The efficacy of smartphone-based mental health interventions for depressive symptoms: a meta-analysis of randomized controlled trials

Joseph Firth[1,2], John Torous[3,4], Jennifer Nicholas[5,6], Rebekah Carney[1], Abhishek Pratap[7,8], Simon Rosenbaum[5,6], Jerome Sarris[1,9]

[1]NICM, School of Science and Health, Western Sydney University, Campbelltown, Australia; [2]Faculty of Biology, Medicine and Health, Division of Psychology and Mental Health, University of Manchester, Manchester, UK; [3]Department of Psychiatry and Division of Clinical Informatics, Beth Israel Deaconess Medical Center, Boston, MA, USA; [4]Harvard Medical School, Boston, MA, USA; [5]Black Dog Institute, University of New South Wales, Sydney, Australia; [6]Faculty of Medicine, School of Psychiatry, University of New South Wales, Sydney, Australia; [7]Sage Bionetworks, Seattle, WA, USA; [8]Department of Biomedical Informatics and Medical Education, University of Washington, Seattle, WA, USA; [9]Department of Psychiatry, University of Melbourne, Professorial Unit, The Melbourne Clinic, Melbourne, Australia

The rapid advances and adoption of smartphone technology presents a novel opportunity for delivering mental health interventions on a population scale. Despite multi-sector investment along with wide-scale advertising and availability to the general population, the evidence supporting the use of smartphone apps in the treatment of depression has not been empirically evaluated. Thus, we conducted the first meta-analysis of smartphone apps for depressive symptoms. An electronic database search in May 2017 identified 18 eligible randomized controlled trials of 22 smartphone apps, with outcome data from 3,414 participants. Depressive symptoms were reduced significantly more from smartphone apps than control conditions (g=0.38, 95% CI: 0.24-0.52, p<0.001), with no evidence of publication bias. Smartphone interventions had a moderate positive effect in comparison to inactive controls (g=0.56, 95% CI: 0.38-0.74), but only a small effect in comparison to active control conditions (g=0.22, 95% CI: 0.10-0.33). Effects from smartphone-only interventions were greater than from interventions which incorporated other human/computerized aspects along the smartphone component, although the difference was not statistically significant. The studies of cognitive training apps had a significantly smaller effect size on depression outcomes (p=0.004) than those of apps focusing on mental health. The use of mood monitoring softwares, or interventions based on cognitive behavioral therapy, or apps incorporating aspects of mindfulness training, did not affect significantly study effect sizes. Overall, these results indicate that smartphone devices are a promising self-management tool for depression. Future research should aim to distil which aspects of these technologies produce beneficial effects, and for which populations.

Key words: Smartphone technology, mental health interventions, depression, e-health, mhealth, apps, cognitive training, mood monitoring, cognitive behavioral therapy, mindfulness training

Depression is now recognized as a leading cause of global disability, impacting over 300 million people around the world[1]. In countries like the US, 9% of the population may have depression at any one time[2]. Beyond the personal suffering, depression is associated with unemployment, poor physical health, impaired social functioning and, in its most severe forms, suicide[3]. Thus, the disorder carries a high cost for both the individual and the society, particularly when considering the economic burden incurred through clinical care and lost productivity[4].

Depression is a potentially treatable condition, with a range of available medications and psychological interventions that are supported by robust clinical evidence. While the choice of pharmacotherapy or psychotherapy depends on many factors, for most individuals with mild or moderate depression they may be nearly equivalent[5].

However, there are many barriers towards both of these treatment methods. For instance, access to mental health care remains limited, as almost half of the world's population lives in countries where there is less than one psychiatrist per 100,000 people[6], and continued shortage in mental health care staff is expected for both the near and long term future[7,8]. Additionally, medications and psychotherapies may carry some level of stigma (particularly among younger people), which further limits their effectiveness[9,10].

Furthermore, although these therapies demonstrate high clinical efficacy for reducing symptoms, they may not always bring about full and sustained remission in those treated. Finally, many people experience either subclinical depression or residual depressive symptoms even after achieving clinical response to treatment. Therefore, novel primary and/or adjunctive methods for reducing depression on a population scale are urgently needed.

Digital technologies may represent a novel and viable solution. Mobile phones are among the most rapidly adopted innovations in recent history, and smartphone ownership continues to increase in both developed and developing countries[11]. Through providing ubiquitous Internet connectivity, along with the capacity to download and run externally created applications ("apps"), smartphone technology presents an opportunity to transform mobile phones into devices which could provide global, cost-effective and evidence-based mental health services on demand and in real time[12].

This clear therapeutic potential has triggered a wave of interest and investment in mental health apps from governments, technology companies, advocacy groups, and research groups internationally[13,14]. But in the enthusiasm to realize the potential of apps for depression, it has become difficult to separate actual efficacy from overzealous aspirational claims[15]. With thousands of mental health apps readily available through Apple or Google marketplaces, finding a useful tool supported by robust evidence to manage one's depression is clearly a challenge for a lay person[16,17]. The increasing media promotion and accessibility of apps for mental health now presents a "duty of

care" issue towards ensuring that people have information and understanding of evidence-based digital treatments for depression.

Recent meta-analyses have documented that various smartphone interventions can have positive effects on physical diseases, such as diabetes[18], and mental health conditions, such as anxiety[19]. However, the clinical effect of smartphone interventions on symptoms of depression has yet to be established. Thus, our aim was to examine the efficacy of delivering mental health interventions via smartphones for reducing depressive symptoms in both clinical and non-clinical populations. We also sought to use subgroup and meta-regression analyses in order to explore which aspects of smartphone interventions are associated with greater or lesser efficacy for depressive symptoms. The results of these meta-analyses provide the first overall estimate of effects from such interventions, along with informing treatment choices and future research in this area.

METHODS

This systematic review and meta-analysis followed the PRISMA statement for transparent and comprehensive reporting of methodology and results[20]. In order to eliminate researcher bias, the search strategy, inclusion criteria and data extraction, as well as the overall and pre-planned subgroup analyses, strictly adhered to those adopted in a previous systematic review of smartphone interventions for anxiety[19], as specified in a registered online protocol (CRD42017064882).

Search strategy

We conducted an electronic search of the following databases: Cochrane Central Register of Controlled Trials, Health Technology Assessment Database, Allied and Complementary Medicine (AMED), Health Management Information Consortium (HMIC), Ovid MEDLINE, Embase, and PsycINFO, from inception to May 1, 2017. The search applied the PICO framework[21], using a range of relevant terms to capture all potentially eligible results relating to smartphone mental health interventions for depressive symptoms. An additional search of Google Scholar was implemented, and reference lists of retrieved articles were checked to identify any further eligible studies.

Eligibility criteria

Only English-language articles were included. Eligible studies were all randomized controlled trials (RCTs) examining the effects of mental health interventions delivered via smartphone devices with at least one outcome measure for depressive symptoms. We aimed to examine the effects of smartphone interventions on primary depression, comorbid depression and subclinical depressive symptoms. No restrictions were placed on diagnosis or any other clinical or demographic characteristics of eligible samples.

Three independent investigators judged article eligibility (JF, JN and JT), with any disagreements resolved through discussion. "Smartphones" were defined as mobile phones with 3G or 4G Internet connectivity, along with the ability to download, install and run external applications ("apps"). RCTs of interventions delivered solely or in part via smartphone devices matching this definition, aimed at improving mental health or well-being (with depression as a primary or secondary outcome), were included in the review.

Studies using either "inactive" or "active" control groups were eligible for inclusion. "Inactive" control groups were classified as those in which participants received no intervention during the trial period (or were put into a waitlist until pre-and-post measures had been collected from both groups). "Active" control groups were categorized as those which attempted to control for the time and attention given to people in the smartphone intervention condition, by using apps not aimed at treating depression, in-person interventions, or other forms of activities or patient contact. RCTs comparing smartphone interventions to antidepressant medications were also eligible for inclusion. All eligible studies had a duration of at least one week (thus excluding studies measuring changes in mood following a single use of smartphone apps).

