Original Article

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Effectiveness of Mobile Health Application Use to Improve Health Behavior Changes: A Systematic Review of Randomized Controlled Trials

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Objectives: The purpose of this study was to examine the effectiveness of mobile health applications in changing health-related behaviors and clinical health outcomes. Methods: A systematic review was conducted in this study. We conducted a comprehensive bibliographic search of articles on health behavior changes related to the use of mobile health applications in peer-reviewed journals published between January 1, 2000 and May 31, 2017. We used databases including CHINAHL, Ovid-Medline, EMBASE, and PubMed. The risk of bias assessment of the retrieved articles was examined using the Scottish Intercollegiate Guidelines Network. Results: A total of 20 articles met the inclusion criteria. Sixteen among 20 studies reported that applications have a positive impact on the targeted health behaviors or clinical health outcomes. In addition, most of the studies, which examined the satisfaction of participants, showed health app users have a statistically significant higher satisfaction. Conclusions: Despite the high risk of bias, such as selection, performance, and detection, this systematic review found that the use of mobile health applications has a positive impact on health-related behaviors and clinical health outcomes. Application users were more satisfied with using mobile health applications to manage their health in comparison to users of conventional care.

Keywords: Health Behavior, Mobile Applications, Smartphone, Review, Mobile Health

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I. Introduction

The global mobile health (mHealth) application (app) market has been growing at a tremendous rate, and it is expected to continue to flourish [1]. These mHealth apps provide quick and easy access, transfer, and tracking of health information as well as interactive displays and interventions that can allow users to be highly engaged in promoting health outcomes and changing health-related behaviors [2]. Thus, health-related apps have a great potential to aid a wide range of target audiences with a variety of health issues [3].

Despite the evolution and widespread use of these mHealth apps, the factors involved in smartphone and health app use and their effectiveness are not yet fully understood and the field of research related to mHealth apps is still in a nascent

stage [4]. Kitsiou et al. [5] mentioned that a wide range of mHealth apps have not been strictly evaluated. For this reason, most consumers use mHealth apps without any concrete information about their effectiveness or harm and evidence of the effectiveness of mHealth has been inconclusive and not fully understood [4].

Thus, research determining the effectiveness of health apps is urgently needed [2]. This study aimed to demonstrate the effectiveness of mHealth apps in changing health-related behaviors and clinical health outcomes through a systematic review of randomized controlled trials (RCTs).

II. Methods

1. Search Strategy

We searched the electronic literature of RCTs published from January 1, 2000 to May 31, 2017, using four databases: the Cumulative Index to Nursing and Allied Health Literature (CINAHL), PubMed, Excerpta Medica dataBASE (EM-BASE), and Ovid Medline.

A university librarian was consulted who was a subject expert in the field of teaching and learning of systematic review. Searches used the following medical subject headings terms and keywords in various combination. We derived three broad themes that were then combined using the Boolean operator 'AND'. The first theme 'mobile' was created using the Boolean operator 'OR' to combine text words (mobile*, OR smartphone*). The second theme 'application' was created using the Boolean operator 'OR' to combine text words (app*, OR application*). The third theme 'health behaviors', was created using the Boolean operator 'OR' to combine text words (health behaviors*, OR health behaviors change*, OR behaviors change*).

In this study, to secure the quality of this systematic review, the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines were used [6]. This is a tool developed to verify the quality of evidence obtained during systematic reviews.

2. Study Selection

Two investigators independently reviewed the titles first and then examined the abstracts. Data extraction was conducted by one reviewer (MH Han) and was rechecked for by another reviewer to confirm the accuracy (EJ Lee). The same investigators read and screened the full texts to make the final decision. The reasons for inclusion and exclusion were recorded.

We included articles with the following characteristics:

(1) published in English, (2) published during the period from 2000 to 2017, (3) results related to changes in health behaviors, (4) RCTs designed for app-based interventions to improve any health-related behaviors. The exclusion criteria were being other kinds of study than RCTs; qualitative studies; books; conference proceedings; reviews; dissertations; protocols; or studies examining text messages, Web, emails, Twitter, social network services, or personal digital assistantbased health interventions. We also excluded studies lacking behavior change indicators or outcomes, not using apps as the primary intervention tools; or focusing primarily on app design and development. Conference abstracts, protocol papers, reviews, editorials, and commentary were also excluded.

