# **BMJ Open** Effect of a real-time feedback smartphone application (TCPRLink) on the quality of telephone-assisted CPR performed by trained laypeople in China: a manikin-based randomised controlled study

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## ABSTRACT

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**Objectives** To determine the effect of a free smartphone application (TCPRLink) that provides real-time monitoring and audiovisual feedback on chest compressions (CC) on trained layperson telephone-assisted cardiopulmonary resuscitation (T-CPR) performance.

Design A manikin-based randomised controlled study. Setting This study was conducted at a multidisciplinary university and a community centre in China.

Participants One hundred and eighty-six adult participants (age 18-65 years) with T-CPR training experience were randomly assigned to the TCPRLink (n=94) and T-CPR (n=92) groups with age stratification. Interventions We compared the participants performance for 6 min of CC in a simulated T-CPR scenario

both at the baseline and after 3 months.

Primary and secondary outcome measures The primary outcomes were the CC rate and proportion of adequate CC rate (100–120 min<sup>-1</sup>). The secondary outcomes included the proportion of participants counting the CC rhythm, time to first CC, CC depth, hands-off time and CC full-release ratio.

**Results** Participants in the TCPRLink feedback group more consistently performed CC with higher rate, both initially and 3 months later (median 111 (IQR 109-113) vs 108 (103–112) min<sup>-1</sup>, p=0.002 and 111 (109–113) vs 108 (105-112) min<sup>-1</sup>, p<0.001, respectively), with less need to count the rhythm (21.3% vs 41.3%, p=0.003% and 7% vs 22.6%, p=0.004, respectively) compared with the T-CPR group. There were no significant differences in time to the first CC, hands-off time or CC full-release ratio. Among 55-65 year group, the CC depth was deeper in the TCPRLink group than in the TCPR group (47.1±9.6 vs 38.5±8.7 mm, p=0.001 and 44.7±10.1 vs 39.3±10.8 mm, p=0.07, respectively).

Conclusions The TCPRLink application improved T-CPR quality in trained laypersons to provide more effective CCs and lighten the load of counting out the CC with the dispatcher in a simulated T-CPR scenario. Further investigations are required to confirm this effectiveness in real-life resuscitation attempts.

# Strengths and limitations of this study

- ▶ The effectiveness of a real-time feedback smartphone application (TCPRLink) was evaluated in a telephone-assisted cardiopulmonary resuscitation (T-CPR) simulation among participants from the Chinese general population.
- Trained adult laypersons (age range 18-65 years) participated in this study to facilitate the identification of discrepancies in T-CPR performance among different age groups.
- The study included a 3-month follow-up T-CPR per-formance test to investigate the participants' skill retention.
- The Hawthorn effect could not be excluded in the simulation scenario, with the possibility of a motivation bias.

# INTRODUCTION

Bystander-provided immediate and adequate cardiopulmonary resuscitation (CPR) can directly impact patient outcomes following an out-of-hospital cardiac arrest.<sup>1–3</sup> The updated guidelines of the American Heart Association (AHA) and European Resuscitation Council (ERC) state that telephone-assisted CPR (T-CPR) has a positive effect on the entire resuscitation process by getting more callers to start CPR and through coaching the callers to provide effective CPR.45 Despite significant advances in the T-CPR instructions during the resuscitation procedures, here exists a blind zone between the dispatcher and caller. The dispatcher is voice connected to the caller via the phone, but is unable to see the patient and evaluate the quality of bystander CPR. Therefore, new strategies to address this challenge are needed.

The ubiquitous presence and utilisation of smartphones suggest a novel opportunity to improve resuscitation care through the measurement of bystander CPR metrics.<sup>6–12</sup> In a recent statement from the AHA and ERC, the use of digital strategies, such as mobile devices, was encouraged to provide bystanders with an accelerometer to measure CPR metrics.<sup>13 14</sup> In adherence to these guidelines, an audiovisual smartphone application (TCPRLink) was developed to facilitate high-quality bystander-provided CPR and assist the dispatcher to evaluate the CPR quality in real time.<sup>15</sup> The TCPRLink application utilises the smartphone front camera to detect chest compressions (CC) and displays the CC rate to the bystanders and simultaneously sends the real-time CC rate and the time without compressions via the internet to a monitor that is in front of the dispatcher.