Data extraction

A systematic extraction form was used for each article to collect the following data: a) study information (sample size, mean age of participants, diagnostic information or relevant inclusion criteria, study length and trial quality); b) intervention features (app/program name, regularity of instructed use, smartphone program summary, any additional intervention components, details of the control condition); c) effects on depressive symptoms (changes in total depressive symptoms scored before and after smartphone and control interventions using any clinically validated rating scale). For studies which used more than one measure of depression, a mean total change was calculated by pooling outcomes from each measure.

Statistical analyses

All analyses were conducted by Comprehensive Meta-Analysis 2.0[22], using a random-effects model[23] to account for between-study heterogeneity. The total difference in changes in depressive symptoms between smartphone interventions and control conditions were pooled to compute the overall effect size of the former (as Hedges' g), with 95% confidence intervals (CI). For RCTs comparing smartphone interventions to both inactive and active control conditions, the comparative effects with active control groups were used in the primary
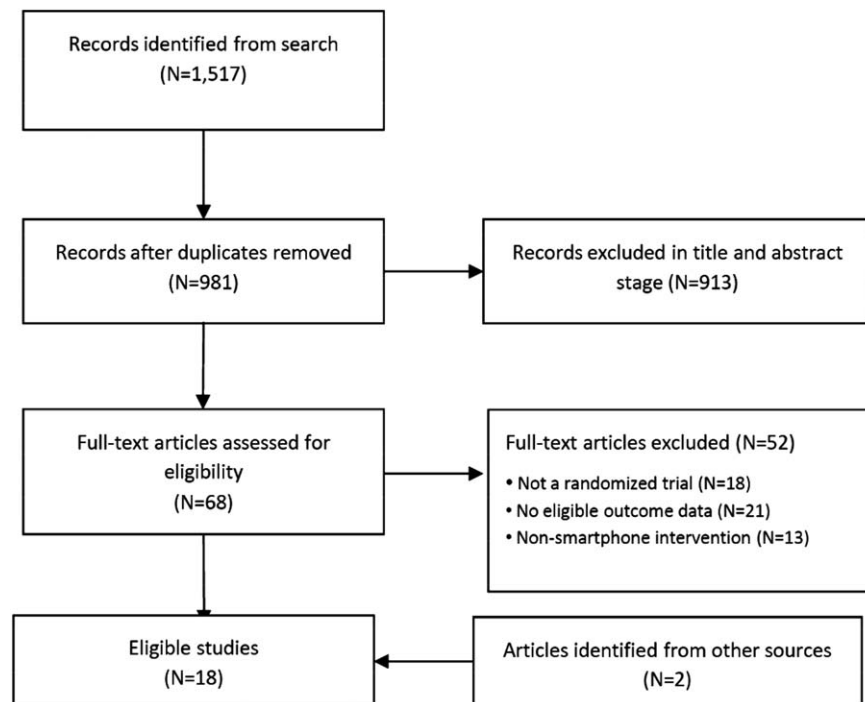
Figure 1 PRISMA flow chart of study selection

analysis. After computing main effects, a sensitivity analysis was applied to investigate effects of smartphone interventions in RCTs which used intention-to-treat analyses or had complete outcome data.

To quantify the degree to which statistical heterogeneity in the meta-analyses arose due to between-study differences, rather than due to chance, Cochran's Q (with p value) and I^2 were used. Included studies were also assessed using the Cochrane Collaboration's Risk of Bias tool. This examined study quality in six areas of trial design (sequence generation, allocation sequence concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, selective outcome reporting), ranking each area as high, low or unknown for risk of bias[24].

Risk of publication bias was examined using a funnel plot of study effect sizes, and Egger's regression test was applied to all aforementioned analyses. Furthermore, a Duval and Tweedie's trim-and-fill analysis was conducted to re-calculate the pooled effect size after removing any studies which may introduce publication bias (i.e., small studies with large effect sizes from the positive side of the funnel plot). Additionally, a "fail-safe N" was used to account for the file draw problem[25], estimating the number of non-significant unpublished trials which would be needed to cause the observed p value to exceed 0.05.

Pre-planned subgroup analyses were conducted to examine whether effects of smartphone interventions differed when comparing them to inactive or active control conditions. Additionally, we carried out a range of exploratory post-hoc subgroup and meta-regression analyses in order to examine which

factors may impact the effectiveness of smartphone interventions, particularly with regards to sample details (i.e., clinical population, age, gender) and treatment characteristics (i.e., psychological basis, technological features and length of smartphone interventions).

RESULTS

The search returned a total of 1,517 records; 981 after duplicates were excluded. Title and abstract screening removed a further 913 articles. Full versions were retrieved for 68 papers, of which 16 met eligibility criteria. Two further articles were retrieved following an additional search of Google Scholar. Thus, 18 unique RCTs were included in the meta-analysis, assessing the effects of 22 different smartphone-delivered mental health interventions. The article inclusion/exclusion process is shown in Figure 1.

Characteristics of included studies

Full details of each study are displayed in Table 1. Outcome data were available from 18 RCTs. Two papers reported outcome data in a format not suited for meta-analysis, but the corresponding authors provided the raw data to enable inclusion[26,30]. Mean sample ages ranged from 18 to 59 years (median 39 years). All but two studies[32,34] used some indication of mental health issues as inclusion criteria. For clinical populations, two studies

Table 1 Details of included studies

Study	Sample type	N (each condition)	Age (years, mean)	Design	Other intervention aspects	Outcome measure
Arean et al[26]	Self-reported mild-to-moderate depression	211,209,206	33.9	12 weeks of Project EVO (cognitive training app) vs. iPST (problem-solving therapy app) vs. Health Tips control app	None	PHQ-9
Birney et al[27]	Self-reported mild-to-moderate depression	150,150	40.7	6 weeks of MoodHacker (CBT-based depression app) vs. links to approved depression websites	Daily e-mails to provide additional digital content and prompt engagement	PHQ-9
Depp et al[28]	DSM-IV bipolar disorder	41,41	47.5	10 weeks of PRISM (mood monitoring and self-management app) vs. paper and pencil equivalent	Both groups received four sessions of individual therapy	MADRS
Enock et al[29]	Self-reported high social anxiety	158,141	34.8	4 weeks of CBM Active (cognitive bias modification training app) vs. inactive training or waitlist control	None	DASS
Faurholt-Jepsen et al[30]	ICD-10 bipolar disorder	33,34	29.3	6 months of MONARCA (self-monitoring app) vs. regular smartphone use	Patients could also contact their clinicians directly using the smartphone, in case of deterioration	HAM-D
Horsch et al[31]	Self-reported mild insomnia	74,77	39.7	6 to 7 weeks of Sleepcare (CBT-based insomnia app) vs. waitlist control	None	CES-D
Howells et al[32]	General population	57,64	40.3	10 days of Headspace (mindfulness app) vs. list-making app control	None	CES-D
Ivanova et al[33]	Self-reported social anxiety	50,51,51	35.3	10 weeks of guided ACTsmart (acceptance and commitment therapy app) vs. unguided ACTsmart vs. waitlist control	Participants also provided with pen-and-paper booklet for completing written assignments and a CD with ACT exercises	PHQ-9
Kahn et al[34]	US veterans	44, 41,42, 46	NA	16 weeks of Mission Reconnect program (using mindfulness and awareness techniques) vs. Prevention and Relationship Enhancement program vs. both programs together vs. waitlist control	Strategies for applying learnt techniques in challenging situations, and additional audio exercises	BDI-II
Kuhn et al[35]	Self-reported traumatic event + PTSD symptoms	62,58	39	3 months of PTSD Coach (app providing psychoeducation, symptom tracking and self-management strategies) vs. waitlist control	None	PHQ-8
Ly et al[36]	DSM-IV major depression	46,47	30.6	10 weeks of Behavioral Activation app plus 4 face-to-face behavioral activation sessions vs. 10 face-to-face behavioral activation sessions	None	BDI-II
Moell et al[37]	Self-reported data to diagnose ADHD	26,27	36.8	6 weeks of LivingSMART (app facilitating life organization and improving attentional control) vs. waitlist control	Computer-aided training on how to use the apps; participants were also allocated a coach to help with app usage	HADS