References that clearly did not meet all criteria were excluded. Full-text articles that appeared to be relevant were retrieved and independently assessed by two reviewers. Disagreements were resolved through a meeting. The initial search revealed 1,247 articles: 66 in CINAHL, 481 in Ovid-Medline, 626 in EMBASE, and 74 in PubMed. Following the PRISMA guidelines (Figure 1), we removed duplicates and screened the titles and abstracts, which narrowed the list down to 57 relevant articles. Two investigators reviewed these 57 articles, 37 of which conducted other interventions with apps, compared two app-functions, examined protocols, or had unclear outcomes. Thus, finally, 20 articles were included for this review.

3. Data Collection and Analysis

From the 20 included articles, the following information was retrieved and analyzed: first author, year of publication, country, study design, themes, participants' character, sample size, mean age, intervention tool, follow-up duration, intervention characteristics, outcome measurements, as well as reported outcomes and significant levels. The search was broad with no limited target health-related behaviors in the search strategy. In this study, apps were considered effective if statistically significant results of health-related behavior changes were reported for them.

4. Risk of Bias Assessment

Risk of bias assessment for included studies was conducted by two authors using a modified version of the Scottish Intercollegiate Guideline Network (SIGN) checklist for RCT [7]. Specifically, the SIGN checklists were applied to grade the level of evidence of each study. The evaluation items are divided into 7 categories as follows: random sequence generation (selection bias), allocation concealment (selection

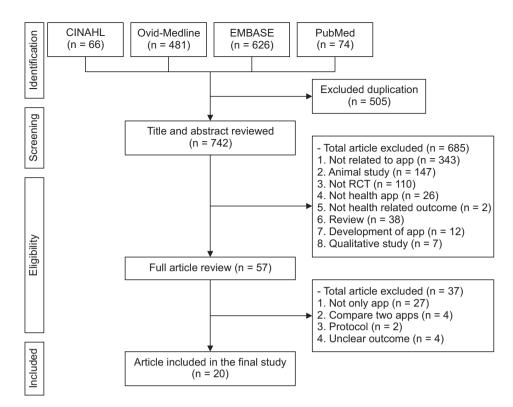


Figure 1. PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) flow diagram.

bias), blinding of participants and personnel (performance bias), blinding of outcome assessment (detection bias), incomplete outcome data (attrition bias), selective reporting (reporting bias), and other bias. Each domain was classified as having a low, high, or unclear risk of bias. This assessment was conducted by two researchers, and disagreements were resolved by discussion.

III. Results

1. Characteristics of Included Studies

The RCTs included in this review were published between 2014 and 2017. Most (n=16) were simply RCTs; the rest were an open-labelled RCT, an unmasked RCT, a cluster RCT, and a single-blinded parallel 3-arm pilot cluster RCT. The longest study duration was 8 months [8].

Five studies had a large number of participants [8-12]. The cluster RCT study had the largest sample size (n = 1,192) [11]. Two studies had a moderate number of participants [13,14]. The other 13 studies had a small number of participants. All 20 studies had between 80% and 100% retention rates; 18 (90%) studies achieved high (80–100%) retention rates in the intervention group, and only two (10%) studies [8,14] had a moderate retention rate.

The 20 selected studies were analyzed in this systematic review, and the following 16 themes related to health behaviors were created: physical activity (4), alcoholism (1), dietary

change (1) adherence of medication or therapy (2), preparation of clinical procedure (1), PTSD management (1), weight loss (2), prenatal education and engagement (1), adherence to follow-up clinic appointments (1), improvement of CPR skill performance (2), suicide prevention (1), prevention of CHD (1), smoking cessation (1), and knowledge improvement of pap testing (1). The comprehensive characteristics of included articles are summarized in Table 1.

2. Risk of Bias of Selected Studies

The evaluation of risk of bias for all 20 studies was conducted using the SIGN checklist for RCTs. The results were summarized using the risk of bias table of RevMan 5.3 software. A total of 15 studies properly reported random sequence generation. Only one article did not mention random sequence generation [15].

For allocation concealment, only 6 studies explicitly mentioned that allocation was concealed [8,10,16-18]. However, 8 studies did not discuss allocation concealment adequately. Participants were blinded in 4 studies [16,17,19,20]. However, due to the traits of mHealth apps, some studies could not be conducted with perfect blinding. For the remaining 16 studies, either they were not blinded or information on blinding was not clearly provided in the reporting.