This study was conducted to evaluate the effectiveness of the TCPRLink application with real-time audiovisual feedback in dispatcher-assisted CPR during a cardiac arrest simulation. We hypothesised that this smartphonebased CC rate feedback application would improve the quality of CPR in the general population compared with the use of conventional T-CPR instructions.

#### **METHODS**

#### Study design and ethics

This study was a simulation-based randomised experimental trial that was carried out from 1 September 2018 to 30 May 2019. All participants were verbally informed about the purpose of the study and provided written informed consent. They were informed that their T-CPR performance would be tested and video-recorded in a simulated scenario after training and, again, 3 months later.

# **Study population**

We randomly recruited 186 participants from those who participated in the 'WeCan CPR' training programme<sup>16</sup> an initiative of the China Resuscitation Academy. College

students and adult laypersons (age range 18–65 years) who had completed the training programme within 1 week were eligible for study enrolment. Physicians, nurses, dispatchers and other healthcare professionals were excluded from the study.

The WeCan CPR course is a video-based, 1-hour training programme on applying dispatcher-telephoneguided CPR training in combination with practical and basic CPR training that is targeted at potential bystanders. Participants learn how to call the emergency dispatch centre, follow the procedure of the T-CPR instructions and perform hands-only CPR. All trainees performed at least 550 CC on instrumented feedback manikins (QCPR Classroom, Laerdal Medical, Norway) during the training.

## Patient and public involvement

Patients or the public were not involved in the design, or conduct, or reporting or dissemination of our research.

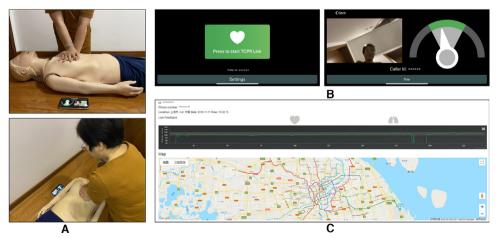
### **Randomisation**

Randomisation was stratified by age groups (18–24, 25–54 and 55–65 years) and conducted to ensure equal distribution of participants across study arms. Participants were randomised into either the control arm (conventional T-CPR group) or interventional arm (T-CPR with the TCPRLink group). All participants were informed the purpose of the study, which was to assess the impact of the TCPRLink App on resuscitation performance, and were not blinded to the study-arm allocation due to the nature of the intervention.

## **TCPRLink application**

TCPRLink (University of Stavanger and Laerdal Medical, Norway) is a free, CPR audiovisual feedback smartphone application that was designed to measure the CC rate and hands-off time and to provide feedback to the bystander and the *dispatcher*. The accuracy and validation of the TCPRLink app has been demonstrated earlier.<sup>17</sup>

The illustration of the application in use is presented in figure 1. By clicking the 'Press to start TCPR Link'



**Figure 1** The illustration of TCPRLink application in use. (A) Illustration photo of TCPRLink in use in a simulated T-CPR situation. (B) Screenshots of TCPRLink. Front page to the left and bystander feedback example to the right. (C) Screenshot of the web server available for the dispatcher. T-CPR, telephone-assisted cardiopulmonary resuscitation.

button, the application activates the speaker, establishes a telephone connection with the dispatcher, activates the TCPRLink app which captures and analyses the CPR movement via the front facing camera of the smartphone in real time, and simultaneously sends the location and real-time compression data to a web server which is available for the dispatcher (web server: http://tcprlink. azurewebsites.net/?\_country=china).

At the bystander interface, a speedometer displayed on the smartphone screen next to a preview frame allows the bystander to keep track of the CC rate, which is obtained by analysing body movement. Thus, the individual receives real-time objective feedback via the speedometer (with the indicator in the green or yellow range of 100–120 and <100 or >120 compressions/min, respectively).