Table 1 Details of included studies (continued)

Study	Sample type	N (each condition)	Age (years, mean)	Design	Other intervention aspects	Outcome measure
Oh et al[38]	Older adults with self-reported memory complaints	18,19,16	59.3	8 weeks of SMART vs. Fit Brains (two cognitive training apps) vs. waitlist control	None	CES-D
Proudfoot et al[39]	Self-reported mild-to-moderate depression	126,195, 198	39	7 weeks of MyCompass (app enabling self-monitoring of problematic moods, thoughts and behaviors, tracking their severity, and receiving feedback advice and mental health management tips by SMS) vs. attention-matched and waitlist control	Computer modules provided to deliver evidence-based interventions	DASS
Reid et al[40]	Youth mental health patients	68,46	18	2 to 4 weeks of MobileType (app tracking mental health relevant thoughts and behaviors) vs. using a control app which tracks irrelevant behaviors	Participants reviewed information gathered by Mobile-Type with their general practitioner, and were given guides for managing mental health	DASS
Roepke et al[41]	Clinically significant depression	93,97,93	40.2	1 month of SuperBetter (app supporting self-esteem and self-acceptance) vs. Super-Better Plus (app adopting principles of CBT and positive psychology) vs. waitlist control	None	CES-D
Tighe et al[42]	Recent suicidal thoughts	31,30	26.3	6 weeks of ibobbly (app based on acceptance and commitment therapy principles) vs. waitlist control	24-hour helpline details available through the app in case of suicidality	PHQ-9
Watts et al[43]	DSM-IV major depression	10,15	41	8 weeks of Get Happy (CBT-based depression app) vs. computerized CBT program	Clinician contact during first two weeks to check and promote adherence	BDI-II PHQ-9

CBT – cognitive behavioral therapy, PTSD – post-traumatic stress disorder, ADHD – attention-deficit/hyperactivity disorder, PHQ – Patient Health Questionnaire, MADRS – Montgomery-Åsberg Depression Rating Scale, DASS – Depression Anxiety Stress Scale, HAM-D – Hamilton Rating Scale for Depression, CES-D – Center for Epidemiological Studies – Depression, BDI-II – Beck Depression Inventory II, HADS – Hospital Anxiety Depression Scale, NA – not available

recruited people with major depression[36,43], two individuals with bipolar disorder[28,30], one young people in primary care with any mental health condition[40]. Others recruited individuals from the general population with self-reported mild-to-moderate depression[26,27,39,41], suicidal thoughts/tendencies[42], probable attention-deficit/hyperactivity disorder (ADHD)[37], anxiety disorders[29,33], insomnia[31], or symptoms of post-traumatic stress disorder (PTSD)[35]. One further study examined older adults with memory complaints[38].

Smartphone interventions lasted between 4 and 24 weeks. Depressive symptoms were measured as a primary outcome in 12 studies, and as a secondary outcome in six. The following tools were used: the Depression Anxiety Stress Scale[44] depression subscale in three studies[29,39,40]; the Center for Epidemiological Studies Depression scale[45] in four[31,32,38,41]; the Beck Depression Inventory II[46] in three[34,36,43]; the Patient Health Questionnaire[47] in six[26,27,33,35,42,43]; the Hamilton Rating Scale for Depression[48] in one[30]; the Hospital Anxiety Depression Scale[49] in one[37]; and the Montgomery-Åsberg Depression Rating Scale[50] in one[28].

The results from the Cochrane Risk of Bias assessments are displayed in Table 2. This shows that the most frequent risk factor for bias was inadequate blinding of participants, with only five of 18 studies using intervention-matched comparators for which the participants would not be aware of their treatment/control status or of the hypothesized outcomes of the trial.

Overall effects of smartphone interventions on depressive symptoms

Figure 2 displays the pooled effect size from smartphone interventions on depressive symptoms, along with individual

Table 2 Quality assessment in included studies

Study	1	2	3	4	5	6	7
Arean et al[26]	+	+	+	+	+	+	–
Birney et al[27]	+	+	–	+	+	+	–
Depp et al[28]	+	+		+	+	+	+
Enock et al[29]			+	+	+	+	+
Faurholt-Jepsen et al[30]	+	+	–	+	+	+	+
Horsch et al[31]	+	+	–	–	+	+	–
Howells et al[32]	+	+	+	+	–	+	
Ivanova et al[33]	+	+			+	+	–
Kahn et al[34]	+			+	+	+	–
Kuhn et al[35]	+	–	–	+	+		
Ly et al[36]	+	+	+	+	+	+	
Moell et al[37]		–	+	+	+		
Oh et al[38]			–		–	+	+
Proudfoot et al[39]	+	+		+	+	+	+
Reid et al[40]	+	+	+	+	+	+	+
Roepke et al[41]	+	+	–	+	+	+	–
Tighe et al[42]	+	+	–	–	+	+	+
Watts et al[43]	+	+			–	+	

1 – random sequence generation, 2 – allocation concealment, 3 – blinding of participants and personnel, 4 – blinding of outcome assessment, 5 – incomplete outcome data, 6 – selective outcome reporting, 7 – other bias

effects from each app trialled. A random-effects meta-analysis revealed a small-to-moderate positive effect size of smartphone mental health interventions for reducing depressive symptoms in comparison to control conditions (18 studies, N=3,414, g=0.383, 95% CI: 0.24-0.52, p<0.001).

Although there was heterogeneity across the study data (Q=80.8, p<0.01, I^2=74.0%), there was no evidence of publication bias (p=0.255 in Egger's regression test), and the fail-safe N was 567 (estimating that 567 unpublished "null" studies would need to exist for the actual p value to exceed 0.05). A trim-and-fill analysis identified no outlier studies, and thus did not change the observed effect size.

When considering only the studies which used intention-to-treat analyses and/or reported complete outcome data, we found a similar effect of smartphone interventions on depressive symptoms (16 studies, N=3,320, g=0.399, 95% CI: 0.25-0.55, p<0.001; Q=80.0, I^2=77.5%).

In our pre-planned subgroup analyses, we found that effect sizes were significantly greater when comparing smartphone interventions to inactive conditions than when using active control conditions (Q=9.76, p=0.002; Figure 3). Compared to inactive control conditions, the pooled effect size across 13 smartphone interventions (N=1,674) was g=0.558 (95% CI: 0.38-0.74), indicating a moderate effect on depressive symptoms. However, when compared to active control conditions, smartphone interventions had only a small effect size on depressive symptoms (12 studies, N=2,381, g=0.216, 95% CI: 0.10-0.33).

Both studies with active and inactive controls had significant heterogeneity, but no evidence of publication bias (Table 3).

Population characteristics and effects on depressive symptoms

We also applied post-hoc subgroup analyses to studies that had used mood disorder inclusion criteria, in order to explore which populations smartphone interventions may be most effective for. As shown in Table 4, the only populations in which smartphone interventions significantly reduced depressive symptoms were those with self-reported mild-to-moderate depression (5 studies, N=1,890, g=0.518, 95% CI: 0.28-0.75, p<0.001; Q=36.6, I^2=83.6). There was no significant effect among the smaller samples with major depressive disorder, bipolar disorder and anxiety disorders (two studies each).

Mixed-effects meta-regressions were applied to explore whether continuous moderators of average age, gender distribution and sample size affected study findings, but found no indication that these factors influenced observed effect sizes (all p>0.2).