Among the studies in this review, three reported the blinding of outcome assessment [16,17,19]. For reporting bias, 9 studies had a low risk of bias, and 11 were evaluated as un-

Table 1. Summary characteristics of articles included

			Study					Follow-		Outcome measure-	
Study	Year	Study Year Country	design	Themes	Participants	Group (N, Age)	Intervention	dn	Features	ment	Outcomes
Glynn	2014	West of	2014 West of Open-la-	Physical Patient		-Intervention	Smart-	2 mo	-Automatic feedback -Mean daily step	-Mean daily step	-Mean of daily step count
et al.		Ireland	Ireland bel RCT	activity	activity referred by	group $(n = 37, 46)$ phone app	phone app		Tracking of step	count	-Intervention group
[19]					their pri-	± 11 years)	(SMART		count and calories	-Improvement of	(n = 4,365)
					mary care	-Control group	MOVE)		burnt	daily step count	-Control group
					health pro-	$(n = 40, 42 \pm 11)$			-Graphic display of	in the interven-	(n = 5,138)
					fessional or	years)			step-count history	tion group	-Mean improvement in the
					self-referred				-Goal setting		intervention group (n =
											1,029; 95% CI, 213-1843;
											p = 0.009
Gus-	2014 USA		Unmasked Alcohol- Those who	Alcohol-		-Intervention	Smartphone 8 mo		-Education (audio-	-Count the fewer	-Fewer risk of drinking
tafson			RCT	ism	met the	group $(n = 170,$	app (A-		guided relaxation)	risk of drinking	day: intervention group
et al.					criteria	$38.3 \pm 9.5 \text{ years}$	CHESS)		-Interactive features	days	(1.39),
[8]					for DSM-	-Control group			(counselor discuss		control group (2.75)
					IV alcohol	$(n = 179, 38.4 \pm$			with users)		(Mean difference = 1.37 ;
					dependence	11.2 years)					95% CI, $0.46-2.27$; $p =$
											0.003)
Ipjian	2017 USA		RCT	Dietary Healthy		-Intervention	Smartphone 1 mo		-Received feedback	-24-hour sodium	-24-hour sodium excretion:
and				change	adults	group $(n = 15,$	app (My-		on sodium content	excretion	intervention group (-838
John-						35.5 ± 14.9	FitnessPal)		of foods	-Satisfaction	\pm 1,093 mg/day), control
ston						years)			-Entering the data		group $(236 \pm 1,333 \text{ mg/})$
[25]						-Control group:			(daily food and bev-		day) $(p = 0.010)$
						Journal group			erage entry to moni-		-Intervention group report-
						$(n = 15, 33.3 \pm$			tor dietary sodium		ed significantly greater
						16.8 years)			levels)		satisfaction than the
									-Instruction and edu-		journal group $(p = 0.010)$
									cational materials		

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Year Country	_ ≥	Study	Themes	Participants	Group (N, Age)	Follow- Intervention	Features	Outcome measure-	Outcomes
2	2015 USA	RCT	Adher- ence to medica- tion	College students who had a current prescription for an antidepressant and regularly used a smartphone device	-Intervention group (n = 30, 20.3 ± 4.0 years) -Control group (n = 27, 20.9 ± 4.7 years)	Smartphone 1 mo app	-Reminder (par- ticipant entered the prescribed infor- mation regarding dosing; participants were asked to use the medication reminder app to indicate when they had taken their medication by responding to the message received)	Percent adherence -	Percent adherence -Participants in the treatment group were 3.5 times more likely to adhere to their medication regimen than those in the control group (95% CI, 0.945–12.966; X^2 [1, $n = 40$] = 3.64; $p = 0.057$; $\phi = 0.30$).
0	Kang et 2016 China al. [12]	RCT	Preparation before procedure (bowel preparation)	Colonoscopy -Intervention outpatients group (n = 3 44.4 ± 13.2 years) -Control grou (education c standard for (n = 383, 45 13.0 years)	Intervention group (n = 387, 44.4 ± 13.2 years) -Control group (education of standard form) (n = 383, 45.5 ± 13.0 years)	Smartphone 6 mo app (We- Chat)	-Social media instrucLevel of adequate tion bowel preparation (Ottawa score<6).		A higher proportion of patients in the intervention group had adequate bowel preparation than in the control group (82.2% vs. 69.5%, $p < 0.001$). A higher proportion of patients in the intervention group had cecal intubation and have adenomas than in the control group (97.2% vs. 93.2%, $p = 0.014$; 18.6% vs. 12.0%, $p = 0.012$).