Similarly, at the dispatcher's interface, real-time objective feedback is presented during the emergency call via a sliding window from a website presented on a computer screen that shows the history and progression of the CC rate. Guided by the indicator on web server, the dispatcher can further guide the bystander-rendered CC rate through direct instructions to 'push faster', 'push slower' or 'don't stop'.

# **Study procedures**

The T-CPR performance of all participants was evaluated twice. The first evaluation (phase I test) was conducted within 1 week of WeCanCPR training in a cardiac arrest T-CPR simulated scenario, and the second occurred 3 months later (phase II test) and corresponded to the same setting as the initial test.

The simulations were performed in a quiet, isolated, designated room with a manikin placed on the floor. Individuals were asked to enter the room alone, make an emergency call to an assigned phone number and try their best to rescue the manikin in a cardiac arrest T-CPR simulated scenario. T-CPR instructions were strictly standardised using the Medical Priority Dispatch System (MPDS V.12.1, Salt Lake City, USA) out-of-hospital cardiac arrest (OHCA) dispatch protocol.<sup>18</sup> One dispatcher who had 6 years of T-CPR experience from working at the local emergency dispatch centre acted as dispatcher in the simulation.

During T-CPR calls, individuals were asked for their current address, patient's age and gender, patient's consciousness level and breathing status in accordance with the MPDS protocol. Then, individuals were instructed by the dispatcher to activate the speaker and place their phone on the floor by the manikin. The dispatcher followed a standard procedure to initiate CPR and let the participant perform hands-only CPR for 6 min. For encouragement, the dispatcher counted the CC rhythm with the participants and said 'good job, push harder' every 30s during the simulation.

For the conventional T-CPR group, the participants received no visual feedback from the smartphone and were guided only by the dispatcher instructions. For the TCPRLink group, individuals were asked to call for help using the TCPRLink app. The participants' behaviour and performance during the simulation exercise were recorded by a separate video camera that faced towards the manikin and was located 80 cm above the ground and 1.5 m away for a panoramic shot.

## **Outcome measures**

The primary outcomes measured were the CC rate and the proportion of the adequate CC meeting the guidelinerecommended rate  $(100-120 \text{ min}^{-1})^{19}$  <sup>20</sup> during 6 min of hands-only CPR. The secondary outcomes were CC depth, the proportion of CC with the adequate CC depth (5–6 cm), the proportion of CC with complete recoil (complete release recoil of the chest between compressions) and the absolute hands-off time (the sum of all periods during which there was no hand compression of the chest) during the 6 min of hands-only CC. The abovementioned parameters of CCs effectiveness were monitored using the proprietary software for the ResusciAnne QCPR manikin (Laerdal Medical, Norway).

The video recording of the simulation scenario was used to evaluate individual participant behaviours, including the communication with the dispatcher (counting the CC rhythms with the dispatcher) and time to first CC (time interval from call connection to first CC). We documented the age, sex, education level, self-reported body weight and height of all participants.

## Sample size estimation

The sample size calculation was followed to sequentially recruit 68 participants (34 in the TCPRLink group with 12, 11 and 11 participants in the 18-24, 25-54 and 55-65 years age range, respectively, and 34 in the T-CPR group with 11, 12 and 11 participants in the 18-24, 25-54 and 55-65 years age range, respectively) in the phase I test. A change in the proportion of adequate CC by >5% was considered to be a relevant difference. With a statistical power of 90% and two-sided alpha level of 0.05, the minimum numbers of participants required in the TCPRLink/T-CPR group among the different age groups were 20 (18-24 years), 26 (25-54 years) and 18 (55-65 years), respectively. Considering the possibility of 20% loss to follow-up and the participants' availability, we recruited 54, 75 and 57 participants in the age ranges of 18-24, 25-54 and 55-65 years, respectively.

