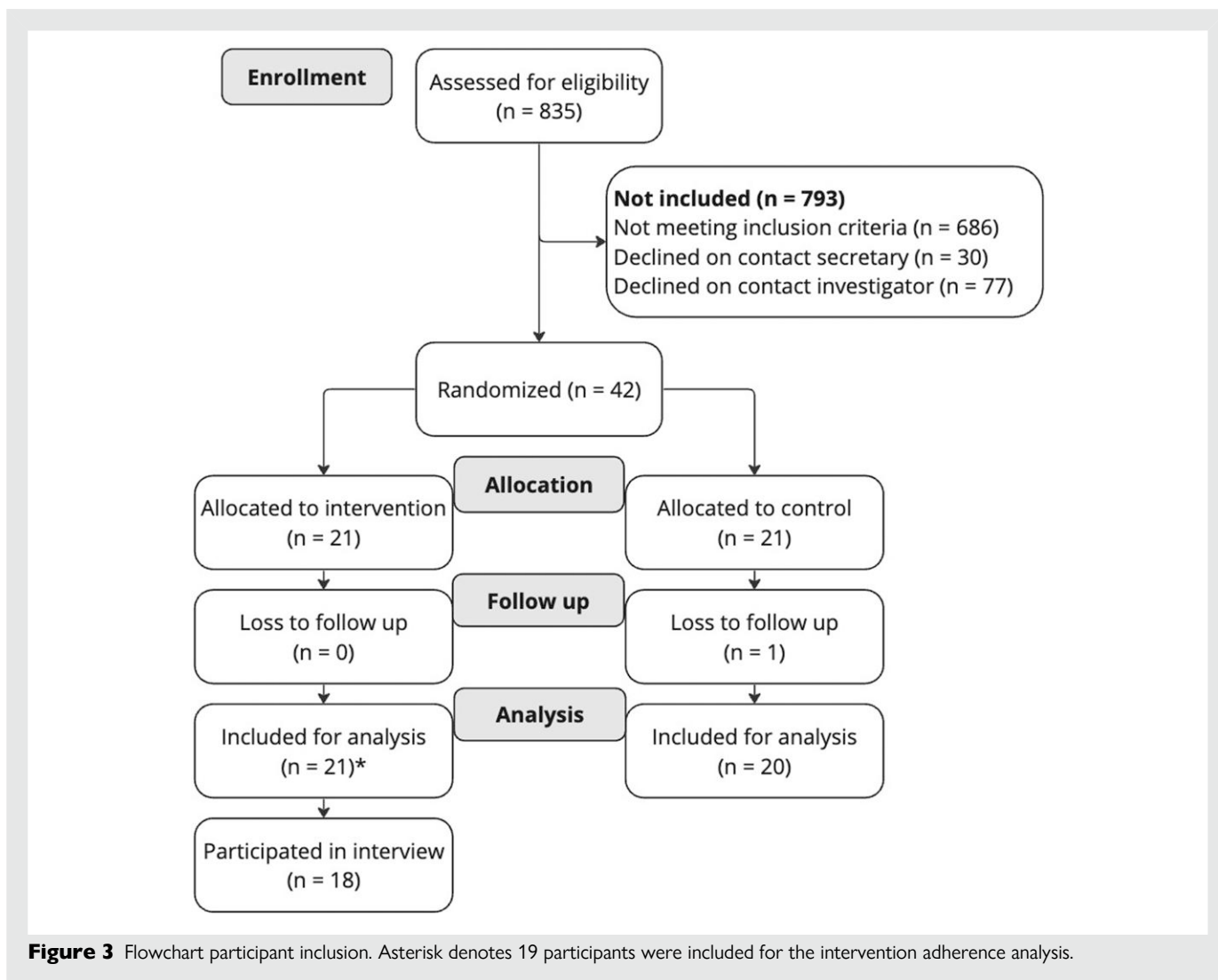
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**Figure 2** Key interface screens from the CapriXpress intervention. From left to right: journey-based progression home page, done-pile tracker and travel bag, and multimedia messages from healthcare professionals.

participant in the control group dropped out during the study due to the burden of participation. Eighteen participants from the intervention group participated in a semi-structured interview, and 19 participants from the intervention group sent their usage data for the adherence analysis. The majority of the sample was male (80%), with a median

(IQR) age of 62 (14) years. Ischaemic heart disease was the most common condition (63%). The median (IQR) waiting time from hospital discharge to the start of CR was 55 (43) days and 29 (13) days from enrolment at the CR facility to the beginning of the programme (see Table 2 for more details).



