# Cost-Utility Analysis of a Cardiac Telerehabilitation Program: The Teledialog Project

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This study has been registered at ClinicalTrials.gov with clinical trial registration number NCT01752192.

### Abstract

Background: Cardiac rehabilitation can reduce mortality of patients with cardiovascular disease, but a frequently low participation rate in rehabilitation programs has been found globally. The objective of the Teledialog study was to assess the cost-utility (CU) of a cardiac telerehabilitation (CTR) program. The aim of the intervention was to increase the patients' participation in the CTR program. At discharge, an individualized 3-month rehabilitation plan was formulated for each patient. At home, the patients measured their own blood pressure, pulse, weight, and steps taken for 3 months. Materials and Methods: The analysis was carried out to-

gether with a randomized controlled trial with 151 patients during 2012-2014. Costs of the intervention were estimated with a health sector perspective following international quidelines for CU. Quality of life was assessed using the 36-Item Short Form Health Survey. Results: The rehabilitation activities were approximately the same in the two groups, but the number of contacts with the physiotherapist was higher among the intervention group. The mean total cost per patient was €1,700 higher in the intervention group. The qualityadjusted life-years (QALYs) gain was higher in the intervention group, but the difference was not statistically significant. The incremental CU ratio was more than €400,000 per QALY gained. Conclusions: Even though the rehabilitation activities increased, the program does not appear to be cost-effective. The intervention itself was not costly (less than €500), and increasing the number of patients may show reduced costs of the devices and make the CTR more cost-effective. Telerehabilitation can increase participation, but the intervention, in its current form, does not appear to be cost-effective.

Key words: cost-utility, heart patients, telerehabilitation, randomized study, economic evaluation

## Introduction

ardiovascular diseases (CVD) is the cause of 30% of all deaths in 2010. Cardiac rehabilitation (CR) has documented a 20% reduction in all-cause mortality and a 27% reduction in cardiac mortality in systematic reviews. A CR program is defined as a secondary preventive program that includes interventions such as exercise and patient education on risk factors in order to achieve and maintain a healthy lifestyle. Globally, low participation rates for patients in CR programs have been reported. Cardiac telerehabilitation (CTR) is defined as CR using information and communication technology. CTR appears to be an effective strategy to increase both patients' attendance in CR and long-term adherence to recommendations, as well as being in

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particular a promising new tool to include patients who are not able to attend such a program for various reasons.<sup>6,7</sup> However, there is a lack of multidisciplinary evaluations of CTR programs, including health economic studies.<sup>6</sup>

This study conducted a cost-utility (CU) analysis (CUA) of the Teledialog research project. The aim of the project was to develop and test an individualized CTR program designed to increase participation in rehabilitation, improve patient quality of life, reduce the number of admissions, and reduce the need for acute care. As part of the Teledialog project, a CTR program was developed and tested in a randomized controlled trial of cardiac patients diagnosed with artery sclerosis, coronary artery bypass surgery, valve surgery, and heart failure.

## **Materials and Methods**

## THE INTERVENTION

A CTR program was developed and tested in 2012–2014 in a randomized controlled trial (n=151). Two hospitals, four municipal healthcare centers, and a call center took part in the program. On being discharged from hospital, the cardiac patients were enrolled by a project nurse, who included the patients based on various inclusion and exclusion criteria ( $Table\ 1$ ).

CRITERIA	SPECIFICS					
Inclusion	Women and men above 18 years of age					
	Must have signed the "Informed Consent" document					
	Must be able to understand the study information					
	Must live in Hjoerring or Frederikshavn Municipality					
	Must have a mobile network					
	Must be able to use IT or have a close family member who car use IT					
	<ul> <li>Patients with clinical diagnosis of artery sclerosis (myocardial infarction, angina pectoris), coronary artery bypass surgery, valv surgery, and heart failure</li> </ul>					
Exclusion	Patients who, in the investigator's judgment, will not be able to participate in the study					
	Lack of ability to speak and understand Danish					
	Pregnancy or nursing					
	Neurological disease					
	Use of wheelchair/lack of ability to walk					
	Participation in other studies that could influence the outcome of this study					

The project nurse performed an interview with each patient and identified the patient's need for rehabilitation and the place where rehabilitation would be carried out (hospital, healthcare center, or call center). The important part of the CTR was to design an individual rehabilitation based on the patient's needs.