## **Statistical analysis**

Data are presented as frequencies with percentages for categorical variables and mean±SD or median (IQR; M ( $P_{25}-P_{75}$ )) for continuous variables. Normal distribution was confirmed using the Kolmogorov–Smirnov test. Intergroup differences in the outcomes for the categorical variables were assessed using the  $\chi^2$  or Fisher's exact test. Independent Student's t-tests were conducted to explore the effect of the intervention for continuous variables with normal distribution and Mann–Whitney U-test was used for variables with non-parametric distribution between the control and intervention arm. All analyses

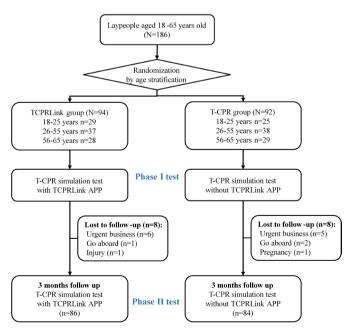
Table 1 Demographic characteristic	s of the participants		
	Total (n=186)	TCPRLink group (n=94)	T-CPR group (n=92)
Male (n, %)	83 (44.6)	42 (44.7)	41 (44.6)
Age, years			
18–24	54 (27.0)	29 (30.9)	25 (27.2)
25–54	75 (37.5)	37 (39.4)	38 (41.3)
55–65	57 (28.5)	28 (29.8)	29 (31.5)
Education status (n, %)			
≤High school/junior college	68 (36.6)	30 (31.9)	38 (41.3)
College	75 (40.3)	44 (46.8)	31 (33.7)
Masters and PhD	43 (23.1)	20 (21.3)	23 (25.0)
Height, m, mean±SD	1.68±0.1	1.67±0.1	1.68±0.1
Weight, kg, mean±SD	64.5±11.4	63.3±10.3	65.6±12.4
BMI, kg/m <sup>2</sup> , mean±SD	22.9±3.1	22.7±2.8	23.0±3.4

BMI, body mass index; T-CPR, telephone-assisted cardiopulmonary resuscitation.

were conducted using SPSS V.22.0. All p-values were two-sided, and p<0.05 was considered to be statistically significant.

## RESULTS

A total of 186 participants (94 in T-CPR with TCPRLink group and 92 in conventional T-CPR group) were included in this study. The demographic characteristics are shown in table 1. Age, gender, education level and body mass index did not differ between the groups. Eight participants in each study arm were lost to follow-up after the initial test (figure 2).



**Figure 2** Flow diagram of the participants. T-CPR, telephone-assisted cardiopulmonary resuscitation.

During the 6min of hands-only CPR, individuals in the TCPRLink group performed CC with a higher rate, both initially (median 111 (IQR 109–113) vs 108 (103– 112) min<sup>-1</sup>, p=0.002) and at the 3-month retest (111 (109–113) vs 108 (105–112) min<sup>-1</sup>, p<0.001), compared with the conventional T-CPR group, respectively (table 2 and figure 3). In the TCPRLink group where the CC rate speedometer was displayed, individuals were less likely to count out the CC rhythms with the dispatcher (21.3% vs 41.3%, p=0.003% and 7% vs 22.6%, p=0.004, respectively) (table 2 and figure 4). Hands-off times, CC fullrelease ratio and time to first CC did not statistically differ between the study groups either initially or at 3 months follow-up.

The depth of CCs in the TCPRLink group was significantly deeper in the age group of 55–65 years ( $47.1\pm9.6$  vs .  $38.5\pm8.7$  mm, p=0.001) than in the control group in the phase I test (table 3). However, the CC depth showed a tendency to be deeper in TCPRLink group but the difference was not statistically significant in the phase II test conducted 3 months later ( $44.7\pm10.1$  vs  $39.3\pm10.8$  mm, p=0.07; table 4).