Table 1. Continued 2

Study	Study Year Country	Sountry	Study	Themes	Participants	Group (N, Age)	Follow- Intervention	Features	Outcome measure-	Outcomes
Kuhn et al. [13]	2017 USA		RCT S	Self -man- PTSD agement of PTSD symp- toms		Intervention group (n = 62, 39.43 \pm 15.16 years) -Control group: Waitlist condition (n = 58, 39.12 \pm 14.08 years)	Smartphone 3 mo app (PTSD coach)	-Instruction/Educa- tion: Offer sound psychoeducational information and evidence-based cognitive behavior tools	symp-	-Intervention group par- ticipants had significant- ly greater improvement in PTSD symptoms (<i>p</i> = 0.035), depression symptoms (<i>p</i> = 0.005), and psychosocial func- tioning (<i>p</i> = 0.007) than waitlist participants. -A greater proportion of PTSD coach partici- pants achieved clini- cally significant PTSD symptom improvement (<i>p</i> = 0.018) than waitlist participants.
Laing et al. [10]	2014 USA		RCT	weight	Overweight primary care patitients	-Participants (n = 212, 43.3 ± 14.3 years): Intervention group (n = 107) and control group (usual care; n = 105)	Smartphone 6 mo app (My- FitnessPal)	-Self-monitoring -Goal setting -Feedback	-Weight loss at 6 months -SBP -Satisfaction with the app	-Weight change: No difference between groups (-0.30 kg [95% CI, -1.50-0.95; p = 0.63]) -Systolic pressure: No difference between groups (-1.7 mmHg; 95% CI, -7.1-3.8; p = 0.55) -Most users reported satisfaction.

Table 1. Continued 3

group: rmation ancy more p = 0.04), de- ater patient p = 0.02 group. e was de- erpersonal munication.	VAS & NDI: In the intervention group, VAS (p = 0.003) and NDI (p = 0.005) improved significantly after the appbased neck exercise. MVES (p = 0.013), improved significantly in the intervention group, the level of exercise adherence was high.
Intervention Record inform about pregn frequently (veloped greativation (than control -No differenc tected in int	-VAS & NDI: In the intervention group, VAS (p = 0.003) and NDI (p = 0.005) improved significantly after the appbased neck exercise. -MVES (p = 0.013), improved significantly -In the intervention group, the level of exercise adherence was high.
-The frequency of recording information	-VAS for pain intensity -NDI for functional disability -MVFS/MVES -Level of exercise adherence
Entering the data (record information: pregnancy experi- ence—weight, blood pressure, experience between prenatal appointments)	-Education of neck exercise
Smartphone 5 mo app	e 2 mo
-Intervention group (n = 65, 43.3 ± 14.3 years) -Control group: Notebook (n = 62, 29.29 ± 4.80 years)	-Participants (43.3 Smartphon ± 14.3 years): In- app (My- tervention group FitnessPa (n = 11) and control group (Brochure; n = 9)
Pregnant mothers	Office workers
Prenatal educa- tion and engage- ment	Neck exercise (pain manage)
RCT	RCT
. USA	2017 USA
al. [4]	al. [1]
	Prenatal Pregnant -Intervention Smartphone 5 mo -Entering the data - Geduca mothers group (n = 65 , app (record information: tion and 43.3 \pm 14.3 pregnancy experienged ment years) engage - Control group: hond pressure, experience Notebook (n = 62 , 29.29 ± 4.80 preparable appointments) years)

Table 1. Continued 4

Study	Study Year Country	Study y design	Themes	Participants	Group (N, Age) Intervention		Follow- up	Features	Outcome measure- ment	Outcomes
yu et	2016 China	RCT	Clinical	Patients diagIntervention		Smartphone	om 9	-Entering the data	-Time consump-	-Time consumption for
al.			follow-	nosed with	group $(n = 53,$	app (We-			tion for follow-	follow-up delivery:
[18]			up with	head and	61.0 ± 13.0	chat)			up delivery -Total	WFU group (23.36 ±
			dis-	neck tumor	years)				economic cost	6.16 minutes) was sig-
			charged	,	-Control group:				-Lost to follow-up	-Lost to follow-up nificantly shorter than
			patients		Telephone (TFU)				rate -Satisfaction	that in the TFU group
					$(n = 46, 61.5 \pm$				with methods	$(42.89 \pm 7.15 \text{ minutes})$
					9.3 years)					(p < 0.001).
									•	-Total economic cost in
										WFU group (RMB 90
										Yuan) was much lower
										than in the TFU group,
										9.80% (5/51).
									•	-Lost follow-up rate:
										Intervention group
										(7.02%, 4/57) and TFU
										group (9.80%, 5/51)
									•	-Satisfaction: Intervention
										group (94.34%, 50/53)
										and TFU group (80.43%,
										37/46) (95% CI, 0.057-
										0.067; p = 0.034)