The patient was registered as a user in the information technology platforms and was equipped with a tablet computer giving him or her Web access to his or her own personal health record and measurements. The patient received training in the use of the different devices and the digital rehabilitation plan, and a doctor prescribed how often the patient needed to measure blood pressure, pulse, and weight, most often twice a week, whereas steps were measured every day. The data were transmitted via a secure transmission line. Patients, relatives, and healthcare professionals from the hospital and healthcare centers were able to communicate and share data from the personal health record. In addition, patients and relatives had access to Activeheart.dk (a digital toolbox with information on rehabilitation topics, activities, and videos showing patients describing their experiences of being a heart patient and appropriate exercises for after surgery).

After 2 weeks in the program, each patient was visited by a project assistant in order to make sure that the patient and relative were confident in using the technology. Healthcare professionals monitored the measured values every week, contacted the patients if the values were abnormal, and discussed rehabilitation activities with the patient. The telerehabilitation program lasted for 3 months. *Table 2* gives further description of the equipment.

The control group followed a traditional rehabilitation program at the hospital or healthcare center based on CR guidelines. Aalborg University was responsible for the delivery, installation, and removal of the equipment for the patients in the intervention group.

## RANDOMIZATION

When the patient had signed the informed consent form, a project nurse performed the randomization. Randomization into groups was done using an automatically generated list for each block with an equal number of intervention and control numbers in random order, so as to ensure an even distribution in each group in varying sizes (10, 20, and 14). In total, 151 cardiac patients were enrolled.

## **OUTCOME MEASUREMENT**

Quality of life was assessed using the 36-Item Short Form Health Survey (SF-36). To estimate the changes in the number

EQUIPMENT	USAGE							
MyMedic	MyMedic is a telehealth monitor produced by Tunstall Healthcare A/S that transfers data via a mobile Internet connection to a central server. Devices like the sphygmomanometer, digital weight scale, and ECG are transmitting data via the telehealth monitor. Leasing of the telehealth monitor and devices is €33.51 per patient.							
Triage Manager	The Triage Manager is a software administrative module used to administer data on the patients being monitored.  The Triage Manager is provided by Tunstall Healthcare A/S and used by the healthcare professionals.							
Sphygmomanometer	Approved for medical use. The meter was paired with the telehealth monitor in advance.							
Digital weight scale	Approved for medical use. The weight scale was paired with the telehealth monitor in advance.							
ECG	The ECG is CE-approved and approved for medical use. It measures the first derived ECG scan for 40 s and then transmits the measured values to the telehealth monitor via wireless Bluetooth.							
Fitbit Ultra	The Fitbit Ultra is a digital pedometer registering the number of steps. The patients with access to their data can follow their data on the telehealth monitor or at shared care. Cost is €30.83 per patient.							
ActiveHeart.dk	An interactive Web page with information on health, the heart, and heart diseases communicated using text and videos. The videos have instructions on exercises and short narratives by patients and relatives about their experiences with daily life after being diagnosed with heart disease. On the Web page, the patients have the possibility to call SOS International which administers a hotline for the heart patients. The hotline is open daily between 9:00 to 16:00 h, and nurses are available within this time frame for any questions from the patients. Finally, there is a Web forum, where patients can interact with each other. The Web page was available for patients in the intervention group and their relatives. The Web page has been designed to suit patients' different learning styles. The information is available 24/7. Cost is €72.97 per patient.							
Shared Care	Shared Care is an Internet-based rehabilitation plan delivered by IBM. The patients, relatives, and healthcare professionals have access to the plan and can obtain an overview of patient data, such as rehabilitation plan, GP, diary, medications, activities, monitored values, etc. Was available free in the project							
CareConnect	CareConnect is a data platform provided by KMD for integrating and connecting the different systems in the project.  CareConnect receives data from Medcom correspondence, MyMedic, Fitbit, and Shared Care. Cost is €5.36 per patient.							
Tablet	Each patient had an Android tablet (Samsung) to access data. The patients had the table for 3 months. The tablet was reused for more patients. Cost is €5.36 per patient.							

