A mobile app for postoperative wound care after arthroplasty: Ease of use and perceived usefulness

H. Scheperab,⁎, R. Deroega, R. Mahdadb, R.J.P. van der Walc, R.G.H.H. Nelissend, L.G. Vissera, M.G.J. de Boera

a Department of Infectious Diseases, Leiden University Medical Centre, Albinusdreef 2, 2333ZA, the Netherlands
b Department of Orthopaedic Surgery, Alrijne Hospital, Leiderdorp, the Netherlands
c Department of Orthopaedics, Leiden University Medical Centre, Albinusdreef 2, 2333ZA, the Netherlands

corresponding author.
E-mail address: h.scheper@lumc.nl (H. Scheper).

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1. Introduction

A prosthetic joint infection (PJI) is a feared complication for patients with a total joint arthroplasty. The reported incidence of PJIs ranges between 0.5–1.0% and 0.5–2.0% for hip and knee arthroplasty, respectively. This incidence is largely underestimated due to inadequate registration of infections [1]. Inadequate treatment of wound complications results in hospital readmission, revision surgery, long term antibiotic treatment and, in the worst case, removal of the prosthesis [2]. In The Netherlands, most patients are discharged the first or second postoperative day after arthroplasty, which is associated with faster functional recovery and lower costs [3]. Consequently, patients are responsible for monitoring their post-operative wound at home.

This put them at risk for a delayed diagnosis of wound infections. This delay may lead to chronic PJI with extensive revision surgery with removal of the implant [4].

A mobile woundcare app used by patients after joint arthroplasty underscores the importance of adequate wound monitoring. Daily revision of the wound by patients may lead to improved monitoring, increased awareness for complications and, consequently, earlier consultation of the treating physician. There is evidence for distant post-discharge monitoring of postoperative patients. Reports have shown that post-operative telephone review is cost-effective and acceptable for patients with no underreporting of complications [5,6]. Another report showed a significant reduction in unnecessary emergency room visits by using email with smartphone photography in post-

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hypsipedia patients [7]. The use of smartphones for monitoring recovery in post-operative patients at home has been shown to be feasible and acceptable to patients and surgeons, although patients were concerned about the lack of timely responses from healthcare [8,10]. To the best of our knowledge, no studies have been performed yet in which a mobile woundcare app was used with an integrated alert system for patients when to contact their physician. We hypothesized that a mobile woundcare app after joint implantation is useful for patients. We hypothesized that using such an app may lead to increased patient involvement, early detection of wound problems and prevention of chronic PJI [9]. In this prospective study we investigated the ease of use and perceived usefulness of using such a mobile woundcare app in patients after joint arthroplasty.

2. Methods

All patients having a primary or revision total joint arthroplasty during the period July to December 2017 were eligible for participation in a prospective cohort study conducted at an academic hospital (Leiden University Medical Center) and a large regional teaching hospital (Alrijne Hospital). The primary objective was to investigate the ease of use and the patient’s perceived usefulness of the woundcare app. Secondary objectives were the number of alerts, the number of calls to the treating physician during the study period, the amount of days the app was actually used, patient-reported wound infection and the concordance between patient-reported outcome and physician-reported outcome. The study was approved by the institutional Medical Ethical Committee (protocol nr. P17.091).

All patients scheduled for total joint arthroplasty were asked to participate during their hospital admission. Inclusion criteria were at least 18 years old, able to provide written informed consent and ownership of an android or iOS 9.0 or newer smartphone. Informed consent was obtained by the study coordinator who also guided each patient with downloading of the app. Instructions were given to patients how to use the app and how to fill in the daily review tasks. The study coordinator was available for the first 2–3 postoperative days for practical assistance and could be called during the study if needed. People who were unable to understand or read Dutch were excluded. After 30 days, patient files were reviewed to check for concordance between patient-reported and physician-reported outcome with respect to wound complications. All patients were seen in the outpatient clinic two and six weeks postoperatively. Clinicians were instructed about the underlying algorithm in the app and the alert system that could prompt patients to call them. It was left to the judgment of the treating clinicians to decide whether patients needed a clinical review or that a telephonic review was sufficient. The nurses on the ward were instructed about postoperative use of the app so they could help patients with filling in. Statistical analysis was done using SPSS (IBM SPSS Statistics version 24.0, Armonk, USA).