### DISCUSSION

This study evaluated a novel, digital invention that integrated an audiovisual feedback smartphone application and a web-based system, thereby combining real-time dispatcher instructions and real-time feedback to ensure the appropriate quality of CPR. We compared the quality of T-CPR performed by potential bystander-rescuers in the age range of 18–65 years in a cardiac arrest simulation scenario with or without the smartphone application. The results of this study showed that real-time, audiovisual feedback using a smartphone application and webbased system in combination with dispatcher instructions

Table 2 Participants' CPR p	performance asse	ssment in the T-C	PR simulation	scenario		
	Phase I (n=186)			Phase II (n=170	))	
T-CPR performance	TCPRLink group (n=94)	T-CPR group (n=92)	P-value	TCPRLink group (n=86)	T-CPR group (n=84)	P-value
Counting with the dispatcher (n, %)	20 (21.3)	38 (41.3)	0.003	6 (7.0)	19 (22.6)	0.004
Time from call connected to	o: (seconds, mea	an±SD)				
Cardiac arrest identification	98.2±12.8	99.1±16.9	0.68	101.7±13.0	104.2±15.0	0.25
First chest compression	143.6±17.8	140.0±25.8	0.27	149.7±16.6	146.0±20.2	0.19
CPR parameters (M (P <sub>25</sub> -P <sub>75</sub>	<b>) or mean±SD</b> )					
Total number of compressions	661 (643–674)	648 (615–674)	0.035	661 (644–675)	646 (630–667)	0.002
Average compression rate $(min^{-1})$	111 (109–113)	108 (103–112)	0.002	111 (109–113)	108 (105–112)	<0.001
Percentage of adequate rate (100–120 min <sup>-1</sup> , %)	96 (89–98)	82 (50–97)	<0.001	95 (78–98)	93 (67–97)	0.11
Average compression depth (mm)	45.4±8.8	43.6±8.8	0.17	43.9±9.1	42.9±11.5	0.59
Percentage of adequate depth (50–60 mm, %)	20 (3–74)	12 (0–51)	0.14	17 (4–54)	13 (0–57)	0.26
Percentage of fully released (%)	97 (72–100)	97 (69–100)	0.79	95 (54–100)	96 (51–100)	0.40
Average hands-off time (s)	0 (0–1)	0 (0–1)	0.24	0 (0–1)	0 (0–1)	0.72

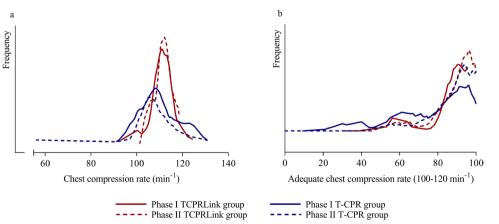
Phase I tests were cardiopulmonary resuscitation (CPR) performance and capabilities assessment using the telephone-assisted CPR (T-CPR) simulation scenario among individuals who have undergone CPR training with/without the TCPRLink application.

Phase II tests were CPR skill retention assessments among individuals with/without TCPRLink application after 3 months.

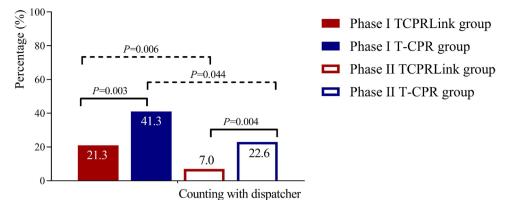
T-CPR, telephone-assisted cardiopulmonary resuscitation .

augmented the interaction between dispatchers and bystanders with a resultant positive effect on the quality of bystander-rendered CPR.

Dispatchers may coach callers to perform CPR, although they rely on audio communication alone to understand what is happening. With no other means of feedback, depending on the dispatcher's instructions may lead to lower quality CC and more hands-off time.<sup>21</sup> Several experimental manikin studies have demonstrated the potential benefits and drawbacks of video-assisted communication between rescuers and dispatchers compared with that of the conventional audio-instructed practice with regard



**Figure 3** Distribution of the chest compression rate and the proportion of the adequate chest compression rate (100–120 min<sup>-1</sup>) in TCPRLink group and T-CPR group. Phase I test was conducted in T-CPR trained individuals with/without TCPRLink App after inclusion. Phase II test was conducted in the same individuals with/without TCPRLink App after 3 months. T-CPR, telephone-assisted cardiopulmonary resuscitation.