of quality-adjusted life-years (QALYs), the patients' responses to the SF-36 have been transformed using the Brazier algorithm and the SF-6D methodology.<sup>8</sup> This routine applies a nonparametric model to estimate SF-6D health state utility values using Bayesian methods.<sup>9</sup> The SF-6D comes with a set of preference weights obtained from a sample of the British population using the standard gamble method.<sup>9</sup> Patients who complete the questionnaires may be uniquely classified in terms of QALY values according to the SF-6D.<sup>8</sup>

#### COST MEASUREMENTS

The CUA is based on a health sector perspective. The economic evaluation follows international guidelines for conducting a CUA of a randomized controlled trial.<sup>10</sup> All prices and unit costs are in euros, calculated from Danish kroner (DKK) at the exchange rate of €100=DKK 746.<sup>11</sup>

Only direct healthcare costs are considered (i.e., costs related to the intervention [equipment, training by professionals] and to the use of healthcare [hospital admissions, ambulatory contacts, general practitioner contacts, emergency physician contacts, and other primary services]). Data on costs of the intervention, including prices of equipment used in the intervention, were obtained from companies and from worktime records of the clinical staff. Data on use of healthcare were collected for each patient from the National Patient Register and is measured in the years 2013–2014. Estimates of prices of use of inpatient and outpatient hospital care for each patient were derived from the National Patient Register and based on Diagnose Related Groups from the Danish reimbursement system.

## ANALYSIS OF DATA

The statistical analysis of data is based on the intention-to-treat method. Data from the SF-36 were entered into EpiData (version 1.4.1) and transferred using EpiData Manager (version 1.3.2.1.) to a statistical package (R version. 3.1.2) for analysis. The intervention and control groups were compared using descriptive statistics and tested by t test.

These tests are applied to approximately normal data as baseline characteristics and QALY measures. Data on use of resources and costs are skewed and do not follow a normal distribution. This means that the groups are compared by their means but tested using the nonparametric Mann–Whitney U test for differences. Statistical significance was defined at a 5% significance level. The results are presented with means for all variables included in the economic evaluation.

#### ETHICAL PERMISSION

The project was approved by the Danish Ethical Committee (number 20120051). The ethical principles for medical research involving human subjects, described in the Helsinki Declaration and the Act on Processing of Personal Data, were followed.

## **Results**

Altogether, 369 patients were screened (*Fig. 1*), and 151 patients were eligible and randomized. Baseline data were collected for 141 of the eligible patients (72 for the intervention group and 69 for the control group). Attrition was acceptable, as 88% of the patients completed the 6-month follow-up assessment (90% from the intervention group and 83% from the control group), and 84% completed the 12-month follow-up (89% from the intervention group and 80% from the control group).

*Table 3* shows the patients' characteristics at baseline. Tests of association conducted for each variable across randomization group confirmed that baseline characteristics did not differ statistically.

## **ESTIMATED COSTS**

Data about the use of resources to carry out the intervention and the use of rehabilitation and healthcare services in the intervention and control groups is shown in *Table 4*.

With regard to rehabilitation services, we found that those in the intervention group used the physiotherapist an average of 5 times and the team training an average 2.2 times; among the control group the corresponding figures were 2.9 and 3.4, respectively. However, these differences are statistically insignificant. Generally, the use of healthcare services is higher in the intervention group except for contacts to emergency doctors. However, the average number of contacts in the control group may be severely skewed due to an outlier in the control group who had 27 contacts, and the nonparametric test confirms that the difference is not significant.

The estimated mean costs per patient are presented in *Table 5*. The intervention in itself costs  $\leq$ 392 per patient, including both the costs of devices and staff resources. In the

intervention group, the overall costs are for rehabilitation services about €100 higher and for healthcare services about €1,200 higher, but neither difference is statistically significant. Nevertheless, the cost of reimbursed transportation to rehabilitation is significantly higher in the control group.

The total costs of healthcare utilization are  $\in$ 1,200 higher on average in the intervention group, although not statistically significant. This is due mainly to the higher number of admissions. However, the total average cost per patient in the intervention group is significantly higher ( $\in$ 5,709) compared with the control group ( $\in$ 4,045), although this difference may be due to the added cost of the intervention.

#### **ESTIMATED OALYS**

The reported values in *Table 6* are QALYs derived from the respondents' answers to SF-36 during the project. With an increase of 0.089 QALY in the intervention group and 0.085 QALY in the control group during the first 12 months, the difference is not significant. However, the estimated QALY show a significant increase over time in both groups.