2.1. Mobile woundcare app

A woundcare app (Fig. 1) was developed by a digital innovation company (Innovidattic, Delft, The Netherlands) with intellectual input from the authors.

All data entered in the app were pseudonymised and stored on a local ISO 27001 certified data management server at the coordinating hospital. A key for disclosure was stored on a local data safety folder. The app consisted of an introductory page collecting basic patient characteristics followed by daily short questionnaires regarding the patient’s wound. Patients recorded redness, pain (by visual analogue score,VAS), wound leakage, fever and a picture of the wound could be taken (Appendix 1). After 30 days, the patient-reported outcome was scored by the patient (i.e. PJI). Based on the daily questionnaires, an algorithm created daily a risk-score. A threshold score, developed by consensus meetings of the authors (HS, MB, RG, LV) defined above which the wound was thought to be at risk for being infected (Appendix 2). If the score exceeded this threshold, an alert message on the smartphone advised patients to contact their treating physician within 24 h. The orthopaedic ward could be called directly via a push button in the app. Prior to the study, caregivers were instructed to register every contact in the electronic patient files. Apart from using the app, post-operative wound care did not differ between study participants and patients who were not included.

2.2. Ease of use and perceived usefulness

The questionnaires that were used to test for perceived usefulness and ease of use (Likert scale) were adapted from questionnaires that were developed for user acceptance of information technology [10] (Appendix 3). The app provided a link to the online questionnaires on day 15 and day 30 of the study. Additionally, patients received a reminder for the questionnaire by email. Responses followed a 5-point Likert scale from “strongly agree” to “strongly disagree.” Results of day 15 and day 30 were compared for both questionnaires with a paired-samples t-test. Patients who did not manage to fill in one of the questionnaires were contacted by telephone after 30 days to grade the app and to explore the reasons for not filling in the questionnaire.

3. Results

Of 127 eligible patients, thirty patients (24%) did not own a smartphone. Of the remaining 97 patients, 69 patients (71%) were included (Fig. 2).

The median age was 68 years (range 33–90) (Table 1). Forty-one patients (59.4%) used the app until day 30. Nine patients (13.0%) stopped using the app immediately after the first or the second day of use. On average, the app was used by 43 patients per day. In total, the app was used on 1317 postoperative days (64% of the total amount of 30 postoperative days in 69 patients). The overall amount of responses tended to decline slowly over time (Fig. 3).

3.1. Perceived ease of use and perceived usefulness

The additional questionnaires about ease of use and usefulness were filled in by 31 patients (44.9%) on day 15 and by 37 patients (53.6%) on day 30. Fifteen patients (21.7%) filled in both questionnaires. The mean score for ease of use at day 15 was 4.2 (on a scale of 1–5) and 4.1 for perceived usefulness (Figs. 4 and 5).

The scores on day 30 were comparable to day 15 for ease of use (score 4.2, p = 0.43) and perceived usefulness (score 4.0, p = 0.40). The average satisfaction with the app at day 15 was 8.2 (on a scale of 1–10; range 6–10). Sixteen patients (23%) who did not fill in a questionnaire at all were contacted by telephone, to have information on user-friendliness or hick-ups when using the app. Eight of them could be reached and were interviewed with predefined questions. The mean satisfaction-score of the app among them was 7.9 (range 7–10). The majority of these patients had stopped using the app prior to reaching the day of the questionnaire (day 15). Reasons for discontinuation were malfunction of the smartphone (n = 1), the app had stopped giving reminders (n = 2) or patients had forgotten to fill in the app (n = 6).