## Adherence to the intervention

The median (IQR) *use period length* was 16 (10) days. During this period, the median (IQR) percentage of the days the participants accessed the application was 100% (25), with a median (IQR) *daily use time* of 4 (2) min. Sixty-seven per cent of the participants opened the application every day. Regarding content interaction, the median (IQR) *total number of messages viewed* was 38 (24) out of 43. Half of the participants viewed all the messages available. [Figure 4](#) presents the relationship between the number of days since first usage and the cumulative messages completed by participants. The trend line shows continuous completion of messages over time with a slight decrease in the number of messages completed each day after ~2 weeks. Two qualitative themes related to these adherence patterns emerged (see [Supplementary material online, Appendix S3](#), for a complete overview of the qualitative themes). First, almost three-quarters of the participants stated that the intervention aligned well with their daily routines. As one participant expressed:

We used to sit in the morning for coffee. Yeah, we would sit down for a while, and I received them [the messages], and then I had my phone in my hand. Well, I went through it; I even turned it on so that the lady could listen along and that way. Yeah, it's also at a fixed time. You have to be careful not to leave it for a whole

week and then review it after a week. Because that will not work, I think. If you throw everything together, it is just a matter of sifting through it and fulfilling a duty. [Male, 73]

Second, we found that more than half of the interviewed participants cited curiosity as their driving factor for usage. As one participant stated:

I was curious about it every day. I also opened it every day. I went through the entire program. I was, well, actually, looking forward to seeing what news they had to say today. Yeah, it was actually more curiosity. [Male, 69]