#### ESTIMATED INCREMENTAL CU RATIO

The incremental cost-effectiveness ratio (ICER) is calculated according to the following formula:

$$ICER = \frac{C_{I} - C_{C}}{QALY_{I} - QALY_{C}}$$

where  $C_i$  (i = I, C) is equal to the mean costs per patient in the two groups, where I represents the intervention group and C represents the control group. The denominator consists of the change in QALY from baseline (0 month) to the 12-month follow-up for both intervention and control groups. Based on the estimates in *Tables 5* and 6, ICER can be estimated to  $\in$ 518,280 per QALY gained. Thus, the estimated total costs of investing in Teledialog are  $\in$ 518,280 per gained QALY.

Uncertainty of the estimated ICER was evaluated with probabilistic sensitivity analysis using a 5,000 bootstrap resample with a 95% bias-adjusted confidence interval (CI). The results of the resample are illustrated in the cost-effectiveness plane in *Figure 2*. Based on the resample, the mean costs per patient in the intervention and control groups were estimated to be €5,724 (CI: 3,709; 953) and €4,057 (CI: 2,445; 689), respectively, and the mean QALY gain to be 0.089 (CI: 0.05; 0.13) and 0.085 (CI: 0.05; 0.12), whereas the mean ICER was €483.608.

Several sensitivity analyses were carried out to examine the robustness of the results. First, patients with very high costs (costs of more than €33,500) were excluded (4 patients). This

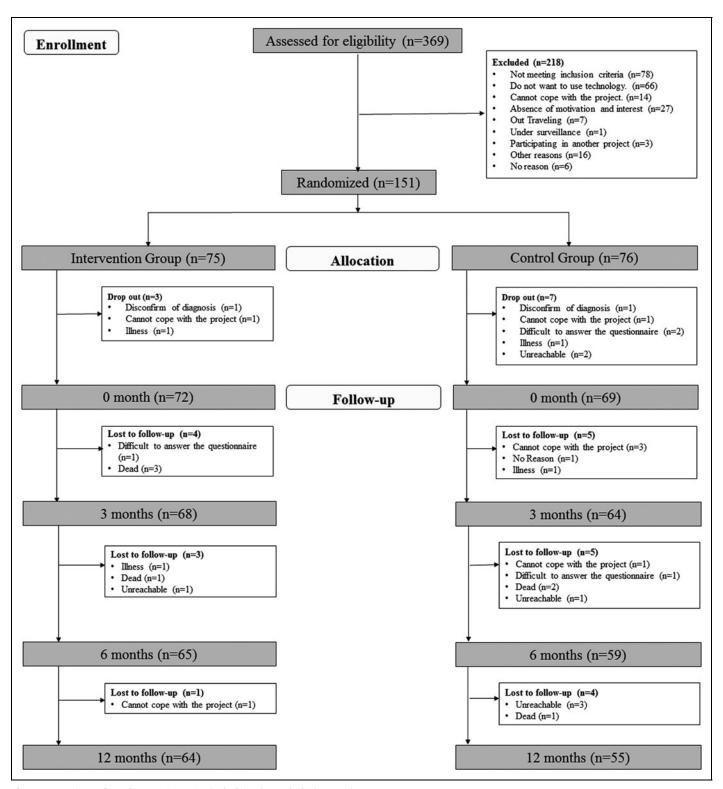


Fig. 1. Overview of cardiac patients included in the Teledialog project.