Intervention characteristics and effects on depressive symptoms

In order to gain insight into which aspects of smartphone interventions make them effective for depressive symptoms, we performed further comparative subgroup analyses after separating studies on the basis of common characteristics, such as intervention components, feedback types, and therapeutic approaches applied. The common features examined, and the results of all subgroup comparisons, are detailed in full in Table 5.

These analyses showed that smartphone interventions which involved "in-person" (i.e., human) feedback had small, non-significant effects on depressive symptoms (g=0.137, 95% CI: −0.08 to 0.35, p=0.214), whereas those which did not use in-person feedback had moderate positive effects (g=0.465, 95% CI: 0.30-0.63, p<0.001). The difference between these subgroups was statistically significant (p=0.017).

Additionally, the effects of smartphone interventions which were delivered entirely via the smartphone device (10 studies, N=2,178, g=0.479, 95% CI: 0.27-0.69, p<0.001) appeared larger than those which were not self-contained smartphone-only interventions (8 studies, N=1,236, g=0.241, 95% CI: 0.09-0.39, p=0.002), although the difference between these subgroups fell short of significance (p=0.07).

Similarly, interventions which provided "in-app feedback", such as summary statistics and progress scores, had greater effect sizes (g=0.534, 95% CI: 0.26-0.81, p<0.001) than those which did not have in-app feedback (g=0.266, 95% CI: 0.14-0.39, p<0.001), although again the difference between subgroups was non-significant (p=0.082).

The only other notable finding was that the studies of cognitive training apps had a significantly (p=0.004) smaller effect size on depression outcomes (four studies, N=836, g=0.123,

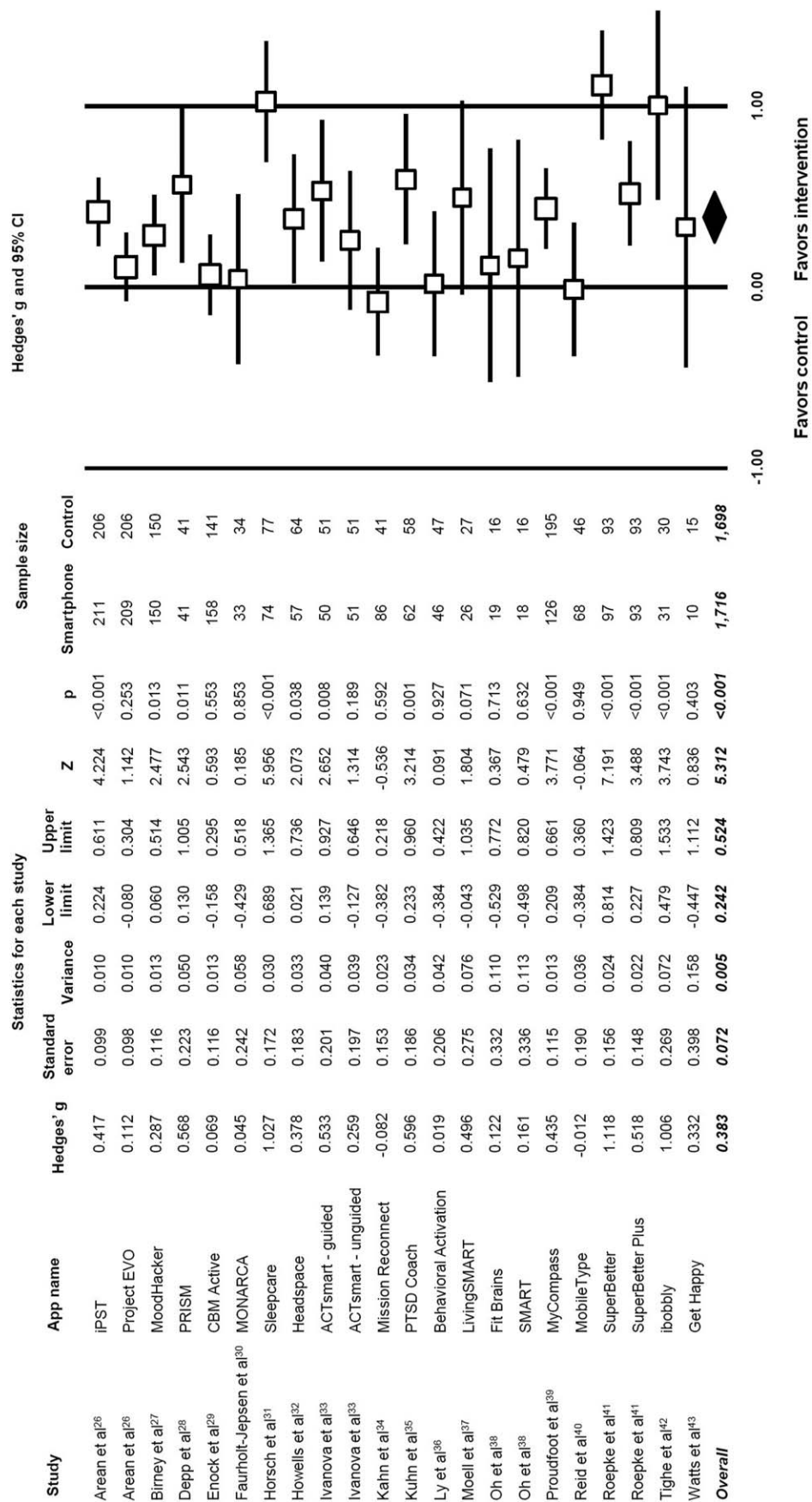| Study	App name	Hedges' g	Standard error	Variance	Lower limit	Upper limit	Z	p	Smartphone	Control
Arean et al[26]	iPST	0.417	0.099	0.010	0.224	0.611	4.224	<0.001	211	206
Arean et al[26]	Project EVO	0.112	0.098	0.010	-0.080	0.304	1.142	0.253	209	206
Birney et al[27]	MoodHacker	0.287	0.116	0.013	0.060	0.514	2.477	0.013	150	150
Depp et al[28]	PRISM	0.568	0.223	0.050	0.130	1.005	2.543	0.011	41	41
Enock et al[29]	CBM Active	0.069	0.116	0.013	-0.158	0.295	0.593	0.553	158	141
Faurholt-Jepsen et al[30]	MONARCA	0.045	0.242	0.058	-0.429	0.518	0.185	0.853	33	34
Horsch et al[31]	Sleepcare	1.027	0.172	0.030	0.689	1.365	5.956	<0.001	74	77
Howells et al[32]	Headspace	0.378	0.183	0.033	0.021	0.736	2.073	0.038	57	64
Ivanova et al[33]	ACTsmart - guided	0.533	0.201	0.040	0.139	0.927	2.652	0.008	50	51
Ivanova et al[33]	ACTsmart - unguided	0.259	0.197	0.039	-0.127	0.646	1.314	0.189	51	51
Kahn et al[34]	Mission Reconnect	-0.082	0.153	0.023	-0.382	0.218	-0.536	0.592	86	41
Kuhn et al[35]	PTSD Coach	0.596	0.186	0.034	0.233	0.960	3.214	0.001	62	58
Ly et al[36]	Behavioral Activation	0.019	0.206	0.042	-0.384	0.422	0.091	0.927	46	47
Moell et al[37]	LivingSMART	0.496	0.275	0.076	-0.043	1.035	1.804	0.071	26	27
Oh et al[38]	Fit Brains	0.122	0.332	0.110	-0.529	0.772	0.367	0.713	19	16
Oh et al[38]	SMART	0.161	0.336	0.113	-0.498	0.820	0.479	0.632	18	16
Proudfoot et al[39]	MyCompass	0.435	0.115	0.013	0.209	0.661	3.771	<0.001	126	195
Reid et al[40]	MobileType	-0.012	0.190	0.036	-0.384	0.360	-0.064	0.949	68	46
Roepke et al[41]	SuperBetter	1.118	0.156	0.024	0.814	1.423	7.191	<0.001	97	93
Roepke et al[41]	SuperBetter Plus	0.518	0.148	0.022	0.227	0.809	3.488	<0.001	93	93
Tighe et al[42]	ibobbly	1.006	0.269	0.072	0.479	1.533	3.743	<0.001	31	30
Watts et al[43]	Get Happy	0.332	0.398	0.158	-0.447	1.112	0.836	0.403	10	15
Overall		*0.383*	*0.072*	*0.005*	*0.242*	*0.524*	*5.312*	*<0.001*	*1,716*	*1,698*