3.2. Alerts

An alert was sent to patients on 29 (2.2%) of the 1317 days the app was used. Ten alerts were sent because the score exceeded five points, three alerts because the score exceeded four points on two consecutive days and 16 alerts because the score exceeded three points on three consecutive days (see also Appendix 2). Thirteen patients responded on the question of the online questionnaire specifically asking if the hospital took their calls, based on alerts, seriously (score 3.7 on day 15 and 3.6 on day 30, Fig. 4). No single record of patient calls was found in the electronic patient files. Also, it appeared that in the iOS version of the
app there was a technical flaw in the algorithm resulting in only sending alerts when the score exceeded five points. Due to this flaw, 28 out of 57 alerts were not sent to the patient.

Fig. 1. Screenshots of the woundcare app (with Dutch language).

English translation. Screen 1: Woundcare. Three days to go. You can now fill in the daily questionnaire. Screen 2: Does the wound leak? No, minimal (less then 2 x 2 cm on the bandage), a little (more than 2 x 2 cm on the bandage), fair (exchange of two bandages), strong (exchange of more than two bandages). Screen 3: Give your pain a score (Visual Analogue score 0–10). Screen 4. Advice: Your scores of today may fit with a wound complication. We advise you to contact your orthopaedic surgeon within 24 h or (if out-of-office hours) with the emergency department.

Fig. 2. Selection and inclusion of patients.

Table 1
Baseline characteristics of 69 patients who used the Woundcare app.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Count</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (median, range)</td>
<td>68</td>
<td>33–90</td>
</tr>
<tr>
<td>Female/Male</td>
<td>46/23</td>
<td></td>
</tr>
<tr>
<td>University Medical Center (n)</td>
<td>19</td>
<td>28%</td>
</tr>
<tr>
<td>Regional hospital (n)</td>
<td>50</td>
<td>72%</td>
</tr>
<tr>
<td>Operating System Mobile Device</td>
<td></td>
<td></td>
</tr>
<tr>
<td>iOS (n, %)</td>
<td>33</td>
<td>48%</td>
</tr>
<tr>
<td>Android (n, %)</td>
<td>36</td>
<td>52%</td>
</tr>
<tr>
<td>Joint arthroplasty</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hip</td>
<td>32</td>
<td>46%</td>
</tr>
<tr>
<td>Knee</td>
<td>37</td>
<td>54%</td>
</tr>
<tr>
<td>Past medical history</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>9</td>
<td>13%</td>
</tr>
<tr>
<td>Rheumatoid arthritis</td>
<td>8</td>
<td>12%</td>
</tr>
<tr>
<td>Megaprosthesis</td>
<td>2</td>
<td>3%</td>
</tr>
</tbody>
</table>

Fig. 3. Number of patients completing daily forms in the app.

3.3. Postoperative course

Forty-one patients filled in the outcome score on complications on day 30. Concordance of patient-reported and physician-reported outcome was reached in 33 patients (80%) (Table 2).

Discordance occurred in seven patients who did not have a complication, but scored “I don’t know” as outcome. The only patient (1.5%) in our study that developed a PJI on day 30 scored a “suspected PJI, but appeared to be no infection”.

One patient (1.5%) had revision surgery because of repeated dislocations of the hip joint. Two patients (2.9%) developed a deep venous thrombosis of the leg. Four patients (5.8%) reported a temperature > 38.0 °C at least once during the 30 day postoperative period. Postoperative wound leakage was reported by thirty-seven patients (53.6%); the majority of the patients reported this on the second and third postoperative day (Fig. 6).

Wound leakage duration was present during a mean 2.2 days (range 1–11). From day 18 onwards, five patients reported new wound leakage for one to five days. The leakage reported in the fourth postoperative week corresponded with the patient who developed a prosthetic joint infection. This patient scored an unchanged wound for four weeks and leakage and fever since one day before admission with a PJI.

4. Discussion

We found that introduction of a mobile woundcare app resulted in a high perceived usefulness and ease of use. Patients felt engaged with their health and with the care provided by the hospital. This
involvement was consistent during the use of the app. The number of patients completing daily forms in the app declined only mildly, confirming patient engagement with their own woundcare.