VARIABLE	TELEREHABILITATION GROUP (N=72)	CONTROL GROUP (N=69)	P VALUI
Age (years) by gender			
Men	63.890 ± 11.650 (31–88) [57–72] (n=56)	61.270±11.780 (39–83) [53–70] (n=55)	0.241
Women	57.440 ± 14.940 (34–84) [49.500–72.250] (n = 16)	68.140±10.110 (52-85) [60-78] (n=14)	0.032
Men and women	62.460 ± 12.310 (31–88) [54–72]	62.670±11.720 (39–85) [54–72]	0.919
Clinical parameters			•
Weight (kg)	86.440 ± 18.160 (43.900-140.000) [76.350-98.230]	85.800 ± 20.350 (51.000-140.000) [71.000-99.000]	0.846
Blood pressure (mm Hg)		,	1
Systolic	125.690 ± 15.730 (85–170) [115.000–138.000]	125.660±15.150 (95-169) [115.500-135.000]	0.990
Diastolic	75.670±11.100 (50-115) [70.000-80.000]	73.060 ± 11.780 (40–103) [63.000–80.500]	0.185
Heart rate (beats/minute)	70.860±13.880 (50-110) [60.000-77.750]	70.740 ± 14.880 (46–111) [60.000–77.500]	0.960
Missing values (n)	2	5	
Primary diagnoses			0.383
Acute coronary syndrome	33	38	
Heart surgery	26	15	
Heart failure	9	9	
Artery sclerosis and heart failure	6	6	
Civil status			0.572
Single	12	9	
Married or living with a partner	58	57	
Missing	2	3	
Education			0.898
Elementary school	21	16	
High school	8	7	
Skilled worker	26	26	
Short education	1	3	
Bachelor's degree	10	10	
Master's degree	3	3	
Missing	3	4	
Work status			0.325
Works under 20 h/week	2	4	
Works 20–36 h/week	1	4	
Works full-time 37 h/week	10	9	
Sick leave	16	12	
Unemployed	0	2	
Retired	36	28	
Missing	7	10	

Table 3. Basic Characteristics of the Telerehabilitation and Control Groups continued								
VARIABLE	E TELEREHABILITATION GROUP (N=72) CONTROL GROUP (N=69)							
Place of rehabilitation			0.560					
Hospital	28	29						
Healthcare center	31	40						
Call center	13	0						

 $Age \ and \ clinical \ parameters \ are \ mean \ \pm \ standard \ deviation \ values \ (minimum-maximum) \ [interquartile \ range]. \ Other \ values \ are \ number \ of \ patients.$ 

	MEAN USE PER PATIENT IN						
		INTERVENTION GROUP			CONTROL GROUP		
TYPE OF COSTS	N	NUMBER	CI (95%)	N	NUMBER	CI (95%)	P VALUE
Resource use (min) in Teledialog							
Staff time (information to patients)		85			60		
Staff time (dialogue with patients at home)		120			-		
Visit at patient's home		45			-		
Collecting and cleaning devices		45			_		
Use of rehabilitation services							
Contacts (n) with							
Nurse	68	2.480	(1.880; 3.090)	66	2.860	(2.230; 3.500)	0.380
Doctor	68	0.710	(0.480; 0.930)	66	0.700	(0.470; 0.920)	0.984
Physiotherapist	68	5.000	(2.800; 7.200)	66	2.890	(1.230; 4.560)	0.351
Dietician	68	0.340	(0.090; 0.580)	66	0.360	(0.120; 0.600)	0.390
Psychologist	68	0.180	(0,000; 0.440)	66	0.200	(0.000; 0.560)	0.976
Participations (n) in team training	68	2.220	(0.830;3.610)	66	3.440	(1.750; 5.130)	0.251
Use of healthcare							
Inpatient days (n)	68	1.020	(0.530; 1.520)	66	0.600	(0.250; 0.950)	0.429
Re-admissions (n)	68	0.490	(0.250; 0.600)	66	0.350	(0.200; 0.500)	0.496
Outpatient visits ( <i>n</i> )	68	4.290	(2.350; 6.230)	66	3.820	(2.190; 5.450)	0.868
Length of stay (days) for each admission	33	3.880	(2.440; 5.320)	24	2.290	(1.200; 3.380)	
Contacts (n) with							
GP	68	15.120	(12.070; 18.160)	66	14.670	(11.230; 18.100)	0.680
Emergency physician	68	0.720	(0.400; 1.040)	66	1.450	(1.130; 1.780)	0.980
Reimbursed transportation visits (n)	68	9.010	(6.160; 11.870)	66	9.270	(6.790; 11.750)	0.301