Figure 2 Meta-analysis of the effects of smartphone interventions on depressive symptoms. Box size represents study weighting. Diamond represents overall effect size and 95% CI.
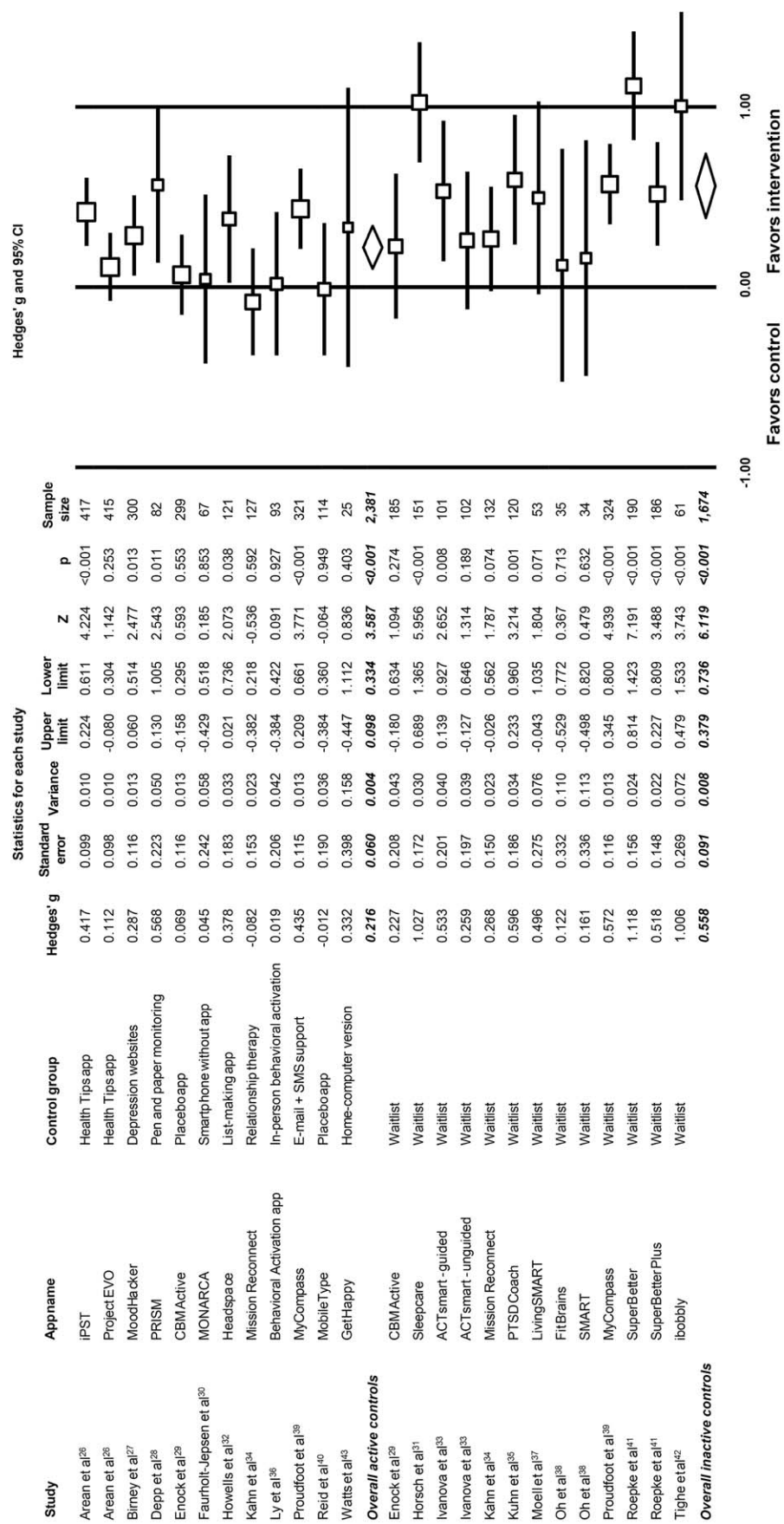
Study	Appname	Control group	Hedges' g	Standard error	Variance	Lower limit	Upper limit	Z	p	Sample size
Arean et al[26]	iPST	Health Tips app	0.417	0.099	0.010	0.224	0.611	4.224	<0.001	417
Arean et al[26]	Project EVO	Health Tips app	0.112	0.098	0.010	-0.080	0.304	1.142	0.253	415
Birney et al[27]	MoodHacker	Depression websites	0.287	0.116	0.013	0.060	0.514	2.477	0.013	300
Depp et al[28]	PRISM	Pen and paper monitoring	0.568	0.223	0.050	0.130	1.005	2.543	0.011	82
Enock et al[29]	CBM Active	Placebo app	0.069	0.116	0.013	-0.158	0.295	0.593	0.553	299
Faurholt-Jepsen et al[30]	MONARCA	Smartphone without app	0.045	0.242	0.058	-0.429	0.518	0.185	0.853	67
Howells et al[32]	Headspace	List-making app	0.378	0.183	0.033	0.021	0.736	2.073	0.038	121
Kahn et al[34]	Mission Reconnect	Relationship therapy	-0.082	0.153	0.023	-0.382	0.218	-0.536	0.592	127
Ly et al[36]	Behavioral Activation app	In-person behavioral activation	0.019	0.206	0.042	-0.384	0.422	0.091	0.927	93
Proudfoot et al[39]	MyCompass	E-mail + SMS support	0.435	0.115	0.013	0.209	0.661	3.771	<0.001	321
Reid et al[40]	MobileType	Placebo app	-0.012	0.190	0.036	-0.384	0.360	-0.064	0.949	114
Watts et al[43]	GetHappy	Home-computer version	0.332	0.398	0.158	-0.447	1.112	0.836	0.403	25
Overall active controls			*0.216*	*0.060*	*0.004*	*0.098*	*0.334*	*3.587*	*<0.001*	*2,381*
Enock et al[29]	CBM Active	Waitlist	0.227	0.208	0.043	-0.180	0.634	1.094	0.274	185
Horsch et al[31]	Sleepcare	Waitlist	1.027	0.172	0.030	0.689	1.365	5.956	<0.001	151
Ivanova et al[33]	ACTsmart-guided	Waitlist	0.533	0.201	0.040	0.139	0.927	2.652	0.008	101
Ivanova et al[33]	ACTsmart-unguided	Waitlist	0.259	0.197	0.039	-0.127	0.646	1.314	0.189	102
Kahn et al[34]	Mission Reconnect	Waitlist	0.268	0.150	0.023	-0.026	0.562	1.787	0.074	132
Kuhn et al[35]	PTSD Coach	Waitlist	0.596	0.186	0.034	0.233	0.960	3.214	0.001	120
Moell et al[37]	LivingSMART	Waitlist	0.496	0.275	0.076	-0.043	1.035	1.804	0.071	53
Oh et al[38]	FitBrains	Waitlist	0.122	0.332	0.110	-0.529	0.772	0.367	0.713	35
Oh et al[38]	SMART	Waitlist	0.161	0.336	0.113	-0.498	0.820	0.479	0.632	34
Proudfoot et al[39]	MyCompass	Waitlist	0.572	0.116	0.013	0.345	0.800	4.939	<0.001	324
Roepke et al[41]	SuperBetter	Waitlist	1.118	0.156	0.024	0.814	1.423	7.191	<0.001	190
Roepke et al[41]	SuperBetter Plus	Waitlist	0.518	0.148	0.022	0.227	0.809	3.488	<0.001	186
Tighe et al[42]	ibobbly	Waitlist	1.006	0.269	0.072	0.479	1.533	3.743	<0.001	61
Overall inactive controls			*0.558*	*0.091*	*0.008*	*0.379*	*0.736*	*6.119*	*<0.001*	*1,674*

Figure 3 Meta-analysis showing effects of smartphone interventions on depressive symptoms in comparison to active and inactive controls. Box size represents study weighting. Diamonds represent overall effect size and 95% CI.

