The safety and effectiveness of minor injuries telemedicine

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Objectives: To determine the safety of minor injuries telemedicine compared with on-site specialist care, current practice, and a robust gold standard, and to assess the clinical effectiveness of this new technique.

Methods: Patients presenting to a peripheral hospital within 10 days of injury were separately assessed by each of: an emergency medicine specialist based at a district general hospital using telemedicine, a second on-site emergency medicine specialist, and an on-site general practitioner (representing current practice). The primary outcome measure was discrepancies between these three medical assessments and a gold standard. All patients were subsequently randomised to follow one of the independent treatment plans generated by the above assessments. Secondary outcomes were recovery and further use of healthcare services measured seven days after recruitment, and consultation duration.

Results: 600 patients were recruited over a 12 month period. Overall, 73 discrepancies were identified, with 12 important over-treatments and 11 important under-treatments. No consultation modality was clearly superior to any other, and there were no statistically significant differences in the secondary outcomes of clinical effectiveness measured at seven days. The mean duration of a telemedicine consultation (6.0 min) was almost twice as long as an on-site specialist (3.1 min) or on-site general practitioner consultation (3.4 min) (p<0.0001 in both cases).

Conclusions: Minor injuries telemedicine is safe and clinically effective, providing care that is equivalent to specialist on-site assessment and the current practice of treatment by a general practitioner. There is no evidence that telemedicine provides superior care, and there are a number of process issues that may impede successful implementation of this new technique.
Box 1 Inclusion and exclusion criteria

Inclusion criteria
- The patient required, in the view of the nurse assessing them, an opinion from a doctor.
- The onsite specialist was present at the peripheral site.
- The patient had sustained an injury within the previous 10 days.

Exclusion criteria
- Patients unable or unwilling to be followed up. This was required as a safety measure, and to establish a gold standard in each case.
- Patients unable to provide informed consent.
- Patients with major injuries who clearly required immediate transfer.
- Patients who had already participated in the study.

available. The onsite specialist played no routine part in the peripheral unit, and was present only for the purposes of the trial. The onsite specialist attended the peripheral unit from 8 am to 8 pm on a total of 156 days spread throughout a 12 month period. Recruitment was not conducted overnight as patient attendances were very infrequent, and there was no senior doctor immediately available for telemedicine at the main hospital. An experienced nurse, who was also able to request radiographs of the peripheral skeleton, assessed each patient on arrival according to normal practice. Patients fulfilling the inclusion and exclusion criteria (box 1) were referred to the onsite specialist for recruitment, with the nurse prospectively recording what would have happened to the patient had they not entered the trial.

Written consent was obtained from all patients, or from a parent or legal guardian for children. Approval for the study was gained from the East Gloucestershire Ethics Committee.

Study design
To evaluate safety the study directly compared three independent medical assessments with a gold standard. In addition, a prospective, randomised, and blinded trial was conducted to measure clinical effectiveness. Process measures relating to consultation duration, radiograph requests, and follow up were also collected prospectively. Figure 1 is a summary trial flow diagram.

Patients were independently assessed by each of: the onsite emergency medicine specialist, a second emergency medicine specialist (either a consultant, staff grade, or specialist registrar) from the main hospital using telemedicine, and an onsite GP (if available). GPs visited the peripheral unit on a rota basis from their individual surgeries, and were therefore not always immediately available to assess recruited patients. Participants were generally unwilling to wait more than one or two hours to see a GP if two emergency medicine specialists had already seen them, and GP assessment was therefore omitted if a GP was not readily available in the peripheral unit.

Assessment by telemedicine entailed an initial telephone call to the main emergency department to indicate that a teleconsultation was required. When a senior doctor became available they started the telemedicine link and undertook a remote consultation with the patient, facilitated by the nurse.

Radiographs requested by the nurse before medical consultation were made available to all doctors during their assessment. If any doctor felt that further radiographs were required they requested these as part of their treatment plan, but these radiographs were not made available to the other doctors unless they had also independently requested them.

Each doctor made an independent written record of their diagnosis, radiological interpretation, and itemised treatment plan with follow up arrangements. The doctor did not discuss these with the patient. When all treatment plans had been recorded patients were randomised to follow one of these plans using one-to-one-to-one block randomisation with a variable block size. A second nurse opened a sealed, opaque envelope corresponding to the patient’s study number. This indicated which plan should be followed. The treatment instructions from this plan were transcribed onto a loose sheet, which was returned to the first nurse to enact. This ensured that neither the treating nurse nor patient knew which plan was being followed. Where the GP treatment plan was indicated a second randomised option was also stated for use if the GP assessment had been omitted.

All treatments were carried out by nursing staff, with the patient referred to a GP or transferred to the main hospital if further medical intervention was indicated on the treatment plan.

Participants were reviewed seven days after their initial assessment, as substantial improvement is common by this time. All participating doctors were equally aware of the trial protocol and arrangements for review. Patients remained blinded to which treatment plan they had followed. A standardised questionnaire was completed by each participant (or consenting adult in the case of a younger child). A gold standard diagnosis was then established by direct clinical assessment of each patient by the principal investigator, the report of a consultant radiologist blinded to the study and referral to a consultant specialist or further follow up until complete resolution.

