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Do mobile device apps designed to support medication adherence demonstrate efficacy? A systematic review of randomised controlled trials, with meta-analysis

Laura Catherine Armitage, Aikaterini Kassavou, Stephen Sutton

ABSTRACT

Objectives To estimate the efficacy of app-based interventions designed to support medication adherence and investigate which behaviour change techniques (BCTs) used by the apps are associated with efficacy.

Design Systematic review of randomised controlled trials (RCTs), with meta-analysis.

Setting Medline/PubMed, PsycINFO, Cumulative Index to Nursing and Allied Health Literature, Embase and Web of Science were searched from 1990 to November 2018 for RCTs conducted in any healthcare setting.

Participants Studies of participants of any age taking prescribed medication for any health condition and for any duration.

Intervention An app-based intervention delivered through a smartphone, tablet computer or personal digital assistant to help, support or advise about medication adherence.

Comparator One of (1) usual care, (2) a control app which did not use any BCTs to improve medication adherence or (3) a non-app-based comparator.

Primary and secondary outcome measures The primary outcome was the pooled effect size of changes in medication adherence. The secondary outcome was the association between BCTs used by the apps and the effect size.

Results The initial search identified 13 259 citations. After title and abstract screening, full-text articles of 83 studies were screened for eligibility. Nine RCTs with 1159 recruited participants were included. The mean age of participants was >50 years in all but one study. Health conditions of target populations included cardiovascular disease, depression, Parkinson’s disease, psoriasis and multimorbidity. The meta-analysis indicated that patients who use mobile apps to support them in taking medications are more likely to self-report adherence to medications (OR 2.120, 95% CI 1.635 to 2.747, n=988) than those in the comparator groups. Meta-regression of the BCTs did not reveal any significant associations with effect size.

Conclusions App-based medication adherence interventions may have a positive effect on patient adherence. Larger scale studies are required to further evaluate this effect, including long-term sustainability, and intervention and participant characteristics that are associated with efficacy and app usage.

Strengths and limitations of this study

► This systematic review and meta-analysis of randomised controlled trials offers new information on the efficacy of mobile device apps in supporting medication adherence across a range of medical conditions.

► This review is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses statement.

► A sensitive search strategy was designed, and screening of articles and data extraction were performed independently by one clinical researcher and one non-clinical researcher.

► The review protocol was prospectively registered, and the scope of the review, inclusion and exclusion criteria remained unchanged throughout.

► Two reviewers independently considered the abstracts and full-text articles for inclusion, extracted the data, and assessed the included studies for risk of bias in the same way.

Prospero registration number PROSPERO Protocol Registration Number: CRD42017080150.

INTRODUCTION

Adherence to medication is defined as the extent to which patients take medications as prescribed and agreed with their healthcare providers. The therapeutic benefit of prescribed medications is limited owing to an estimated 50% of patients not adhering to medications. The annual cost of non-adherence is estimated to exceed £930 million in England and between $100 and $300 billion in the USA. Furthermore, in 2003 the Adherence to Long-Term Therapies report published by the WHO stated that unless medication adherence is addressed, advances in biomedical technology will not achieve their potential.
Interventions for improving adherence should be reliable, contemporary, acceptable and readily available to the person. Mobile devices can meet these needs, and in 2018 Ofcom reported that 78% of adults and 95% of individuals aged 16–24 years old in the UK own a smartphone, demonstrating that smartphones are a growing part of modern life. Patients who are prescribed multiple medications are commonly in the older age group, where the percentage of smartphone users may be lower. However, smartphone ownership in those aged over 65 more than trebled between the years 2012 and 2015 from 5% to 18%, and is set to continue to increase.7

Hundreds of apps are now available to patients to support them in taking regular medication. Such apps frequently use behaviour change techniques (BCTs) to promote improvements in adherence. A BCT is defined as an active component of an intervention that is designed to change behaviour.9 To date, the efficacy of smartphone apps in supporting patient adherence to regularly prescribed medications has not been evaluated within the scientific literature using rigorous quantitative review. We have performed a novel systematic review with meta-analysis to evaluate the efficacy of smartphone apps in supporting medication adherence.

The objectives of this review are (1) to establish whether apps designed to support medication adherence demonstrate efficacy and (2) to identify the intervention characteristics and BCTs associated with efficacy. The conclusions drawn have potential to inform future large-scale public health interventions for the improvement of medication adherence.