Table 3 Effects of smartphone-delivered mental health interventions on depressive symptoms: pre-planned subgroup analyses

	Studies	Sample size (smartphone/control)	Meta-analysis				Heterogeneity			Publication bias (Egger's regression)	
			Hedges' g	95% CI		p	Q	p	I^2	Intercept	p
Main analysis	18	1,716/1,698	**0.383**	0.242	0.524	**<0.001**	80.8	**<0.01**	74.0	0.80	0.26
Intent-to-treat or complete outcome data	16	1,669/1,651	**0.399**	0.248	0.550	**<0.001**	80.0	**<0.01**	77.5	1.68	0.15
Smartphone vs. active control	12	1,195/1,186	**0.216**	0.098	0.334	**<0.001**	20.8	**0.03**	47.2	−0.49	0.34
Smartphone vs. inactive control	13	891/783	**0.558**	0.379	0.736	**<0.001**	34.9	**<0.01**	65.6	0.25	0.25

Significant values are highlighted in bold prints

Table 4 Post-hoc analyses: mood disorder samples

	Studies	Sample size (smartphone/control)	Meta-analysis				Heterogeneity		
			Hedges' g	95% CI		p	Q	p	I^2
Self-reported mild-to-moderate depression	5	917/973	**0.518**	0.282	0.754	**<0.001**	36.6	**<0.001**	83.6
Major depressive disorder	2	56/62	0.085	−0.273	0.443	0.642	0.49	0.484	0.00
Bipolar disorder	2	74/75	0.314	−0.198	0.827	0.229	2.53	0.112	60.4
Anxiety disorders	2	259/242	0.250	−0.023	0.523	0.073	4.13	0.127	51.6

Significant values are highlighted in bold prints

Table 5 Post-hoc analyses: intervention features

	Studies	Sample size (smartphone/control)	Meta-analysis				Heterogeneity			Between groups tests	
			Hedges' g	95% CI		p	Q	p	I^2	Q	p
Delivered solely via smartphone	10	1,103/1,075	**0.479**	0.271	0.687	**<0.001**	62.05	**<0.01**	80.66		
Not delivered solely via smartphone	8	613/623	**0.241**	0.088	0.394	**0.002**	13.38	**<0.01**	40.22	3.277	0.070
In-app feedback	8	750/816	**0.534**	0.258	0.810	**<0.001**	54.41	**<0.01**	85.02		
No in-app feedback	11	966/882	**0.266**	0.143	0.389	**<0.001**	18.95	**<0.01**	36.68	3.02	0.082
In-person feedback	6	309/246	0.137	−0.079	0.353	0.214	8.66	0.12	42.25		
No in-person feedback	13	1,407/1,452	**0.465**	0.302	0.627	**<0.001**	61.6	**<0.01**	75.645	**5.654**	**0.017**
Mental health focused apps	15	1,286/1,292	**0.438**	0.276	0.601	**<0.001**	2.09	0.72	0.00		
Cognitive training apps	4	430/406	0.123	−0.012	0.258	0.074	63.6	**<0.01**	74.83	**8.517**	**0.004**
Mood monitoring features	9	653/709	**0.336**	0.182	0.489	**<0.001**	16.6	0.06	82.81		
No mood monitoring	9	1,063/989	**0.418**	0.191	0.645	**<0.001**	64.0	**<0.01**	45.71	0.348	0.555
CBT-based intervention	7	541/615	**0.531**	0.339	0.722	**<0.001**	13.5	0.04	55.58		
Not CBT-based	12	1,175/1,083	**0.311**	0.130	0.493	**0.001**	59.0	**<0.01**	76.26	2.661	0.103
Mindfulness aspects	6	615/573	**0.487**	0.214	0.760	**<0.001**	38.3	**<0.01**	81.716		
No mindfulness aspects	12	1,101/1,125	**0.321**	0.160	0.482	**<0.001**	38.9	**<0.01**	66.549	1.049	0.306

CBT – cognitive behavioral therapy
Significant values are highlighted in bold prints

95% CI: −0.012 to 0.26, p=0.074) than those which focused on mental health (15 studies, N=2,578, g=0.438, 95% CI: 0.28-0.60, p<0.001).

The use of mood-monitoring softwares, cognitive behavioral therapy (CBT)-based interventions and mindfulness training did not appear to influence study effect sizes (all p>0.1 between subgroups with vs. without these features).

A mixed-effects meta-regression of study effect size with intervention length (in weeks) found indication of a slight negative relationship between the two, with smaller effects observed from longer interventions, although this correlation fell short of statistical significance (B=−0.025, SE=0.014, Z=−1.72, p= 0.086).

DISCUSSION

To our knowledge, this is the first meta-analysis to examine the efficacy of smartphone interventions for depressive symptoms. Our systematic search identified 18 RCTs, examining 22 mental health interventions delivered via smartphone devices, across a total of 3,414 participants. Thus, the literature base for this particular area has evolved swiftly, and is considerably larger than that found for smartphone interventions in other conditions. Around twice the number of eligible interventions and participants were identified compared to recent meta-analyses of smartphone interventions for diabetes and anxiety[18,19]. Furthermore, 14 of the 18 eligible studies were published within the last two years, which may reflect both the increased research interest in using apps for mental health[13] and the increased ownership, access and use of mental health apps by patients and health care organizations.

The main analysis found that smartphone interventions had a moderate positive effect on depressive symptoms, with no indication of publication bias affecting these findings. However, our subgroup analyses found that the effects of smartphone interventions were substantially larger when compared to inactive (g=0.56) than active (g=0.22) control conditions. The same pattern of effect sizes was observed in our meta-analysis of smartphone interventions for anxiety[19]. Previous reviews of other technological interventions for mental health conditions have reported similar findings, as a meta-analysis of virtual reality interventions for treating anxiety found significant effects in comparison to inactive controls, but no difference from traditional psychological treatments[51]. The extent to which the observed effects on depressive symptoms arise from using the device itself, rather than the psychotherapeutic components of the intervention, should be examined and quantified in future research, to further explore the notion of a "digital placebo" influencing findings[52].

We also explored other factors which may drive the effects of smartphone interventions for depressive symptoms, using a range of post-hoc subgroup analyses. With regards to population type, significant benefits of smartphone apps were only found for those with self-reported mild-to-moderate depression. This may be due to variations in subgroup sample sizes, as the majority of studies were conducted in non-clinical populations, thus leaving the analyses for major depression and bipolar disorder underpowered to detect significant effects. Nonetheless, the nature of smartphone interventions does appear to position them as an ideal self-management tool for those with less severe levels of depression. The observed effects indicate that these interventions are well-placed for delivering low-intensity treatment within a stepped-care approach[53], or even prevention of mild-to-moderate depression among the millions of people affected by subclinical symptoms[54]. The findings that neither age nor gender had any relationship with study effect size indicate that smartphone interventions may be applicable to a broad range of individuals.

With regards to intervention features, we found that those delivered entirely via smartphone devices had significantly greater effects than those which also involved other human/computerized aspects. Similarly, those using "in-person feedback" components had significantly smaller effects than those which did not. It seems counterintuitive that additional features/human feedback would decrease smartphone effectiveness. However, this relationship is likely due to the fact that apps not relying on external components have been designed as more comprehensive and self-contained tools. Indeed, we found some indication that studies which provided in-app feedback were more effective than those without. It should also be noted that the single study which compared a therapist-guided smartphone intervention to the same intervention without therapist support found equal effects across the two groups[33].