	INTERVENTION GROUP				CONTROL GROUP		
TYPE OF COSTS	N	MEAN COST	CI (95%)	N	MEAN COST	CI (95%)	P VALUE
Costs of Teledialog	·						
Time (min)							
Staff time information to patients)		51			36		
Staff time (dialogue with patients)		72		-			
Visit at patient's home		13			_		
Collecting and cleaning devices		13			_		
Devices' cost							
Rent of telemedicine devices (3 months)		101			-		
Rent of tablet		32			_		
Fitbit Ultra		31			-		
Activeheart.dk	79			_			
Total running costs	392			36			
Cost of rehabilitation services	'			•			
Contacts with							
Nurse	68	84	(61; 106)	66	82	(59; 104)	0.978
Doctor	68	16	(10; 20)	66	16	(11; 21)	0.876
Physiotherapist	68	199	(116; 281)	66	92	(9; 176)	0.145
Dietician	68	9	(2; 15)	66	10	(4; 17)	0.456
Psychologist	68	12	(0; 32)	66	10	(0; 30)	0.989
Participations in team training	68	7	(3; 10	66	20	(16; 23)	0.190
Total cost of rehabilitation		326	(232; 419)		229	(160; 299)	0.570
Costs of healthcare services	•						
Admissions	68	4,224	(1,519; 6,927)	66	3,002	(986; 5,019)	0.507
Outpatient visits	68	533	(290; 775)	66	519	(291; 748	0.815
Contacts with	•						
GP	68	179	(139; 218)	66	166	(127; 205)	0.598
Emergency physician	68	12	(6; 18)	66	24	(9; 38)	0.913
Transportation costs	68	43	(22; 65)	66	68	(46; 91)	0.038
Total costs of healthcare utilization		4,991	(2,286; 7,695)		3,780	(1,679; 5,879)	0.936
Total costs		5,709	(2,990; 8,427)		4,045	(1,917; 6,173)	0.0211

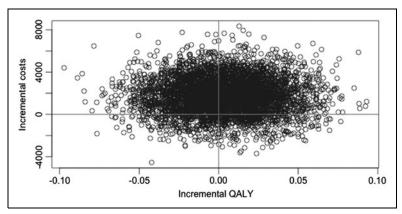
CI, confidence interval; GP, general practitioner.

Table 6. Quality-Adjusted Life-Years Gain in the Intervention and Control Groups									
	BASELINE	3 MONTHS	CHANGE DURING FIRST 3 MONTHS	6 MONTHS	CHANGE DURING FIRST 6 MONTHS	12 MONTHS	CHANGE DURING FIRST 12 MONTHS		
Intervention group	0.610	0.670	0.065	0.680	0.069	0.700	0.089		
Number of patients	61	61	61	61	61	61	61		
Confidence interval	(0.580; 0.640)	(0.640; 0.710)	(0.030; 0.100)	(0.640; 0.710)	(0.030; 0.100)	(0.660; 0.740)	(0.050; 0.130)		
Control group	0.620	0.700	0.081	0.710	0.090	0.710	0.085		
Number of patients	53	53	53	53	53	53	53		
Confidence interval	(0.590; 0.650)	(0.670; 0.740)	(0.040; 0.120)	(0.670; 0.750)	(0.050; 0.130)	(0.670; 0.740)	(0.050; 0.120)		
p value	0.608	0.275	0.512	0.225	0.4353	0.751	0.904		

reduced the difference in the mean costs per patient to  $\{1,011$ . Second, fixed costs related to the investment in telemedicine devices were excluded (costs of rent of telemedicine devices, rent of the tablet computer, purchase of the Fitbit [San Francisco, CA] Ultra tracker, and establishing of the Activeheart.dk Web site) in order to estimate the marginal costs of offering the intervention to additional patients. This reduced the difference in the mean cost per patient to  $\{1,423\}$ . Finally, the mean cost per patient was estimated, excluding the 15 patients for whom 12 months of follow-up data do not exist. This reduced the difference in the mean costs per patient to  $\{1,023\}$ .

## **Discussion**

The aim of the project was to develop and test an individualized CTR program to increase participation in rehabilitation, improve patient quality of life, reduce the number of admissions, and reduce use of acute care. The CUA shows that the intervention group contacted the physiotherapist



**Fig. 2.** Cost-effectiveness plan of incremental cost per quality-adjusted life-year (QALY) (bootstrapping with 5000 replications).

more often while attending fewer group-training sessions compared with the control group, although these differences are not statistically significant. This suggests that patients in the telerehabilitation group are more likely to carry out their rehabilitation activities on an individual basis, thus supporting the main aim of the Teledialog project. Qualitative findings support that patients and healthcare professionals felt that the CTR program in Teledialog was tailored to individual needs. <sup>12</sup>

When examining the use of healthcare in general, the number of emergency contacts to the physician by members of the intervention group is half that of the control group, whereas the numbers of re-admissions, outpatient visits, and general practitioner visits on average are higher in the intervention group, although none of these differences is statistically significant. When comparing the overall average costs across the two groups, we find that the average cost per patient is significantly higher for those in the intervention group,

except for the cost for transport, which is lower in the intervention group than in the control group.