METHODS
This systematic review is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses statement. The protocol for this review is registered with and published on the PROSPERO (International Prospective Register of Systematic Reviews) database of systematic reviews. The scope of the review, inclusion and exclusion criteria remained unchanged throughout.

Patient and public involvement
This research was done without patient involvement. Patients were not invited to comment on the study design and were not consulted to develop patient-relevant outcomes or interpret the results. Patients were not invited to contribute to the writing or editing of this document for readability or accuracy.

Data sources
Medline/PubMed, PsycINFO, Cumulative Index to Nursing and Allied Health Literature, Embase and Web of Science electronic databases were searched from 1990 to November 2018. A sensitive search strategy was designed based on preliminary searches, relevant papers and keywords. Key search terms included adherence, non-adherence, smartphone app and randomised controlled trials (RCTs). An example detailed search strategy for the Medline/PubMed database can be found in online supplementary etext 1. The reference lists of the included studies were screened for additional relevant trials.

Two reviewers (LCA and AK) independently screened all citations by title and abstract and excluded those which were irrelevant or clearly did not meet the eligibility criteria. Full-text articles were obtained for all remaining citations and screened against the eligibility criteria by the same reviewers. Any discrepancies at any stage in the screening process were settled through discussion between LCA, AK and a third reviewer (SS).

Study selection
RCTs relevant to this review were those that investigated the use of an app on a mobile device to support medication adherence. The inclusion criteria were as follows:
► Participants of any age who were taking one or more prescribed medications for any health condition and for any duration.
► An intervention group which received an app-based intervention delivered through a smartphone, tablet computer or personal digital assistant to help, support or advise about medication adherence.
► A comparator group which received usual care, a control condition which did not use an app12 or a control app which did not include any BCTs (eg, an app that was a non-health-related game13).
► Medication adherence data were reported for both the intervention and comparator groups.
The exclusion criteria were as follows:
► Complex interventions of which an app was just one component.
► Interventions delivered by text messaging or telemedicine.
► Outcome assessed was adherence to treatment components other than prescribed medication. For example, one study reported outcome data as a composite score, combining adherence to other aspects of the medical regimen (such as clinic attendance and laboratory work) with adherence to medication.14
► A study population of healthy recruits who were required to adhere to a placebo medication.
► Comparator group received an adherence intervention in another format, such as a different medication adherence app.
Published articles were required to be in the English language.

Data extraction and synthesis
Data were extracted independently by two reviewers (LCA and AK) using a custom data extraction form. Participants’ baseline characteristics were extracted, including age and sex. Raw data on medication adherence were extracted either from the published articles or by contacting the authors of the primary studies by...
email. When two measures of adherence were reported, that which was considered most comparable with the outcome measures of the other included studies was used for meta-analysis. For example, one study reported adherence according to both the number of times an electronically chipped dispenser was actioned by the patient and the weight of the dispensing canister at study follow-up, and the reviewers agreed that the action of dispensing was more comparable with that of the other included studies. Authors were contacted by email for any other data required which were not available in the published articles. Where more than one app intervention was tested, data for the intervention considered most comparable with the other studies were extracted for meta-analysis. For example, one study investigated a ‘basic app’ (which provided a unidirectional once-daily reminder) as well as an ‘advanced app’ (which provided customisable and interactive features) versus usual care. The reviewers agreed that the advanced app was most comparable with the other included studies and so data pertaining to this intervention were extracted. BCTs that were used in the app-based interventions were coded according to the Behaviour Change Technique Taxonomy. This is a structured taxonomy of techniques used by interventions to change behaviour. The BCT taxonomy does not have a code relating to performance of behaviour, and therefore where participants were required to enter whether or not medication was taken, this was coded as ‘Report whether or not the behaviour was performed’. Similarly, the taxonomy does not have a code relating to tailoring of an intervention, and therefore where a study used information gained about a person to define the content, frequency or format of an intervention, this was coded as a ‘Tailored’ intervention.

The included trials were assessed for risk of bias using the Cochrane Risk of Bias Tool. Two reviewers (LCA and AK) individually assessed each of the included articles against the criteria outlined in the tool. Any discrepancies at any stage in the coding process were settled through discussion between the three reviewers (LCA, AK, SS). Studies were considered to be at unclear risk of bias for an item when the primary article did not provide sufficient information for the reviewers to decide categorically between high or low risk of bias.