Smartphone interventions based on CBT significantly reduced depressive symptoms, as did those which incorporated aspects of mindfulness training or mood monitoring. However, we were not able to elucidate which of the features were most effective. A previous study which directly compared smartphone apps based on principles of either behavioral activation or mindfulness also found no overall difference between the two approaches[55]. Nonetheless, results showed that those with more severe depression experienced greater benefits from the behavioral activation app, whereas those with mild depression benefitted more from the mindfulness app. Understanding both which psychological interventions are best delivered via a smartphone and which patient populations will most benefit from smartphone-based interventions will require further research. As smartphone apps for mental health are becoming easier to create, focusing research on specific populations will enable more personalized and likely effective uses.

The trend-level negative correlation between effectiveness and length of intervention indicates that another factor to consider when designing optimal apps is user engagement[56]. Lower rates of user engagement over time have been found in numerous other mental health app studies[57-59]. Higher rates of engagement have also been associated with those apps designed for brief interactions[60], suggesting the need to customize interventions to the ways people use smartphones. While there is early

research on the optimal design and presentation of telehealth platforms[61,62], the impact on patient engagement and outcomes remains an area of nascent exploration. Understanding other factors related to app use, such as socioeconomic status, health literacy[63], technology literacy and health status[64,65], also remain important targets for further research.

A major strength of this meta-analysis is the strict adherence to a registered protocol which exactly described the search strategy, inclusion criteria, data extraction and analytic procedures. However, one drawback is that we only included smartphone interventions which have been evaluated in RCTs. Given the wide availability of mental health apps, ensuring that consumers and clinicians have access to evidence-based interventions is vital for informed decision making. While the sheer number of apps available, and their frequent updating[14,66], makes rating each impossible, research elucidating the components of effective apps and highlighting best practices may offer information immediately useful for clinical care. Of note, future studies must identify and report safety concerns regarding the use of smartphone interventions[67]. The ability of smartphones to immediately register entered mood data, compute if responses exceed a certain threshold, and if so activate emergency response systems, offer real time safety monitoring absent from traditional depression treatment.

Another limitation is the significant heterogeneity found across the analyses. Although this heterogeneity was statistically accounted for by the random-effects models when computing the effect size and respective p values, this still does indicate that significant between-study differences existed, even when subgrouping by sample/intervention type. Due to the extent of differences between studies, it was difficult to establish the single most effective components of smartphone interventions, or determine which populations these interventions are best suited for. Future studies which directly test alternative approaches against each other in non-inferiority controlled trials, while assessing outcome variation between subsamples of participants[55], would add great value to our understanding of what would constitute the optimal smartphone app for depressive symptoms, and in which populations these methods may be most effective.

In conclusion, the evidence to date indicates that mental health interventions delivered via smartphone devices can reduce depressive symptoms. However, delivering treatments via a smartphone introduces several new aspects which need to be considered, beyond the platform change alone. Specifically, we have yet to establish the ways in which user engagement, feedback loops, expectancy effects, and individual patient characteristics influence intervention outcomes. Rather than a barrier, these variables represent new opportunities for further research to optimize and personalize smartphone-based interventions.

Given the early indication of efficacy, and rapidly growing empirical research base, it is possible to envisage that continued technological advances will ultimately lead to scalable and cost-effective digital treatments for depressive symptoms[56,68]. Thus, along with continuing to design and evaluate optimal apps, further research should also be dedicated towards establishing feasible methods for implementing smartphone-based interventions within health care systems.

REFERENCES

1. World Health Organization. Depression and other common mental disorders: global health estimates. Geneva: World Health Organization, 2017.
2. Center for Disease Control and Prevention. Current depression among adults – United States, 2006 and 2008. Morbidity and Mortality Weekly Report 2010;59:1229-35.
3. Hawton K, Casañas i Comabella C, Haw C et al. Risk factors for suicide in individuals with depression: a systematic review. J Affect Disord 2013;147: 17-28.
4. McCrone PR, Dhanasiri S, Patel A et al. Paying the price: the cost of mental health care in England to 2026. London: King's Fund, 2008.
5. Cuijpers P, Sijbrandij M, Koole SL et al. The efficacy of psychotherapy and pharmacotherapy in treating depressive and anxiety disorders: a meta-analysis of direct comparisons. World Psychiatry 2013;12:137-48.
6. World Health Organization. Global health workforce, finances remain low for mental health. www.who.int.
7. Liu JX, Goryakin Y, Maeda A et al. Global health workforce labor market projections for 2030. Human Resources for Health 2017;15:11.
8. Fricchione GL, Borba CP, Alem A et al. Capacity building in global mental health: professional training. Harv Rev Psychiatry 2012;20:47-57.
9. Gulliver A, Griffiths KM, Christensen H. Perceived barriers and facilitators to mental health help-seeking in young people: a systematic review. BMC Psychiatry 2010;10:113.
10. Pedersen ER, Paves AP. Comparing perceived public stigma and personal stigma of mental health treatment seeking in a young adult sample. Psychiatry Res 2014;219:143-50.
11. Poushter J. Smartphone ownership and internet usage continues to climb in emerging economies. Pew Research Center 2016;22.
12. Aboujaoude E, Salame W, Naim L. Telemental health: a status update. World Psychiatry 2015;14:223-30.
13. Firth J, Torous J, Yung AR. Ecological momentary assessment and beyond: the rising interest in e-mental health research. J Psychiatr Res 2016;80:3-4.
14. Larsen ME, Nicholas J, Christensen H. Quantifying app store dynamics: longitudinal tracking of mental health apps. JMIR mHealth uHealth 2016; 4:e96.
15. Torous J, Firth J. Bridging the dichotomy of actual versus aspirational digital health. World Psychiatry (in press).
16. Shen N, Levitan M-J, Johnson A et al. Finding a depression app: a review and content analysis of the depression app marketplace. JMIR mHealth uHealth 2015;3:e16.
17. Powell AC, Torous J, Chan S et al. Interrater reliability of mHealth app rating measures: analysis of top depression and smoking cessation apps. JMIR mHealth uHealth 2016;4:e15.
18. Cui M, Wu X, Mao J et al. T2DM self-management via smartphone applications: a systematic review and meta-analysis. PLoS One 2016;11: e0166718.

19. Firth J, Torous J, Nicholas J et al. Can smartphone mental health interventions reduce symptoms of anxiety? A meta-analysis of randomized controlled trials. J Affect Disord 2017;218:15-22.

20. Moher D, Liberati A, Tetzlaff J et al. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. PLoS Med 2009;6: e1000097.

21. Schardt C, Adams MB, Owens T et al. Utilization of the PICO framework to improve searching PubMed for clinical questions. BMC Med Inform Decis Mak 2007;7:16.

22. Borenstein M, Hedges L, Higgins J et al. Comprehensive Meta-Analysis Version 2.0. Englewood: Biostat, 2005.

23. DerSimonian R, Kacker R. Random-effects model for meta-analysis of clinical trials: an update. Contemp Clin Trials 2007;28:105-14.

24. Higgins JP, Altman DG, Gøtzsche PC et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. BMJ 2011;343:d5928.

25. Orwin RG. A fail-safe N for effect size in meta-analysis. J Educ Stat 1983;8: 157-9.

26. Arean PA, Hallgren KA, Jordan JT et al. The use and effectiveness of mobile apps for depression: results from a fully remote clinical trial. J Med Internet Res 2016;18:e330.

27. Birney AJ, Gunn R, Russell JK et al. MoodHacker mobile web app with email for adults to self-manage mild-to-moderate depression: randomized controlled trial. JMIR mHealth uHealth 2016;4:e8.