We found no significant differences in the improvement of the patients' quality of life between the two groups ( $\Delta$ =0.004), and, taken together, these results indicate that the intervention is not costeffective, with costs per QALY gained at more than €500,000. However, it is possible that the added costs of telerehabilitation may result in other desirable outcomes that were not considered in the current study. The number of cardiac patients not participating in rehabilitation is large, and these patients could benefit from a telerehabilitation approach, although at a greater cost, which may prove to be a better outcome than nonparticipation. Widmer et al. <sup>13</sup> concluded, in a

systematic review and meta-analysis of digital health interventions for the prevention of CVD, that digital health interventions can reduce CVD outcomes and have a positive impact on risk factors for CVD.

In addition, the continuous collection of clinically relevant data carried out using the telerehabilitation approach may provide better support to the physician in offering improved treatment recommendations, as such clinical judgments would be based on ongoing measurements compared with the more static measures obtained from one-off, face-to-face consultations.

The strengths of this study are the wide healthcare perspective, including both primary and secondary care costs. We have not identified any study where costs from the primary and secondary sectors have been included at the same time. In addition, this study has used mainly data from patient registers rather than self-reported patient data and has had a follow-up period of 1 year.

We have identified only three other CU studies of a telerehabilitation program for patients with CVD. Körtke et al. 14 presented a cost-effectiveness analysis of a nonrandomized trial of a transtelephonic guide for ambulatory rehabilitation in cardiac surgery patients. Their study showed that the total cost of the telemedicine program was lower than the conventional inhospital program. Kraal et al. 15 described a protocol for a forthcoming study of a 12-week telemonitoring-guided home-based training program. Frederix et al. 16 conducted a CUA of a 24-week comprehensive CTR program using a sample of 140 patients. The study showed that the mean cost per patient in the telemonitoring program was €564 lower than the mean costs in the control group. At the same time, the mean QALY gain per patient in the telemonitoring group was 0.026, and the conclusion therefore was that CTR tends to be more effective and efficient than centerbased CR alone. The main reason for the result was a statistically significant reduction in the number of days lost due to cardiovascular rehospitalizations in the intervention group.

The Teledialog study has several limitations. First, the project was a small-scale research project. Only healthcare costs were included in the estimation, and potential savings for the patient (e.g., savings due to a lower degree of use of pharmaceuticals) and for the municipality related to the possible improvement in the patients' quality of life were not taken into account. Using a higher number of patients may thus result in reduced costs of the devices used in the intervention and make the CTR program more cost-effective over the long term.

Second, we found an unexpected difference in the costs of outpatient and inpatient visits. Only hospital visits related to heart disease are included in the study, and based on the estimated low impact on the patients' quality of life, no difference in these costs was to be expected. Therefore, only a few

patients with high healthcare costs unrelated to the degree of rehabilitation may have had a large impact on the costs. As a consequence, larger studies are needed to provide further examination of the differences in costs per patient.

Third, the measure of the patients' quality of life, the Brazier algorithm, and the SF-6D methodology may be criticized for being too generic and not sensitive enough for identifying the changes in the patient's health related to rehabilitation.

### **Conclusions**

The current study shows that an individualized CTR program may increase the patient's participation in rehabilitation activities, reduce the cost of transport, and reduce the number of emergency physician visits. However, the mean total healthcare cost per patient in the intervention group was significantly higher, and improvement in the patients' quality of life was not significant, compared with the costs and quality of life of the patients in the control group. Taken together, these findings indicate that the telerehabilitation approach is not costeffective in a small-scale study. It is possible that increasing the number of patients in the sample may show reduced costs of the devices and make the telerehabilitation program more costeffective. In addition, increasing rehabilitation participation and perhaps the kind of clinically relevant data collected may give the telerehabilitation approach an additional advantage. More research is needed to identify the potential for the use of telerehabilitation for cardiac patients.

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## **Disclosure Statement**

No competing financial interest exists.

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