### Statistical analysis

Outcome data on medication adherence were extracted from each of the included studies at the final observation point. Authors were contacted for raw final observation point adherence data if these were not reported in published manuscripts. Trials were considered homogeneous based on the content of the intervention and comparator groups and the outcome measurement of medication adherence. The Comprehensive Meta-Analysis Software was used to estimate the overall effect of app-delivered interventions on medication adherence using a random-effects model and to perform meta-regression using the BCTs used by the apps. The Comprehensive Meta-Analysis Software conversion function was used where required to convert extracted raw data into OR and standardised mean difference between the intervention and control groups.

The characteristics of the nine studies are shown in Table 1. Publication year of the included studies ranged from 2014 to 2018. The median sample size was 102 participants (range 24–412), and the median follow-up period was 12 weeks (range 28 days–16 weeks).

The mean age of participants in the intervention group was reported by eight studies and ranged from 20.3 (SD 4.0) to 73.8 (SD 8), the mean of the mean ages across these eight studies was 56.7 years. One study did not report the mean age but presented the percentage of participants whose age fell within a series of bandwidths. The percentage of participants in the intervention groups who were female ranged from 11% to 89%, and the mean percentage of female participants among the included studies was 45.2%.

### RESULTS

The initial electronic database search identified 13,259 potentially eligible studies, and 2 were identified from the reference lists of identified articles. After removal of duplicates, 9,971 citations were screened by title and abstract. We considered 83 of these to be eligible and retrieved the full-text articles. Of these, nine studies reported outcome data eligible for meta-analysis. The screening process and reasons for exclusion are reported in figure 1.
### Table 1  Study characteristics

<table>
<thead>
<tr>
<th>Source (sample size)</th>
<th>Country of study</th>
<th>Study design</th>
<th>Health problem of target population</th>
<th>Recruitment setting</th>
<th>Participant characteristics at recruitment</th>
<th>Outcome measure</th>
<th>Follow-up interval</th>
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<td><strong>Intervention group</strong></td>
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<td>Intervention group</td>
<td>Comparator group</td>
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<td>Mean age (SD)</td>
<td>Female, n (%)</td>
<td>Mean age (SD)</td>
</tr>
<tr>
<td>Hammonds et al22 (57)</td>
<td>USA</td>
<td>RCT</td>
<td>Depression</td>
<td>College students were recruited via campus-based advertisements and the university research recruitment system.</td>
<td>20.3 (4.0)</td>
<td>25 (83)</td>
<td>20.9 (4.7)</td>
</tr>
<tr>
<td>Labovitz et al23 (28)</td>
<td>USA</td>
<td>RCT</td>
<td>Ischaemic stroke requiring secondary oral anticoagulation</td>
<td>Outpatient clinic.</td>
<td>58.3 (9.79)</td>
<td>6 (40)</td>
<td>55.5 (16.55)</td>
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<tr>
<td>Lakshminarayana et al.24 (215)</td>
<td>UK</td>
<td>RCT</td>
<td>Parkinson's disease</td>
<td>Outpatient clinic.</td>
<td>59.86 (9.13)</td>
<td>34 (36.2)</td>
<td>60.71 (10.26)</td>
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<td>Mertens et al22 (24)</td>
<td>Germany</td>
<td>RCT</td>
<td>Cardiovascular disease</td>
<td>Cardiac rehabilitation groups.</td>
<td>73.8*</td>
<td>12 (50)*</td>
<td>73.8*</td>
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<tr>
<td>Mira et al25 (102)</td>
<td>Spain</td>
<td>RCT</td>
<td>Multicomorbidity</td>
<td>Community health centres.</td>
<td>70.9 (8)</td>
<td>21 (41)</td>
<td>72.9 (6)</td>
</tr>
<tr>
<td>Morawski et al26 (412)</td>
<td>USA</td>
<td>RCT</td>
<td>Hypertension</td>
<td>Social media, mobile apps, targeted advertisements and online communications.</td>
<td>51.7 (10.5)</td>
<td>120 (57.4)</td>
<td>52.4 (10.1)</td>
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<tr>
<td>Santo et al27 (163)</td>
<td>Australia</td>
<td>RCT</td>
<td>Coronary heart disease</td>
<td>Inpatient, outpatient and rehabilitative care settings.</td>
<td>58.4 (9.04)</td>
<td>14 (13)</td>
<td>56.8 (8.64)</td>
</tr>
<tr>
<td>Shah et al28 (24)</td>
<td>USA</td>
<td>RCT</td>
<td>Cardiovascular stent</td>
<td>Patients in the immediate postoperative phase of insertion of a drug-eluting cardiovascular stent.</td>
<td>60.5 (8.9)</td>
<td>6 (46)</td>
<td>60.6 (4.1)</td>
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### Table 1  Continued