Outcome measures
Outcomes were assessed in terms of safety, process, and clinical effectiveness. Safety was assessed by comparing each arm of the trial to the gold standard. All discrepancies in diagnosis, radiological interpretation, and treatment plan were submitted to an expert panel comprising 10 consultants in emergency medicine who were not otherwise involved in the study. Panel members did not know which trial arm had given rise to each discrepancy, and were asked to score these according to the scale shown in box 2, which was developed specifically for this research. The scores allocated by each of the 10 panel members were used to calculate both a mean and median discrepancy score. The mean score was selected for further analysis because it served as a better discriminator between the 44 discrepancies that had a median score of +1 or −1. Process outcomes were the duration of each consultation and the frequency with which radiographs and follow up appointments were requested. Measures of clinical effectiveness were derived from the review questionnaires, and are listed in box 3.

Sample size
The sample size was calculated for the primary outcome measure of safety, rather than the secondary outcome measures of clinical effectiveness that were addressed in the randomised trial. We selected a sample size of 600 because the absence of a significant discrepancy in this number would allow us to state, within 95% confidence limits, that the risk of significant discrepancy in actual clinical practice would lie somewhere between zero and 1 in 200.16 This is comparable to existing and accepted risks in emergency medicine.17 With regards to the secondary outcomes of clinical effectiveness, this sample size is sufficient to detect a 12% difference between telemedicine and the onsite specialist (α = 0.05, two sided, and power 0.8).
Data handling and statistical methods

Data management and analysis were performed using a computerised Stata database (Stata Corporation, Texas, USA). In the randomised trial data were analysed on an intention to treat basis incorporating the two stage randomisation procedure. For process measures, statistical testing took into account the non-independent nature of the data. The numbers available for each pairwise comparison are noted, because not all patients received all three alternatives. For categorical variables the Cochran Q test and McNemar’s $\chi^2$ test were used. Paired $t$ tests were used for normally distributed continuous variables. To take account of the multiple significance testing, a Bonferroni correction was applied. This meant that to maintain a 5% significance level $p$ values from the three pairwise tests between the three groups were only considered significant if they were below 0.017.

For effectiveness data $\chi^2$ testing and the Kruskal-Wallis equality of populations rank test were used. A two sided 5% significance level was used throughout, and 95% confidence intervals (CI) are presented where appropriate.

RESULTS

Recruitment and patient progression

Recruitment of patients took place between December 1999 and November 2000. Figure 2 summarises the flow of patients through the clinical effectiveness trial. A total of 713 patients met the inclusion criteria, of whom 103 were excluded and 10 declined to participate. The “other” exclusion was an 8 year old girl with a penetrating injury to the perineum, who was not considered an appropriate candidate for telemedicine. In 21 recruited patients (11 of whom were subsequently randomised to follow the telemedicine treatment plan) no telemedicine consultation took place because the main hospital was too busy with seriously ill patients to respond within two hours: this meant that for the primary outcome of safety, telemedicine could be compared with onsite specialist assessment in 579 patients, and compared with onsite GP assessment in 165 patients.

In the effectiveness trial, of the 600 patients recruited, 200 were initially randomised to receive the telemedicine treatment plan, 201 the onsite specialist plan and 199 the onsite GP plan. However, 135 patients randomised to the GP plan
had not seen a GP, because no GP was immediately available to assess the patient. Therefore secondary randomisation assigned 74 of these patients to telemedicine and 61 to the onsite specialist plan. This meant that 274 patients were allocated to telemedicine, 262 to the onsite specialist plan, and 64 to the GP treatment plan.

The mean age of recruited patients was 34 years with a range of 9 months to 92 years. A total of 475 patients (79.2%) were registered with a local GP. Table 1 shows the characteristics of all recruited patients by the treatment plan to which they were randomised.

**Safety**

There were 73 cases where a study diagnosis or radiological interpretation, or both, differed from the gold standard. Expert panel assessment of these gave:

- 12 discrepancies with a mean score greater than +1 (“over-treatments”)
- 35 discrepancies with a mean score between +1 and 0
- 15 discrepancies with a mean score between 0 and −1
- 11 discrepancies with a mean score less than −1 (“under treatments”)

The 50 discrepancies with a mean score between +1 and −1 were excluded from further analysis as these were minor, and attributed to variations between clinicians rather than errors likely to have a significant effect on patient outcome.

The 12 “over-treated” and 11 “under-treated” cases are detailed further in tables 2 and 3. The significant undertreatment rate ranged from 0% (95% CI 0% to 1.7%) for general practitioners to 0.5% (95% CI 0.1% to 1.5%) for the onsite specialist, and 1.4% (95% CI 0.6% to 2.7%) for telemedicine. The mean delay between injury and presentation at the peripheral hospital was 1.3 days (95% CI 1.2 to 1.5 days). The mean wait to see a nurse at the peripheral hospital was 4.2 minutes (95% CI 3.4 to 5.0 minutes).

Table 4 shows the action that would have been taken by the assessing nurse, if the patient had not been recruited into the trial.