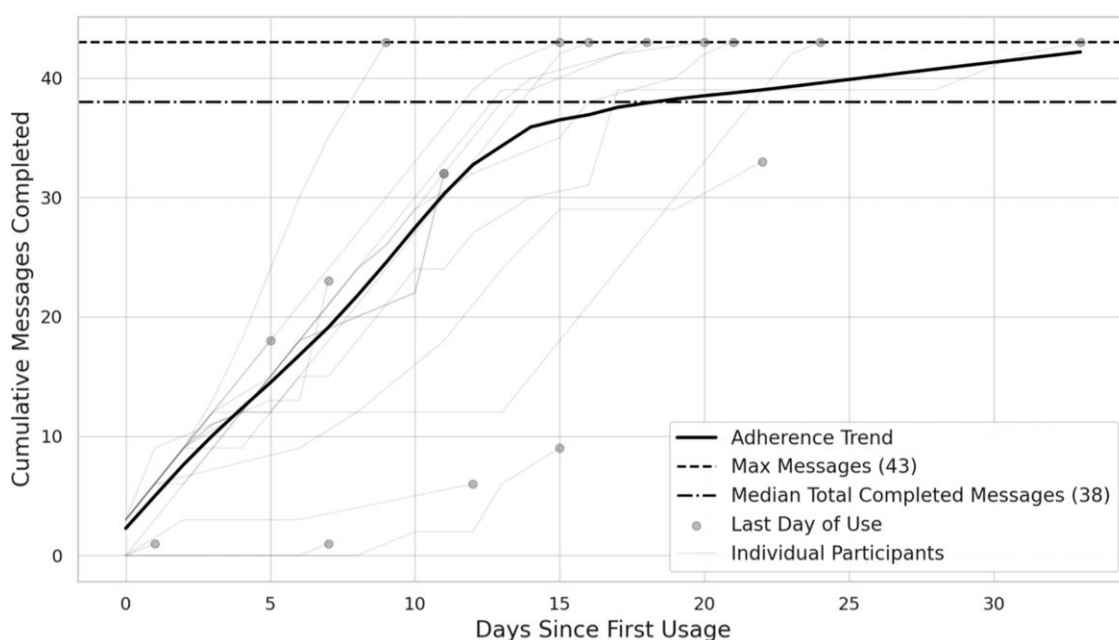
## Acceptance of the intervention

Seventy-one per cent of the participants displayed overall positive acceptance. We found that 86% of the participants were positive about the intervention's usability, and 67% were satisfied. Forty-three per cent felt that the intervention was useful for them (see [Table 3](#) for a complete overview of the acceptance scores). Within the qualitative data, we found that participants mainly appreciated the *ease of use* and the *playful interface*. As one participant expressed:

Well, you know, I found it enjoyable. It's more enjoyable than just a boring list or something, you know. Yeah, it's funny that they

**Table 2 Participant characteristics**

Characteristic	Intervention group (n = 21)	Control group (n = 20)	Sample (n = 41)
<b>Demographics</b>			
Male, n (%)	17 (81)	16 (80)	33 (80)
Age (years), median (IQR)	63 (13)	59 (13)	62 (14)
Low education, n (%)	15 (71)	18 (90)	33 (80)
Employed, n (%)	4 (19)	6 (30)	10 (25)
Unemployed, n (%)	1 (5)	2 (10)	3 (7)
Retired, n (%)	11 (52)	7 (35)	18 (44)
Unfit for work, n (%)	5 (24)	5 (25)	10 (24)
<b>Medical history, n (%)</b>			
Ischaemic heart disease	15 (70)	11 (55)	26 (63)
Cardiac arrhythmia	2 (10)	1 (5)	3 (7)
Other, cardiac disease	4 (20)	7 (35)	11 (27)
Other, non-cardiac disease	0 (0)	1 (5)	1 (3)
<b>Waiting time, days, median (IQR)</b>			
Hospital discharge—start CR	66 (31)	43 (32)	55 (43)
Enrolment CR—start CR	28 (13)	29 (11)	29 (13)



**Figure 4** Cumulative messages completed vs. days since the first usage (median ± interquartile range).

thought of it like, oh yeah, let's pretend it's a journey. With your stories in a suitcase, very amusing. You're on a journey to your rehabilitation. [Female, 60]

We also found qualitative themes that related to the usefulness. More than half of the participants suggested the need for more *personally relevant information* better aligned with their health concerns and the severity of their conditions. As one participant expressed:

All those social workers and such... For me, I think it's not interesting. I only do it to become physically well. That's my goal. I don't think I have any other issues. I think the app is limited in that aspect. [Male, 76]

Additionally, approximately half of the participants suggested a need for *additional depth and detail* in the provided information. As one participant expressed:



**Table 3** Overview of acceptance scores displayed as median interquartile ranges and percentages of participants in the categories positive, neutral, and negative on usability, satisfaction, and usefulness

Question	Median (IQR), minimum–maximum	Positive (4–5) n (%)	Neutral (3) n (%)	Negative (1–2) n (%)
Overall	3.8 (0.8), 2.7–5.0	15 (71)	6 (29)	0 (0)
Usability	4.0 (1.0), 2.7–5.0	18 (86)	3 (14)	0 (0)
Is easy to use	4 (1), 3–5	18 (86)	3 (14)	0 (0)
Required no effort	4 (1), 2–5	19 (90)	1 (5)	1 (5)
Allowed to perform well	4 (2), 2–5	15 (71)	5 (24)	1 (5)
Satisfaction	4.0 (0.7), 2.3–5.0	14 (67)	6 (28)	1 (5)
Is fun to use	4 (1), 3–5	14 (67)	7 (33)	0 (0)
Would recommend to others	4 (1), 2–5	18 (85)	2 (10)	1 (5)
Aligns with needs	4 (1), 1–5	14 (67)	3 (14)	4 (19)
Usefulness	3.3 (1.0), 2.3–5.0	9 (43)	10 (47)	2 (10)
Is useful	4 (1), 3–5	13 (62)	8 (38)	0 (0)
Aligns with needs	3 (1), 2–5	7 (33)	11 (52)	3 (15)
Aligns with expectations	4 (1), 2–5	11 (52)	6 (29)	4 (19)