Table 1. Continued 5

Study	Study Year Country	Study design	Themes	Participants	Group (N, Age)	Intervention	Follow- up	Features	Outcome measure- ment	Outcomes
lord	2016 Sweden Cluster	Cluster	CPR	Students	-Intervention	Smartphone 1 ses-	1 ses-	-Training/Education	-CPR skills	-The DVD-based group
et al.		RCT		(13 years)	group $(n = 549,$	app	sion		-Willingness to act	was superior to the app-
[11]					13 years)		(30		-Improvement	based group in CPR
					-Control group:		min-		of compression	skills: a total score of 36
					DVD-based CPR		utes)		depth	(33–38) vs. 33 (30–36)
					training (n =					directly after training (p
					575, 13 years)					< 0.001) and 33 (30-36)
										and 31 (28-34) at 6
										months ($p < 0.001$).
										-If a friend suffered car-
										diac arrest, 78% (DVD)
										vs. 75% (app) would do
										compression and ven-
										tilations, whereas only
										31% (DVD) vs. 32%
										(app) would perform
										standard CPR if the
										victim was a stranger.
										-At 6 months, the DVD
										group performed sig-
										nificantly better in 8 out
										of 12 CPR skill com-
										ponents. Both groups
										improved compression
										depth from baseline to
										follow-up.

Table 1. Continued 6

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Outcomes	-Proper compression	depth (mm) was shown	in the traditional group	(53.77) compared to	the smartphone group	(48.35) (p < 0.01).	-The proper chest	compression (%) was	formed suitably ($p <$	0.05) in the traditional	group (73.96%) more	than smartphone group	(60.51%).	-The traditional group	(3.83 points) had higher	awareness of chest	compression accuracy	(p < 0.001) than the	smartphone group (2.32	points).
Outcome measure- ment	-The proper com-	pression depth	-The proper chest	compression	-Awareness of	chest compres-	sion													
Features	-Education (CPR curThe proper com-	riculum)																		
Follow- up	l ses-	sion																		
	Smartphone 1 ses-	app																		
Group (N, Age) Intervention	-Intervention	group $(n = 33,$	61.0 ± 13.0	years)	-Control group:	Traditional	group $(n = 31,$	$61.5 \pm 9.3 \text{ years}$												
Participants	Those who	agreed to	complete	CPR cur-	riculum															
Themes	CPR																			
Study design	RCT																			
ountry																				
Study Year Country	2014 Korea																			
Study	Park	[27]																		

Table 1. Continued 7

5, 5, 0 ars) p: 0	vention typ (n = ± 8.7.) trol gro al care s.3 ± 1.5 s)	activity stroke group (n = 15, 56.3 ± 8.7 years) Control group: Usual care (n = 8, 55.3 ± 12.6 years)
dd d	vention up (n = 17) rrol group: dard group 11)	People un- dergoing group (n = 17) ART -Control group: Standard group (n = 11)

Table 1. Continued 8

Outcomes	-Depressive SympDSI-SS were significant tom Inventory- in the ibobbly arm (t Suicidality = 2.40; df = 58.1; p = Subscale (DSI-SS) 0.0195); these differabatient Health ences were not signifiared. Questionnaire cant compared with the (PHQ-9) waitlist arm (t = 1.05; df -Kessler Psycho- = 57.8; p = 0.2962)Parlogical Distress ticipants in the ibobbly Scale (K-10) group showed substan-Barratt Impulsive tial and statistically significant reductions in PHQ-9 and K10 scores compared with waitlistNo differences were observed in impulsivity between intervention and control groups.
Outcome measure- ment	-Depressive Symp1 tom Inventory- Suicidality Subscale (DSI-SS) -Patient Health Questionnaire (PHQ-9) -Kessler Psycho- logical Distress Scale (K-10) -Barratt Impulsive Scale (BIS-11)
Features	-Education (delivered acceptancebased therapy)
Follow- up	5 wk
Intervention	Smartphone 6 wk app (ibob-bly)
Group (N, Age) Intervention	Intervention group (n = 31, 27.48 \pm 9.54 years) -Control group: Waitlisted (n = 31, 24.97 \pm 6.28 years)
Themes Participants	Suicide Indigenous preven- Australian tion youth
Themes	Suicide prevention
Study design	\(\text{CT} \)
Study Year Country	2017 Austra- RCT
Year	
Study	Tighe et al. [20]