28. Depp CA, Ceglowski J, Wang VC et al. Augmenting psychoeducation with a mobile intervention for bipolar disorder: a randomized controlled trial. J Affect Disord 2015;174:23-30.

29. Enock PM, Hofmann SG, McNally RJ. Attention bias modification training via smartphone to reduce social anxiety: a randomized, controlled multi-session experiment. Cogn Ther Res 2014;38:200-16.

30. Faurholt-Jepsen M, Frost M, Ritz C et al. Daily electronic self-monitoring in bipolar disorder using smartphones - the MONARCA I trial: a randomized, placebo-controlled, single-blind, parallel group trial. Psychol Med 2015;45:2691-704.

31. Horsch CH, Lancee J, Griffioen-Both F et al. Mobile phone-delivered cognitive behavioral therapy for insomnia: a randomized waitlist controlled trial. J Med Internet Res 2017;19:e70.

32. Howells A, Ivtzan I, Eiroa-Orosa FJ. Putting the 'app' in happiness: a randomised controlled trial of a smartphone-based mindfulness intervention to enhance wellbeing. J Happiness Stud 2016;17:163-85.

33. Ivanova E, Lindner P, Ly KH et al. Guided and unguided Acceptance and Commitment Therapy for social anxiety disorder and/or panic disorder provided via the Internet and a smartphone application: a randomized controlled trial. J Anxiety Disord 2016;44:27-35.

34. Kahn JR, Collinge W, Soltysik R. Post-9/11 veterans and their partners improve mental health outcomes with a self-directed mobile and Web-based wellness training program: a randomized controlled trial. J Med Internet Res 2016;18:18-40.

35. Kuhn E, Kanuri N, Hoffman JE et al. A randomized controlled trial of a smartphone app for posttraumatic stress disorder symptoms. J Consult Clin Psychol 2017;85:267-73.

36. Ly KH, Topooco N, Cederlund H et al. Smartphone-supported versus full behavioural activation for depression: a randomised controlled trial. PLoS One 2015;10:e0126559.

37. Moell B, Kollberg L, Nasri B et al. Living smart - a randomized controlled trial of a guided online course teaching adults with ADHD or sub-clinical ADHD to use smartphones to structure their everyday life. Internet Interv 2015;2:24-31.

38. Oh SJ, Seo S, Lee JH et al. Effects of smartphone-based memory training for older adults with subjective memory complaints: a randomized controlled trial. Aging Ment Health 2017:1-9.

39. Proudfoot J, Clarke J, Birch M-R et al. Impact of a mobile phone and web program on symptom and functional outcomes for people with mild-to-moderate depression, anxiety and stress: a randomised controlled trial. BMC Psychiatry 2013;13:312.

40. Reid SC, Kauer SD, Hearps SJ et al. A mobile phone application for the assessment and management of youth mental health problems in primary care: health service outcomes from a randomised controlled trial of mobiletype. BMC Fam Pract 2013;14:84.

41. Roepke AM, Jaffee SR, Riffle OM et al. Randomized controlled trial of SuperBetter, a smartphone-based/internet-based self-help tool to reduce depressive symptoms. Games Health J 2015;4:235-46.

42. Tighe J, Shand F, Ridani R et al. ibobbly mobile health intervention for suicide prevention in Australian Indigenous youth: a pilot randomised controlled trial. BMJ Open 2017;7:e013518.

43. Watts S, Mackenzie A, Thomas C et al. CBT for depression: a pilot RCT comparing mobile phone vs. computer. BMC Psychiatry 2013;13:49.

44. Henry JD, Crawford JR. The short-form version of the Depression Anxiety Stress Scales (DASS-21): construct validity and normative data in a large non-clinical sample. Br J Clin Psychol 2005;44:227-39.

45. Radloff LS. The CES-D scale: a self-report depression scale for research in the general population. Appl Psychol Meas 1977;1:385-401.

46. Beck AT, Steer RA, Brown GK. Manual for the Beck Depression Inventory-II. San Antonio: Psychological Corporation, 1996.

47. Kroenke K, Spitzer RL, Williams JB. The PHQ-9. J Gen Intern Med 2001;16: 606-13.

48. Hamilton M. A rating scale for depression. J Neurol Neurosurg Psychiatry 1960;23:56-62.

49. Zigmond AS, Snaith RP. The Hospital Anxiety and Depression Scale. Acta Psychiatr Scand 1983;67:361-70.

50. Montgomery SA, Åsberg M. A new depression scale designed to be sensitive to change. Br J Psychiatry 1979;134:382-9.

51. Opriş D, Pintea S, García-Palacios A et al. Virtual reality exposure therapy in anxiety disorders: a quantitative meta-analysis. Depress Anxiety 2012; 29:85-93.

52. Torous J, Firth J. The digital placebo effect: mobile mental health meets clinical psychiatry. Lancet Psychiatry 2016;3:100-2.

53. Andrews G, Cuijpers P, Craske MG et al. Computer therapy for the anxiety and depressive disorders is effective, acceptable and practical health care: a meta-analysis. PLoS One 2010;5:e13196.

54. Cuijpers P, Smit F. Subclinical depression: a clinically relevant condition? Tijdschrift voor Psychiatrie 2008;50:519-28.

55. Ly KH, Truschel A, Jarl L et al. Behavioural activation versus mindfulness-based guided self-help treatment administered through a smartphone application: a randomised controlled trial. BMJ Open 2014;4:e003440.

56. Anguera JA, Jordan JT, Castaneda D et al. Conducting a fully mobile and randomised clinical trial for depression: access, engagement and expense. BMJ Innovations 2016;2:14-21.

57. Lattie EG, Schueller SM, Sargent E et al. Uptake and usage of IntelliCare: a publicly available suite of mental health and well-being apps. Internet Interv 2016;4:152-8.

58. Owen JE, Jaworski BK, Kuhn E et al. mHealth in the wild: using novel data to examine the reach, use, and impact of PTSD coach. JMIR Mental Health 2015;2:e7.

59. Frisbee KL. Variations in the use of mHealth tools: the VA Mobile Health Study. JMIR mHealth uHealth 2016;4:e89.

60. Mohr DC, Tomasino KN, Lattie EG et al. IntelliCare: an eclectic, skills-based app suite for the treatment of depression and anxiety. J Med Internet Res 2017;19:e10.

61. Alfonsson S, Olsson E, Linderman S et al. Is online treatment adherence affected by presentation and therapist support? A randomized controlled trial. Comput Human Behav 2016;60:550-8.

62. Sarkar U, Gourley GI, Lyles CR et al. Usability of commercially available mobile applications for diverse patients. J Gen Intern Med 2016;31:1417-26.

63. Mackert M, Mabry-Flynn A, Champlin S et al. Health literacy and health information technology adoption: the potential for a new digital divide. J Med Internet Res 2016;18:e264.

64. de Alva FEM, Wadley G, Lederman R (eds). It feels different from real life: users' opinions of mobile applications for mental health. Proceedings of the Annual Meeting of the Australian Special Interest Group for Computer Human Interaction, Parkville, December 2015.

65. Ancker JS, Witteman HO, Hafeez B et al. "You Get Reminded You're a Sick Person": personal data tracking and patients with multiple chronic conditions. J Med Internet Res 2015;17:e202.

66. Nicholas J, Larsen ME, Christensen H et al. Systematic assessment of mobile apps for bipolar disorder: features and content. Bipolar Disord 2015; 17:129.

67. Faurholt-Jepsen M, Munkholm K, Frost M et al. Electronic self-monitoring of mood using IT platforms in adult patients with bipolar disorder: a systematic review of the validity and evidence. BMC Psychiatry 2016;16:7.

68. Hallgren KA, Bauer AM, Atkins DC. Digital technology and clinical decision making in depression treatment: current findings and future opportunities. Depress Anxiety 2017;34:494-501.