<table>
<thead>
<tr>
<th>Source (sample size)</th>
<th>Country of study</th>
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<td>Female, n (%)</td>
<td>Female, n (%)</td>
<td></td>
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<tr>
<td>Svendsen et al (^{16}) (134)</td>
<td>Denmark</td>
<td>RCT</td>
<td>Psoriasis</td>
<td>Hospital posters and the dermatology outpatient clinic.</td>
<td>18–40 29† 27 (40)</td>
<td>18–40 32† 25 (38)</td>
<td>MyPso SmarTop V.1.0 or SmarTop number 0536776.</td>
</tr>
</tbody>
</table>

*Baseline characteristics presented for the whole cohort as study design was a randomised cross-over trial.

†Number of participants in each age group (years) presented as mean age not available.

MMAS, Morisky Medication Adherence Scale; RCT, randomised controlled trial.

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### Outcome measurement methods

A number of measures of medication adherence were used in the included studies. Of the nine studies, four used the Morisky Medication Adherence Scale. A focused meta-analysis was performed, pooling results from the five studies which reported medication adherence according to the Morisky Medication Adherence Scale. The results of this meta-analysis indicated that patients who participated in medication adherence interventions reported higher adherence to prescribed medications (OR 1.83, 95% CI 1.42 to 2.36).

### Meta-analysis

Meta-analysis was performed, pooling results from the nine eligible studies which reported data on 988 participants. The results of this meta-analysis indicated that patients who participated in medication adherence interventions reported higher adherence to prescribed medications (OR 2.120, 95% CI 1.635 to 2.747) than those who did not use such interventions. The I² value was 9.8, indicating low statistical heterogeneity between the studies included in the meta-analysis.

### Behaviour change techniques

The most common BCT used by apps was ‘Tailored’, which was used by all studies. The next most common was ‘Prompts and Cues’, used by seven studies, and ‘Report whether or not the behaviour was performed’, used by six studies. The definitions of each BCT are available in online supplementary tables 3 and 4.
assessed using the kappa statistics. This revealed a ‘very good’ level of agreement (kappa=0.902, 95% CI 0.835 to 0.969). All BCTs coded for each of the study interventions are provided in online supplementary etable 3. Those coded in more than three but less than six of the nine studies were included in the meta-regression; this did not reveal any significant associations between the BCTs used and the effect size.

Risk of bias at primary study level
A summary of the risk of bias analysis is shown in table 2. Overall, all studies were deemed as unclear in their risk of bias.

The most common source of risk of bias among the studies was the absence of blinding of participants and personnel to the outcome measure. The second most common risk of bias arose from a lack of objective measurements of medication adherence. This was due to the use of self-report measures of adherence: six of the studies included in the meta-analysis reported adherence based on a self-assessment scale,12 13 16 24–26 and one study each used patient-entered medication logs in an app,22 an electronically chipped medicated foam dispenser 15 and pill counts at clinic visits.23

Self-assessment scales are a form of self-report and therefore a source of potential bias, including those of social desirability, and those arising from selective recall and the duration of the recall period.29 Two studies included in this review which measured adherence through participant self-report, sought to compare their subjective measurements of adherence with serum measurements of either drug-reactivity units or a medication’s anticoagulant effect.19 25 One study showed that serum measurement of the medication’s anticoagulant effect was similar between both the intervention and comparator groups, despite adherence measures being better in the intervention group.25 The other study revealed that drug-reactivity units were worse in the intervention group than in the control group, which conflicted with measured adherence outcome data.15 However, the presence of a drug biomarker does not equate to compliance and the absence does not equate to non-compliance.29 Indeed, drug metabolism should be taken into account for such measures of medication adherence, as physiological state and metabolic rate vary among individuals.30

Publication bias
A funnel plot of effect size estimates of each of the studies included in the meta-analysis is provided in online supplementary efigure 1. Formal analysis of plot symmetry was not performed owing to the total number of studies included being less than 10.31 Visual inspection reveals moderate symmetry of the plot.