**Figure 4** Counting with dispatcher in TCPRLink group and T-CPR group. Phase I test was conducted in T-CPR trained individuals with/without TCPRLink App after inclusion. Phase II test was conducted in the same individuals with/without TCPRLink App after 3 months. T-CPR, telephone-assisted cardiopulmonary resuscitation.

to the CC rate and hand position.<sup>22–26</sup> In a recent study that compared the real-world effects of video-instructed or audio-instructed T-CPR on the resuscitation outcomes, video-instructed T-CPR caused no delay in initiating CC although it was not associated with improvement in the survival rates.<sup>27</sup>

In dispatch-assisted instructions, the smartphone has secured a role as a promising carrier to improve video resuscitation care with its wide availability and high communication capabilities. Several diversified, advanced smartphone applications have been developed for integration into the links of the chain of survival and have feasibly created a strengthened 'Mobile chain of survival'28 as shown previously. One kind of application guides users in their CPR procedures via text and pictures or provides video examples of CPR with metronomic guidance that a bystander could watch before or during an actual resuscitation.<sup>68</sup> Another application provides measurement of CPR quality and feedback based on motion-sensing which require the user to place the phone on the patient's chest or hold it between the rescuer's hands while performing CPR.<sup>9-12</sup> However, these previous smartphone solutions have neglected the potential to leverage the dispatcher's involvement and, therefore, may be less suitable for reallife emergencies as the phone connection may be accidentally lost when using the phone as a CPR feedback device.

Given its salient differences with regard to the other smartphone applications, the TCPRLink application could improve the effectiveness of T-CPR, both on the dispatcher instruction and bystander operation aspects. The TCPRLink application uses the smartphone front facing camera for continuous quality improvement through real-time feedback for the bystander and the dispatcher. Dispatcher could monitor the hands-off time and encourage the bystander to continue CPR when they experience fatigue. Therefore, this application may be suitable for real-world emergencies when considering the prolonged time to call the dispatch centre and start CC, and that phone connection may be accidentally lost wmhen using the phone as a CPR feedback device.<sup>10</sup>

As the risk of OHCA increases with age,<sup>29 30</sup> older adults are more likely to be bystanders when their spouse or a family member experiences a cardiac arrest. The CPR capability of older adults has always been a significant concern. Another study that evaluated the effectiveness of a smartphone CPR application showed that participants aged over 60 years could not sustain longduration CPR.<sup>9</sup> However, in contrast with the results of that study, our study showed that TCPR Link app used with dispatcher assistance caused extra stimulus among seniors aged 55-65 as indicated by the subgroup analysis, with comparable quality of CPR with that of the younger participants during the 6min of hands-only CPR. Moreover, providing a feasible CPR feedback devices for seniors might be an appropriate approach to increase both their ability and also their willingness and confidence to do CPR.<sup>9</sup> When guided by the TCPRLink application, the CC rate and depth of CPR performed by older participants were both better and in adherence to the guidelines when compared with that in the conventional T-CPR group. These data suggest that, with the two-way metric of CPR quality and dispatcher encouragement, older participants performed CPR equally well as did the younger generation.

Counting aloud is the the most common method by which the dispatcher can ensure an appropriate CC rate in T-CPR. Without feedback from the rescuer, the dispatcher's understanding of the rescuer's situation is poor.<sup>31</sup> Interestingly, we found that visual guidance of the CC rate from the speedometer on the smartphone reduced the need to count the number of CC aloud to maintain an appropriate rate. Thus, rescuers could expend more energy on compression and less on counting. Furthermore, a lesser need for counting in the dispatcher's protocol leaves more time to coach for compression depth and avoiding leaning. Contrary to the common concern that the use of mobile devices or smartphone applications to improve CPR quality might cause a delay in the initiation of CCs,<sup>8 10</sup> the time to the first CC in the TCPRLink group was not prolonged as compared with that in the conventional T-CPR group in this study.