The dietitian gave a very brief explanation of what she does. [...] But she didn't really delve into the topic. For example, what can you tell about your sugar or salt levels being too high? What are the consequences of that? Could you get paralysis? Could you have a heart attack? So, the information was lacking, in my opinion. [Male, 63]

### Effects on certainty and guidance

We found no significant changes in CGQ scores between T<sub>1</sub> and T<sub>2</sub> in both the intervention [ $\Delta = -0.14$  (IQR 0.57),  $P = 0.94$ ] and control group [ $\Delta = -0.07$  (IQR 0.32),  $P = 0.51$ ]. In addition, we did not find a significant difference in the changes in CGQ scores between the two groups ( $P = 0.51$ ). The qualitative data highlighted areas that did address improvements in feelings of certainty and guidance during the waiting period. Participants suggested that the intervention improved their expectancy about the CR, as one participant mentioned:

You know, when all those people introduced themselves and told stories about different participants. Yeah, that was nice because you know beforehand what to expect when you start the rehabilitation. So, that was quite pleasant. [Male, 73]

In addition, it helped them to feel generally better informed about their current condition by providing additional knowledge they usually would not receive. As one participant expressed:

I received a lot of information that I wouldn't normally get. If you haven't had a heart attack, you don't even think about all the information you've received. So, for me, it was a kind of recognition. And it was very good. So, as I said, it made me wise. [Male, 63]

Finally, the participants highlighted the intervention helped to reduce uncertainties by providing guidance during the waiting period gap. As expressed by one participant:

Well, in terms of reducing uncertainties, the app did help me because if you didn't have that app, you would fall into a void between being discharged from the hospital and starting rehabilitation. So, in that sense, the app was able to provide assistance in filling that void at some point. [Male, 74]

### Factors associated with the effect on certainty and guidance

The length of the waiting period had a significant negative correlation with the change in CGQ score in the control group ( $\rho = -0.51$ ,  $P = 0.02$ ) but not in the intervention group ( $\rho = -0.04$ ,  $P = 0.86$ ). In addition, higher CGQ scores at T<sub>1</sub> were negatively correlated with changes in CGQ scores in both the intervention group ( $\rho = -0.56$ ,  $P = 0.01$ ) and the control group ( $\rho = -0.49$ ,  $P = 0.03$ ). Age and education were not significantly correlated with changes in CGQ scores in both intervention ( $P = 0.16$  and  $P = 0.26$ , respectively) and control ( $P = 0.94$  and  $P = 0.66$ , respectively) groups.

### Effects on dropout during CR

Two (10%) of the participants dropped out of the subsequent CR programme in the control group compared to none in the intervention group. This difference was, however, not significant ( $P = 0.23$ ).

## Discussion

### Principal findings

In this study, we evaluated the feasibility and explored the effects on feelings of certainty and guidance and on dropouts of a newly developed eHealth intervention for CR patients with a low SEP during their waiting period before starting a CR programme. We found good adherence with the participants often using the intervention daily and engaging with 88% of the messages. Most participants (71%) displayed positive overall acceptance of the intervention. However, only 43% were positive about usefulness. The intervention did not affect feelings of certainty and guidance (CGQ) or dropout rate. However, while the length of the waiting period was negatively associated with feelings of certainty and guidance in the control group, no such association was observed in the intervention group. Qualitative feedback suggested that the intervention had helped participants to set expectations and be better informed about their condition and CR journey.