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Table 1.	lable 1. Continued 9									
Study	Study Year Country	Study design	Themes	Participants	Group (N, Age)	Intervention	Follow- up	Features	Outcome measure- ment	Outcomes
Van Reijen et al. [9]	2017 New Zea- land	RCT	Neuro- mus- cular training	Athletes	-Intervention group (n = 110, 37.6 ± 13.1 years) -Control group: Booklet (n = 110, 38.1 ± 13.7 years)	app		-Training (exercise program)	-Compliance with the exercise program -Incidence density of self-reported recurrent ankle sprains	-The mean compliance to the exercise scheme was 73.3% (95% CI, 67.7–78.1) in the App group compared with 76.7% (95% CI, 71.9–82.3) in the Booklet group. No significant difference in compliance was found between groups. -The incidence densities of self-reported timeloss recurrences were not significantly different between groups (HR = 3.07; 95% CI, 0.62–15.20).
Zhang et al. [29]	2017 Singa- pore	RCT	CHD pre- Working vention people		Intervention group (n = 40) -Control group: Health promotion website only (n = 40)	Smartphone 6 weeks app (Care-4Heart)		-Education (heart disease prevention [SBCHDP] pro- gram)	-Awareness of CHD -Knowledge of CHD -Blood cholesterol control	-Participants in the intervention group had a better awareness of CHD ($X^2 = 6.486$, $p = 0.039$). -A better overall CHD knowledge level ($t = 3.171$, $p = 0.002$) -Better behavior concerning blood cholesterol control ($X^2 = 4.54$, $p = 0.033$) than participants in the control group.

Table 1. Continued 10

Study	Year	Year Country	Study design	Themes	Participants	Group (N, Age)	Fo Intervention	Follow- up	Features	Outcome measure- ment	Outcomes
arter	2013 New		RCT	Loss	Overweight	-Intervention	Smartphone 6 mo	υo	-Goal setting	-Self-monitoring	-Self-monitoring declined
et al.		Zea-		weight	volunteer	group $(n = 43,$	app (My		-Self-monitoring of -Mean weight	-Mean weight	over time in all groups.
[16]		land				$41.2 \pm 8.5 \text{ years}$	Meal Mate)		diet and activity	change	-Mean weight change at
						-Control group:			-Feedback via weekBMI change	-BMI change	6 months was -4.6 kg
						Website group (n			ly text message	-Change in body	(95% CI, -6.2 to -3.0)
						$= 40, 42.5 \pm 8.3$				fat	in the smartphone app
						years) and Paper					group, -2.9 kg (95%
						diary group (n					CI, -4.7 to -1.1) in the
						$= 42, 41.9 \pm 10.6$					diary group, and -1.3 kg
						years)					(95% CI, -2.7 to 0.1) in
											the website group.
											-BMI change at 6 months
											was -1.6 kg/m^2 (95%
											CI, -2.2 to -1.1) in the
											smartphone group, -1.0
											kg/m^2 (95% CI, -1.6 to
											-0.4) in the diary group,
											and -0.5 kg/m^2 (95%
											CI, -0.9 to 0.0) in the
											website group.
											-Change in body fat was
											-1.3% (95% CI, -1.7 to
											-0.8) in the smartphone
											group, -0.9% (95% CI,
											-1.5 to -0.4) in the diary
											group, and -0.5% (95%
											CI, -0.9 to 0.0) in the
											website group.

Table 1. Continued 11

study	Year	Study Year Country	Study design	Themes	Themes Participants	Group (N, Age) Intervention	Follow- Intervention up	Features	Outcome measure- ment	Outcomes
Cheung 2015 Hong et al. Kon; [17]	2015 I	Kong	Single blinded, parallel, 3-arm pilot cluster RCT	Smoking cessa- tion	Quitters who had stopped smoking recently	.4, p: nline d: d p: (n 10.4	Smartphone 2 mo app (Whats App)	-Group discussion -Training for smoking cessation	-2- and 6-month relapse rates (the proportion of participants who smoked at least 5 cigarettes in 3 consecutive days)	-The WhatsApp group (17%, 7/42) reported more relapse than the control group (42.6%, 23/54) at 2-month (OR = 0.27; 95% CI, 0.10–0.71) and 6-month (40.5%, 17/42 vs. 61.1%, 33/54; OR = 0.43; 95% CI, 0.19–0.99) followups.
Chris- 2014 New tensen Eng [30] land	2014 1	New Eng- land	RCT	Enhance knowl-edge of pap testing	Women attending a univer- sity in New England	-Intervention group (n = 37, 20.41 years) -Control group: Standard pamphlet on paptesting (n = 37, 20.81 years)	Smartphone 2 mo app (Digi- tal health education application on Pap testing)	-Education	-Knowledge of pap testing	-Knowledge of pap -Pap testing knowledge: testing Intervention group (5.26 \pm 1.66 to 10.78 \pm 1.51), control group (5.70 \pm 1.61 to 8.92 \pm 1.88) (p < 0.001); Knowledge scores on the posttest increased significantly in both groups but was significantly higher in the intervention group.