Clinical applications of mobile e-health by patients can be a valuable tool in health care management. With the increasing use of medical apps, it is important to develop e-tools that support patients and clinicians in improving health care. The high inclusion rate in this study stresses patient willingness to use mobile apps for postoperative wound monitoring. This is in line with recent surveys showing that using an app for surgical wound monitoring, including taking digital wound photographs, is supported by most patients [11,12]. Of all eligible patients aged 65 years or more, smartphone ownership in this study was 76%. Most likely, this will increase over the next years resulting in more patients who may benefit from medical apps.

In our woundcare app postoperative follow up care by patients themselves is integrated with an (wound)risk assessment that supports the patient when to contact their physician. Other studies have suggested that the use of mobile e-health led to more engagement of patients with their treatment [13–15]. Importantly, negative experiences might arise when daily asked to monitor a postoperative wound; however, these were not reported by patients. The response rate for the questionnaires on day 15 and 30 of only 77% might introduce a selection bias with skewed positive responses. Therefore, patients who did not fill in a questionnaire were interviewed later by telephone and, using the same grading system, those non-responders showed comparable high satisfaction rates as responders.

### 4.1. Cost-effectiveness

If postoperative infections can be treated at an earlier stage, devastating chronic PJI can be prevented. The costs of revision surgery for one patient (estimated costs around 30,000 euro) are about the same as the costs for the development of this app [16]. The app may be cost-effective by preventing diagnostic delay but larger studies need to be done to show cost-effectiveness. The app worked well for the only patient who developed a PJI; this patient scored eight points on the day of admission (score based on heavy leakage and a high pain score). She had not used the app on the day prior to admission; two days before admission her score was four. Good compliance is needed in order to really benefit from the app. Of all included patients, 59% used the app as intended until day 30. One of the main - understandable - reasons for discontinuation was that patients deemed further use of the App irrelevant, since their postoperative recovery went uneventful. For these patients, further use of the app would obviously not have resulted in improved clinical outcome.

### 4.2. Patient-reported and physician-reported outcome

Concordance between patient-reported and physician-reported outcome on wound healing is important in order to estimate the

<table>
<thead>
<tr>
<th>Physician-reported outcome</th>
<th>Day 15 (n=31)</th>
<th>Day 30 (n=37)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I don’t know</td>
<td>1 2 3 4 5</td>
<td>1 2 3 4 5</td>
</tr>
<tr>
<td>No infection</td>
<td>4.2 (3-5)</td>
<td>3.7 (3-5)</td>
</tr>
<tr>
<td>Suspicion PJI</td>
<td>4.0 (3-5)</td>
<td>3.9 (3-5)</td>
</tr>
<tr>
<td>PJI</td>
<td>3.7 (3-5)</td>
<td>3.6 (3-5)</td>
</tr>
</tbody>
</table>

*16 patients scored ‘not applicable’

Table 2

Concordance between patient-reported and physician-reported outcome in 41 patients who used the app until day 30.
Zekerheid.

at the LUMC and by a grant of a healthcare insurance company Zorg &
Funding

wound care with this app may have an additional value.

addressed in a large cohort study. A causal relationship would under-

wound leakage and a postoperative PJI. This association should be

powered for finding an association between the length and severity of

removal of the plaster at two weeks postoperative. This study was not

patients completed the app every day. The recurrence of leakage on day

wound leakage, the majority on the second and third postoperative day.

The true incidence of wound leakage may be higher, since not all pa-

symptom of a PJI is essential. Maathuis et al. reported that 10% of all

leakage as being part of normal postoperative course or being a

criteria for wound infection after arthroplasty (pain, fever, leakage,

redness) that is easy to use for patients (Appendix 2). To avoid false-
negative results the threshold for sending an alert was put low, resulting

in alerts in ten individual patients, while only one patient developed a

PJI. Most of these alerts were based on a high VAS score; for these

patients a mobile app may lower the threshold to contact the treating

physician to optimise their pain medication.