Usage data indicated consistent adherence over time, although there was a slight reduction in daily message interactions after the first 2 weeks. This decrease aligns with the intervention's dynamic content distribution system, which recalibrates the frequency of messages once participants enter their CR start dates. When the starting date

is further in the future, the system automatically reduces the number of messages provided daily to extend the usage period. Despite this, continued message views suggest sustained adherence, which contrasts with the relatively low eHealth adherence often observed in people with a low SEP.<sup>33,34</sup> Many participants cited the intervention's integration into daily routines as crucial. Past studies indicate that individuals with a low SEP often face stressful daily challenges, limiting their time and cognitive capacity for engaging with eHealth interventions.<sup>50</sup>

Additionally, curiosity was reported as a key factor in the patient's adherence to the intervention, aligning with the gamification theory that presents curiosity as a strategy to enhance engagement with a system.<sup>51</sup> This facet of the intervention might have been an important contributor to the observed adherence. Regarding acceptance, the intervention's well-received usability contrasts with findings in existing literature. Typically, individuals with a low SEP encounter more challenges with the usability of eHealth interventions.<sup>52–55</sup> The intervention's consistent adherence and positive overall acceptance could be attributed to its participatory design, which followed the leG's recommendations for equitable eHealth development. Failing to achieve adherence and acceptance could negatively influence overall effectiveness, irrespective of any inherent benefits of the intervention.<sup>56</sup> Given our promising outcomes on adherence and acceptance, we recommend future researchers to apply the leG and engage in tailored participatory approaches to develop eHealth interventions for individuals with a low SEP in different settings.

While we did not find a significant intervention effect in this feasibility study on feelings of certainty and guidance or dropout in subsequent CR, we did find some trends pointing towards potential intervention effects. First, the qualitative findings suggest that the participants felt that the intervention contributed to their feelings of certainty and guidance. The interview results suggested improved expectations for future CR, better information, and guidance during the waiting period. These insights hint at the intervention enhancing participant readiness and motivation for CR. Second, the finding that the length of the waiting period was negatively associated with the change in feelings of certainty and guidance in the control group but not in the intervention group suggests that the intervention could serve as an emotional buffer for patients facing longer waiting periods. Qualitative feedback further supports this, with participants reporting that the intervention helped to set expectations and provide information regarding their rehabilitation journey. Although it did not directly improve certainty and guidance, the intervention might have fostered a sense of readiness for rehabilitation by giving information and early engagement with the programme. In future versions of the intervention, its content should be focused more directly on improving the patient's feeling of certainty and guidance. Lastly, although the difference in dropout rates between the intervention and control group was not significant, the 10% dropout rate in the control group is consistent with the general dropout rate in CR.<sup>57</sup> The absence of dropouts in the intervention group could suggest that the intervention may have boosted participant's commitment to CR. This should, however, be confirmed in a sufficiently powered trial.

Our study found participants preferred more personally relevant content and additional depth and detail in information. This suggests that the intervention's one-size-fits-all approach may not meet varying needs for content depth and relevance. This desire for personalized content also aligns with previous research findings.<sup>58,59</sup> Personalized information, as opposed to generic information, has demonstrated a greater positive impact on well-being,<sup>60</sup> health plan decision-making,<sup>61</sup> and lifestyle behaviour.<sup>62</sup> Within CR, several studies have shown to be effective that employed dynamic personalization techniques, such as using initial screenings<sup>63</sup> or artificial intelligence algorithms to adapt the content and delivery in real-time based on user's interactions and responses.<sup>60,64</sup> Future research could explore developing personas or patient profiles reflecting diverse content needs based on health concerns, condition severity, and motivation.<sup>65</sup> These profiles would guide

the creation of tailored pathways for pre-CR content, accommodating different patient types during their waiting period. Pathways may vary by exercise difficulty aligned with disease severity and information delivery adjusted to individual knowledge and health literacy levels.