RCT: randomized controlled trial, DSM-IV: Diagnostic and Statistical Manual of Mental Disorders 4th edition, PTST: posttraumatic stress disorder, SBP: systolic blood pressure, VAS: Visual Analogue Scale, NDI: Neck Disability Index, MVFS: maximal voluntary flexion strength, MVES: maximal voluntary extension strength, CPR: cardiopulmonary resuscitation, ART: antiretroviral therapy, HIV: human immunodeficiency virus, CHD: Coronary heart disease, BMI: body mass index. clear on the presence of bias. A summary of Cochrane's risk of bias table is presented in Figure 2.

3. Content Characteristics of Apps

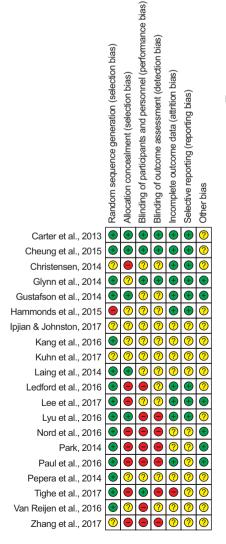
Some app characteristics of contents were categorized according to the behavior change technique taxonomy by Abraham and Michie [21] that is, providing information, planning (goal setting), reminding, providing feedback, or monitoring. Furthermore, additional app characteristics, such as entering data, education/training, and communication were derived from this study. Ten apps have multiple functions to manage health-related behaviors. The most common function of mHealth apps is providing the opportunity for education or training.

All participants in the control groups underwent standardized or usual care. Table 2 shows a summary of app characteristics.

IV. Discussion

These days, mHealth apps seem to be ubiquitous, and the body of research indicating their effectiveness has been growing rapidly. However, evidence for the effectiveness of mHealth apps has been uncertain, and much remains unknown in terms of health-related behavior changes and clinical results.

In this study, 20 RCTs were included to evaluate the effectiveness of mHealth apps for health-related behavior change. Seventeen studies among 20 showed a positive contribution to the enhancement of health-related behaviors. This result is similar to other previous evidence reviews [3]. Therefore, using mHealth apps could be an effective strategy to improve outcomes of users along with the high popularity of smartphone use in the everyday lives of users. However, three studies did not show positive effects of mHealth apps on health-related behavior changes. Laing et al. [10] found that



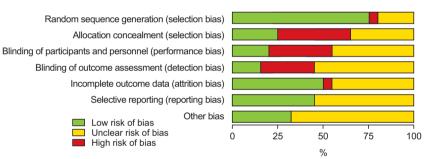


Figure 2. Summary of risk of bias.



Table 2. Summary characteristics of application contents

Ctudu	Providing	Planning	Remind	Feedback	Entering the	Education/	Communica-
Study	information	(goal setting)	Kemina	reedoack	data	Training	tion
Glynn et al. [19]		$\sqrt{}$			$\sqrt{}$		
Gustafson et al. [8]						$\sqrt{}$	$\sqrt{}$
Ipjian and Johnston [25]	$\sqrt{}$			$\sqrt{}$	$\sqrt{}$	$\sqrt{}$	
Hammonds et al. [15]			$\sqrt{}$		\checkmark		
Kang et al. [12]						$\sqrt{}$	
Kuhn et al. [13]	$\sqrt{}$					$\sqrt{}$	
Laing et al. [10]		\checkmark		$\sqrt{}$	$\sqrt{}$		
Leoford et al. [14]					\checkmark		
Lee et al. [1]						$\sqrt{}$	
Lyu et al. [18]					\checkmark		$\sqrt{}$
Nord et al. [11]						$\sqrt{}$	
Park [27]						$\sqrt{}$	
Paul et al. [26]		$\sqrt{}$		$\sqrt{}$			
Perera et al. [28]			$\sqrt{}$				
Tighe et al. [20]						$\sqrt{}$	
Van Reijen et al. [9]						$\sqrt{}$	
Zhang et al. [29]						$\sqrt{}$	
Carter et al. [16]		$\sqrt{}$		$\sqrt{}$	$\sqrt{}$		
Cheung et al. [17]						$\sqrt{}$	$\sqrt{}$
Christensen [30]						$\sqrt{}$	

there was no difference in weight loss between intervention and control groups. In addition, Nord et al. [11] reported that a DVD-based CPR education group had better performance than an app group. Tighe et al. [20] also reported that using an app had no effect on decreasing depression symptoms and impulsivity behaviors. However, it is difficult to assess the effectiveness of apps based on the results of a single study, and more studies with controlled research design are needed.