**DISCUSSION**

**Principal findings**
We conducted a rigorous systematic review of the literature to investigate whether mobile device apps demonstrate efficacy in supporting medication adherence. A random-effects meta-analysis of the pooled results from all nine studies indicated that people who use medication reminder apps are significantly more likely to adhere to their medications than those who do not, but the results should be interpreted with caution owing to the data for six of the nine studies included in the meta-analysis being based on self-reported measures of adherence.

Intervention effects reported between the studies varied, as shown in the forest plot (figure 2). All nine studies reported that their mobile app-delivered interventions demonstrated efficacy in supporting medication adherence, although in five studies the interventions did not have a statistically significant effect.12 13 16 22 25 Although the I² measure of statistical heterogeneity was low, and
Table 2  Summary of risk of bias

<table>
<thead>
<tr>
<th>Source</th>
<th>Allocation sequence adequately:</th>
<th>Free from baseline imbalance?</th>
<th>Blinding of participants and personnel</th>
<th>Objective outcome assessment</th>
<th>Incomplete outcome data adequately addressed</th>
<th>Free from selective outcome reporting</th>
<th>Was the intervention independent of other bias?</th>
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<td>Hammonds et al   22</td>
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<td>Labovitz et al     23</td>
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<td>Lakshminarayana et al  24</td>
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<td>Mertens et al      22</td>
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<td>Mira et al         25</td>
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<td>Morawski et al     26</td>
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<td>Santo et al        16</td>
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●: low risk of bias; ●: unclear risk of bias; ●: high risk of bias.
Strengths and weaknesses of the primary studies

Based on the assessment of the risk of bias undertaken as part of this review, the evidence included in this meta-analysis is of moderate quality. The most common weaknesses in this evidence are small sample size, lack of objectivity of the outcome measurement, lack of blinding of participants and personnel, and limited follow-up time.

Strengths and weaknesses at review level

Our comprehensive literature search and search of the PROSPERO database indicate this is the first systematic review and meta-analysis to investigate the efficacy of mobile apps in supporting medication adherence. This offers new knowledge to the field; however, as such there are no comparable studies with which the results of this review can be compared.

The pooling of data from a group of studies which used a variety of methods for measurement of medication adherence and targeted at patients with different healthcare problems is a limitation of this review. As further studies in this field are published, it will become more feasible to conduct subgroup analysis or meta-regression analyses against variables such as age, health condition, targeted medication, intervention content and app characteristics. A standardised protocol for measuring adherence and reporting the content and characteristics of app interventions may facilitate further analysis of the latter two variables.

This review cannot report on the likely sustainability of the effect on medication adherence given the short study duration of the studies included in this review; across the nine studies, the maximum follow-up period was 16 weeks. Furthermore, the optimal frequency and duration of app use that are required to achieve improvements in medication adherence are not known. Long-term and large-scale research is required in this field to address the question regarding the sustainability of the observed effect found in this meta-analysis.

This systematic review and meta-analysis included studies with a range of ages of participants. The lowest mean age of participants in the included studies was 20.9 years and the highest was 70.9 years. The ubiquity and use of mobile devices vary across age groups, and while the majority of participants in these studies were aged 50 and over further research to establish the acceptability and usability of adherence apps among different age groups of adults would be valuable.

Finally, the reviewers set out to perform a meta-regression of the BCTs used by app interventions in the included studies. However, this did not reveal any significant association between the BCTs used and the observed effect size. This may be owing to the low estimate of heterogeneity and the small number of eligible included studies.

CONCLUSIONS

This meta-analysis indicates that medication adherence interventions delivered by smartphone apps are associated with higher levels of self-reported adherence to prescribed medications. However, when considering whether to recommend an app to patients, these results should be interpreted with caution owing to variance in effect size seen between the studies, the small sample sizes taken from different patient populations, the heterogeneity among included studies and the prominence of self-reported scales to measure medication adherence. Further research should aim to first establish a standard protocol for measuring and reporting of adherence in the context of evaluating digital interventions. Subsequently, further research investigating intervention characteristics associated with an app’s effect and the sustainability of any effect gained would add to this field.

REFERENCES


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