## Strengths and limitations

This feasibility study lays the groundwork for designing effective interventions for patients with a low SEP during their waiting period before starting CR. Such studies are essential for refining intervention designs to improve impact when scaled up.<sup>66</sup> A strength of our research is the mixed methods approach, which offered insightful explanations for our findings and laid a foundation for future research and development. Additionally, it is noteworthy that we have maintained a participant retention rate similar to other trials conducted at Capri Cardiac Rehabilitation,<sup>67,68</sup> despite our emphasis on people with a low SEP, who are typically underrepresented in these earlier trials.

While our study provides important exploratory insights, the results should be approached with caution and seen as suggestive rather than conclusive. It is important to consider the small sample size and single-centre data collection when interpreting these findings, as the limited sample size particularly affects the robustness of the *P*-values. While our use of a self-designed questionnaire may have affected validity, we mitigated this by basing our questions on established instruments and employing a mixed methods approach to triangulate the quantitative data with qualitative data.<sup>69</sup> Additionally, excluding a participant who dropped out from the analysis could further limit our study's integrity. We recommend validating our results in a larger, more robust trial, with validated instruments, conducted across multiple rehabilitation centres.

Another potential limitation of our study lies in the composition of our participant sample, which, due to neighbourhood-level sampling, possibly included individuals with higher SEP. Although our data indicate a low percentage of highly educated individuals within our sample, this metric alone may not provide a comprehensive picture of SEP. Socioeconomic position is a multi-faceted concept influenced by various factors beyond formal education levels. Additionally, our recruitment approach might have favoured those more experienced with digital tools and comfortable in their current situation. Additionally, the interpretability of our results may have been affected by technical difficulties encountered during the early phase of the study, resulting in some participants not receiving messages for a few days. While this issue was promptly resolved, it might have influenced acceptance and adherence scores. As we plan for a larger, more robust trial, it is crucial to thoroughly test the intervention's technical functionality before its commencement.

Finally, a notable limitation of our study concerns the interpretability of the CGQ scores, particularly due to the timeframe of the intervention and its overlap with interactions (e.g. scheduling appointments and intake sessions) in both the intervention and control groups at the CR facility. Most of these facility interactions occur in the final weeks, coinciding with the period when we evaluated the intervention's effect. Moreover, the duration of intervention use, approximately 2 weeks, was relatively brief when contrasted with the average waiting period of 8 weeks in the Capri Cardiac Rehabilitation centre. These limitations might explain the discrepancy between the quantitative and qualitative findings. For future research, initiating the intervention immediately at hospital discharge would be beneficial, thereby exposing patients during the entire waiting period. The final measurements could be conducted before interactions with healthcare providers at the CR centre to minimize their influence on feelings of certainty and guidance.

## Conclusions

The developed eHealth intervention was well adhered to and accepted by the target group. Yet, usefulness should be improved, and we did not

find effects on feelings of certainty and guidance or dropouts. Despite this, the findings from this feasibility study yield important insights into the design of eHealth interventions tailored to people with a low SEP. Through further optimization, for example through personalization and an extended timeframe for offering the intervention, the intervention holds promise as an effective tool to enhance participation in CR and improve adherence among patients with a low SEP, thereby mitigating health disparities in CR and improving its effectiveness. While researchers should acknowledge the limitations of this feasibility study, including its small sample size and focus on a single centre, it represents a first step towards equitable eHealth interventions. Healthcare professionals and intervention developers can leverage these findings to develop and tailor interventions that align with the needs and preferences of individuals with a low SEP, thereby improving their adherence and acceptance.

## Supplementary material

Supplementary material is available at *European Heart Journal – Digital Health*.

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## Author contributions

J.S.F., J.J.K., N.H., V.T.V., and R.J.G.B contributed to the conceptualization and methodology of the work. J.S.F. performed the investigation, project administration, data curation, and analysis and drafted the manuscript. J.J.K., R.J.G.B., V.T.V., H.B.J.B., A.W.M.E., and N.H.C. contributed to the funding acquisition for the project. All authors were actively involved in critically revising the manuscript, provided final approval of the version to be published, and agreed to be accountable for all aspects of the work to ensure its integrity and accuracy.

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## Data availability

Data available on request.

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