Table 3 Age-stratifie	Age-stratified comparison of the participants' CPR performance in the TCPRLink and T-CPR groups (phase I)	e participants' CPR	performan	ce in the TCPRLink	and T-CPR group:	s (phase I)			
	Age 18–24 years			Age 25–54 years			Age 55–65 years		
TCPR performance	TCPRLink group (n=29)	T-CPR group (n=25)	P value	TCPRLink group (n=37)	T-CPR group (n=38)	P value	TCPRLink group (n=28)	T-CPR group (n=29)	P value
Counting with the dispatcher (n, %)	9 (31.0)	12 (48.0)	0.20	7 (18.9)	19 (50.0)	0.005	4 (14.3)	7 (4.1)	0.35
Time from call connection to: (s, mean±SD)	ion to: (s, mean±SD)								
Cardiac arrest identification	98.9±13.5	96.4±19.5	0.58	97.0±13.2	98.2±14.0	0.72	98.9±12.0	102.7±17.8	0.36
First chest compression 141.4±20.2	141.4±20.2	137.8±26.8	0.57	143.6±17.7	135.9±26.9	0.14	145.8±15.4	147.1±22.5	0.79
CPR parameters (M ( $P_{25}-P_{75}$ ) or mean±SD)	5-P <sub>75</sub> ) or mean±SD)								
Total number of compression	663 (640–671)	650 (608–666)	0.21	659 (653–677)	652 (632–674)	0.29	659 (640–676)	640 (612–672)	0.14
Average compression rate (min <sup>-1</sup> )	111 (108–113)	108 (101–112)	0.03	111 (109–114)	109 (106–113)	0.12	110 (107–113)	107 (103–113)	0.06
Percentage of adequate rate (100–120 min <sup>-1</sup> , %)	95 (88–99)	82 (50–96)	0.01	97 (90–98)	89 (51–97)	0.006	95 (88–97)	71 (48–95)	0.003
Average compression depth (mm)	41.8±7.8	43.1±6.6	0.49	46.9±8.2	47.8±8.1	0.67	47.1±9.6	38.5±8.7	0.001
Percentage of adequate depth (50–60mm, %)	8 (0–28)	12 (4–33)	0.37	25 (9–84)	37 (7–86)	0.92	45 (1–99)	1 (0–14)	0.002
Percentage of fully released (%)	100 (95–100)	100 (96–100)	0.66	98 (79–100)	95 (57–99)	0.24	71 (5–100)	96 (37–100)	0.13
Average hands-off time (s)	0 (0–2)	0 (0–1)	0.24	0 (0–1)	(00) 0	0.45	0 (0–1)	0 (0–1)	0.92
Phase I tests were conducted for the evaluation of CPR performance and CPR training with/without the TCPRLink application.	sted for the evaluation of the TCPRLink applicati	of CPR performance ar ion.	nd capabilities	s assessment using a t	elephone-assisted CP	R (T-CPR) sin	capabilities assessment using a telephone-assisted CPR (T-CPR) simulation scenario among individuals who have undergone	g individuals who have	undergone