4.4. Wound leakage and infection

Currently, the importance of postoperative wound leakage as risk

factor for PJI is largely unknown [16]. Differentiation between wound

leakage as being part of normal postoperative course or being a

symptom of a PJI is essential. Maathuis et al. reported that 10% of all

wound leakages resulted in a PJI (unpublished results). Currently, a

multicenter study on the treatment of postoperative wound leakage in

elective hip and knee arthroplasty is done [16]. Immediate extensive

surgical debridement is the cornerstone of treatment for an acute PJI

but if done unnecessary it exposes patients to an additional risk for

infection. Many patients in this study (59.4%) reported postoperative

wound leakage, the majority on the second and third postoperative day.

The true incidence of wound leakage may be higher, since not all pa-

patients completed the app every day. The recurrence of leakage on day

18 in five patients might be explained by easier wound monitoring after

removal of the plaster at two weeks postoperative. This study was not

powered for finding an association between the length and severity of

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accuracy of patients to determine their own diagnosis. The discordance
rate of 20% in this study is probably secondary to outcome options that
were not presented clearly in the app. The ‘I don’t know’ category
(Table 2) was too vague in hindsight and will be omitted in the next
version of the app. We estimate that, when adjusting the options in the
app, the concordance comes close to 100%, but reliable estimation of
concordance can only be addressed in a larger study.

One of the objectives of this study was to determine the number of alerts that led to a call to the treating physician resulting in a change in
treatment. The one patient that developed a PJI did receive an alert and
was admitted to the hospital on the same day. Although physicians
were instructed to report all app-based phone calls by patients in the
 electronic patient files, this apparently did not happen. This can partly
be explained by the reduced number of alerts (due to the technical
problems) but also by underreporting. The technical problems under-

score the importance of pilot studies like this to find and resolve these
issues. Visual integration of all app data into patient’s electronic files
may lead to improved registration, as this supports physicians to in-
terpret a clinical situation more accurate when called by their patients.
Currently, real-life visual integration of the clinical data of the app in
the electronic patient files is implemented in our hospital.

4.3. Scoring system for wound infection

As far as we know, there is no validated grading system to score a
postoperative wound. A systematic review of surgical infection scoring
systems found one scoring system for postoperative sternal wounds, but
this was developed for scoring by physicians and not suited for patient
monitoring [17]. We developed a grading system based on the classical
criteria for wound infection after arthroplasty (pain, fever, leakage,
redness) that is easy to use for patients (Appendix 2). To avoid false-
negative results the threshold for sending an alert was put low, resulting
in alerts in ten individual patients, while only one patient developed a
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Authors’ contributions

HS and RD wrote the original manuscript and analysed the data. RW, RN, MB, LV and RM critically reviewed the manuscript. HS wrote the
final version.

Conflicts of interest

The authors declare they have no conflicts of interest

Author statement

This paper, “A mobile app for postoperative wound care after ar-
throplasty: ease of use and perceived usefulness” has not been published previously. This paper is not under consideration
for publication elsewhere. The publication of this paper is approved
by all authors. If accepted, it
will not be published elsewhere in the same form, in English or in
any other language, including
electronically without the written consent of the copyright-holder

Summery points

What was known:

- Early postoperative discharge after joint arthroplasty may lead
to decreased wound monitoring.
- A mobile woundcare app with an integrated algorithm to
detect complications may lead to improved monitoring and
earlier treatment of complications.

What this study adds:

- A postoperative woundcare app with an alert communication
on possible wound problems resulted in a high perceived
usefulness and ease of use.
- Patient involvement in postoperative wound care is high when
using a mobile app with an integrated alert system

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paration of the study in Alrijne Hospital

Appendix A. Supplementary data

Supplementary material related to this article can be found, in the

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joint infection after 32,896 primary total hip arthroplasties: a prospective cohort
179–188.
increased risk of revision surgery due to deep infection following fast-track hip
monitoring via mobile app on the number of in-person visits following ambulatory