This study differs from other reviews in that we only included RCTs to examine the effectiveness of mHealth apps, because RCTs are considered the 'gold standard' in evaluating the effects of intervention and provide a valuable source of evidence in research and treated as a powerful experimental tool to examine the effectiveness of intervention [22]. Therefore, this systematic review, which only analyzed the results of RCTs, has provided more reliable evidence for the effectiveness of smartphone health apps.

More than half of the reviewed studies had small samples (<60). In addition, 11 reported that the duration of intervention was less than 2 months. According to Man-Son-Hing et al. [23], trials with larger samples and longer intervention

durations or follow-up times are more reliable to appraise the effectiveness of intervention. Based on the results of this review, to demonstrate a certain effect using mHealth apps for health-related behavior changes, more research with long intervention durations and large samples is needed.

According to Zhao et al. [3], the retention rate is defined as the proportion of participants who remain to complete a study. The Cochrane Handbook for Systematic Reviews of Interventions reported that studies with retention rates over 80% are classified as having low attrition, and studies with retention between 60% and 79% are classified as having moderate attrition [24]. Eighteen studies achieved high (>80%) retention rates in the intervention group. In this study, over half of the studies had a moderate-high retention rate. It can be assumed that the reasons for the high retention rate were the high feasibility and acceptability of app use in users' everyday lives. Thus, mHealth apps could be effectively adopted for users to improve health-related outcomes by managing and supporting health-related behaviors of users.

In this review, some studies [8,10,13,15-19,25,26] considered multiple function apps, such as entering data and

providing feedback, education, and reminders. In contrast, other studies [1,9,11,12,14,20,27-30] considered apps that have only one function; most of these apps had an education function. One study reported that having multiple app functions is much better to manage health status and to improve health-related behaviors [4]. However, this result might not be concrete because applying apps to the different types of situations, health behaviors, and participants can yield different results and effectiveness. For example, Laing et al. [10] conducted a study regarding an app with multiple functions for weight loss, and the effect was not significant. However, Zhang et al. [29] reported a significant result in improvement of coronary heart disease (CHD) knowledge and awareness using a single-function app. This suggests that many unconditional features of apps might not work properly, and a customized app is needed that fits the purpose and intent of the user. In this aspect, mHealth apps could provide individualized information via feedback to users and benefit them.

With respect to the risk of bias of included studies, the categories of selection bias (allocation concealment), performance bias (blinding of participants and researchers), and detection bias (blinding of outcome assessment) indicated high risk of bias. Therefore, to enhance the quality of studies and ensure low risk of bias, researchers should consider rigorous study design and reporting. Based on the results of this review, further studies using meta-analysis are needed to identify the effects of mHealth apps with specific outcomes. In addition, the effectiveness of the apps should be verified with the effects or risks being used with verification.

In this review, there were several limitations. First, we used broad key words, such as 'health', 'behaviors', 'smartphone', and 'mobile'. For this reason, many articles related to the use of mHealth apps on specific diseases or health conditions for instance, diabetes, hypertension, or asthma-might not have been included in this study. Second, in this review, only RCTs were included to analyze the effectiveness of mHealth apps. However, some of the RCTs did not fully follow the form of the RCT or applied a modified form of RCT. However, well-structured RCT is needed to verify mHealth app effectiveness. Third, almost all of the studies considered in this review were conducted in developed countries. Hence, it is difficult to generalize our results to developing countries. Fourth, we did not conduct a meta-analysis because of interventions with different kinds of mHealth apps. However, with the results of meta-analysis, the effectiveness of mHealth apps can be verified more clearly.

This systematic review was conducted to examine the ef-

fectiveness of mHealth apps to lead to changes in their targeted health-related behavior. This study summarized the characteristics and changes in targeted health outcomes. To our knowledge, there has been no previous systematic review of RCTs for identifying the effectiveness of mHealth apps in improving health-related behaviors. Similar to previous studies, this systematic review also found that the use of mHealth apps has a positive impact on health-related behaviors, such as physical activity, diet change, adherence to medication or therapy, and knowledge enhancement related to clinical procedures. Moreover, most apps seem to promote better clinical health outcomes. Most app users are satisfied with the use of mHealth apps to manage their health in comparison to users of conventional care. Although most studies analyzed indicated statistically significant effects to improve health, more RCTs with larger samples and longer applied interventions are still needed to confirm the effectiveness of mHealth apps. To assess the efficacy of mHealth apps in greater detail, further research is needed.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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