Table 4 Age-stratified com	nparison of the	participants' CPR	performanc	Age-stratified comparison of the participants' CPR performance between the TCPRLink and T-CPR groups (phase II)	PRLink and T-CPR (	groups (pha	se II)		
	Age 18-24 years	ars		Age 25–54 years			Age 55–65 years		
TCPR performance	TCPRLink group (n=29)	T-CPR group (n=23)	P value	TCPRLink group (n=31)	T-CPR group (n=34)	P value	TCPRLink group (n=26)	T-CPR group (n=27)	P value
Counting with the dispatcher 2 (6.9) (n, %)	2 (6.9)	4 (17.4)	0.40	4 (12.9)	12 (35.3)	0.036	(0) 0	3 (11.1)	0.24
Time from call connected to: (seconds, mean±SD)	(seconds, mea	n±SD)							
Cardiac arrest identification	99.8±16.0	100.4±16.3	0.89	103.8±13.6	103.0±11.9	0.82	101.4±7.3	108.9±16.7	0.04
First chest compression	148.4±20.1	144.2±25.5	0.52	153.2±17.3	143.6±13.9	0.018	146.8±9.8	150.4±21.9	0.45
CPR parameters (M ( $P_{25}-P_{75}$ ) or mean±SD)	or mean±SD)								
Total number of compressions	658 (643–678)	639 (605–653)	0.004	665 (653–675)	648 (640–668)	0.09	663 (644–676)	643 (627–680)	0.32
Average compression rate (min <sup>-1</sup> )	111 (109–113)	107 (101–110)	<0.001	112 (109–113)	109 (107–112)	0.06	111 (107–114)	109 (105–114)	0.28
Percentage of adequate rate (100–120 min <sup>-1</sup> , %)	96 (82–99)	82 (60–98)	0.08	95 (78–98)	95 (84–97)	0.64	92 (77–98)	90 (70–97)	0.61
Average compression depth (mm)	41.7±8.2	43.7±14.1	0.55	45.2±9.0	45.4±9.6	0.92	44.7±10.1	39.3±10.8	0.07
Percentage of adequate depth 10 (2-32) (50-60mm, %)	10 (2–32)	10 (0–54)	0.92	19 (7–55)	22 (2–81)	0.97	19 (3–68)	2 (0–24)	0.04
Percentage of fully released compressions (%)	100 (92–100)	99 (91–100)	0.52	93 (15–100)	90 (44–99)	0.65	68 (33–100)	89 (24–99)	0.84
Average hands-off time (s)	0 (0–2)	0 (0–1)	0.16	0 (0–1)	0 (0–1)	0.48	0-0) 0	0-0) 0	0.89
Phase II tests were conducted for the evaluation of CPR performance and capabilities assessment using telephone-assisted CPR (T-CPR) simulation scenario among individuals who have received CPR training with/without a TCPRLink application at 3 months after the training. T-CPR, telephone-assisted cardiopulmonary resuscitation .	r the evaluation of application at 3 opulmonary resur	of CPR performance a months after the traini scitation .	nd capabilities ng.	s assessment using tel	ephone-assisted CPR (	T-CPR) simul	ation scenario among	individuals who have	eceived CPR

Nevertheless, some limitations of this study need to be mentioned. On the one hand, this study was implemented in a simulated environment which may not reflect the real-world scenario. The Hawthorn effect could not be excluded under the simulation scenario, and could result in a motivation bias. Therefore, this study followed a realistic approach to the simulation of bystander CPR in a cardiac arrest scenario. We invited a senior dispatcher who worked in the emergency dispatch centre to portray the T-CPR scenario. On the other hand, a manikin may not represent the diversity of patients' chests and the changes in chest resistance during extended CPR. Finally, we recruited voluntary participants aged between 18 and 65 years who attended the 'WeCan CPR' training project. Therefore, the participants of this study might have had a selection bias as they had a positive willingness and knowledge of CPR training. We found that elderly individuals older than 65 years were less likely to participate, considering their physical capacity. The mean age of participants was nearly 40 years, which might not be the representative age for bystanders in real life.

## Conclusions

The TCPRLink smartphone application provides realtime feedback to both rescuer and dispatcher to enable more effective CC and lighten the load of counting out the CC with the dispatcher in a simulated T-CPR scenario. Further investigations are required to confirm the effectiveness of this application in the real-life resuscitation scenario.

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**Competing interests** All of the authors declare that they have no potential or actual conflicts of interest. HM and TSB are employees at Laerdal Medical. The manufacturer (Laerdal Medical, Norway) does not have any conflicts of interest with regard to the trial or in the interpretation of the results.

Patient consent for publication Not required.

**Ethics approval** The study protocol was approved by the Joint Research Ethics Board of the Shanghai Jiao Tong University Schools of Public Health and Nursing (approval ID SJUPN-201714).

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