For all telemedicine consultations, the mean delay between a request for telemedicine being made and actual start of the teleconsultation was 7.1 minutes (95% CI 6.3 to 7.9 minutes). Technical failure occurred in two cases (0.3%).

The mean duration of all consultation types was as follows: telemedicine 6.0 minutes (95% CI 5.7 to 6.2 minutes), onsite specialist 3.1 minutes (95% CI 2.9 to 3.3 minutes), and onsite general practitioner 3.4 minutes (95% CI 3.2 to 3.6 minutes). Paired t testing showed no difference between the onsite specialist and GP consultations (p = 0.64, n = 177), but a significant difference between the time taken to perform a telemedicine consultation and either an onsite specialist (n = 578) or GP consultation (n = 171) (p < 0.0001 in both cases). The average time taken to conduct a telemedicine consultation reduced as the study progressed, with a mean of 7.3 minutes (95% CI 6.2 to 8.4 minutes) for the first 100 patients and 5.0 minutes (95% CI 4.7 to 5.4 minutes) for the last 100. However, telemedicine consultations remained significantly (p < 0.0001) longer than either the onsite specialist or the GP for the last 100 patients.

For each consultation type the number of patients who had radiographs and asked to attend for a further follow up appointment is shown in table 5. Overall there were no significant differences in the proportion who had radiographs by the three alternatives (p = 0.11, n = 171, Cochran Q test). There were, however, significant overall differences between the proportion followed up by the three alternatives (p < 0.0001, McNemar’s χ² test). The proportion of patients asked to attend for follow up after a telemedicine consultation than after an onsite specialist consultation (p < 0.0001, n = 578, McNemar’s χ² test). The proportion of patients asked to attend for follow up after a telemedicine consultation

**Box 2 Expert panel discrepancy assessment scale**

1. Over (excessive) treatment leading to major inconvenience (unnecessary treatment restrictions preventing the performance of a normal occupation for greater than seven days, or a less severe restriction for longer)
2. Over (excessive) treatment leading to moderate inconvenience (unnecessary treatment restrictions preventing the performance of a normal occupation for three to seven days, or a less severe restriction for longer)
3. Over (excessive) treatment leading to mild inconvenience (unnecessary treatment restrictions preventing the performance of a normal occupation for one or two days, or an unnecessary visit to the main hospital)
0: No significant difference between the gold standard and trial treatment
−1: Under (inadequate) treatment with mild consequences (likely to cause avoidable symptoms for less than one week)
−2: Under (inadequate) treatment with moderate consequences (likely to cause avoidable symptoms for between one and six weeks)
−3: Under (inadequate) treatment with severe consequences (likely to cause avoidable symptoms for greater than six weeks)

**Box 3 Clinical effectiveness outcomes derived from the review questionnaires**

- Whether the problem had resolved.
- Whether the patient had been able to return to normal activity.
- Whether the patient had taken time off work (where applicable).
- Whether there had been further health services contact regarding the same condition.
- Whether the patient had sought further unscheduled care.
- Whether there had been any change in treatment since the patient entered the trial.

consultations received a score worse than −2 it is possible to state that for minor injuries telemedicine the risk of error leading to severe consequences has a 95% confidence interval between 0% and 0.52%.

**Process measures**

The mean delay between injury and presentation at the peripheral hospital was 1.3 days (95% CI 1.2 to 1.5 days). The mean wait to see a nurse at the peripheral hospital was 4.2 minutes (95% CI 3.4 to 5.0 minutes).

Table 4 shows the action that would have been taken by the assessing nurse, if the patient had not been recruited into the trial.
Clinical effectiveness

Eight patients (1.3%) were lost from the study. The mean time between recruitment and review was 7.0 days (SD 1.4 days, range 3 to 14 days). There was no significant difference in the time to review between the three randomisation groups (p = 0.82, Kruskal-Wallis test).

Table 6 summarises the main measures of clinical effectiveness. No significant differences were detected for any of these outcomes.

Table 1: Baseline characteristics of participants by randomisation of treatment plan

<table>
<thead>
<tr>
<th>Variable</th>
<th>Telemedicine (n = 274)</th>
<th>Onsite specialist (n = 262)</th>
<th>General practitioner (n = 64)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>111 (40.5)</td>
<td>99 (37.8)</td>
<td>20 (31.2)</td>
</tr>
<tr>
<td>Male</td>
<td>163 (59.5)</td>
<td>163 (62.2)</td>
<td>44 (68.8)</td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;16</td>
<td>60 (21.9)</td>
<td>56 (21.4)</td>
<td>18 (28.1)</td>
</tr>
<tr>
<td>16–24</td>
<td>44 (16.1)</td>
<td>53 (20.2)</td>
<td>15 (23.4)</td>
</tr>
<tr>
<td>25–34</td>
<td>52 (19.0)</td>
<td>41 (15.7)</td>
<td>9 (14.1)</td>
</tr>
<tr>
<td>35–44</td>
<td>37 (13.5)</td>
<td>39 (14.9)</td>
<td>5 (7.8)</td>
</tr>
<tr>
<td>45–54</td>
<td>31 (11.3)</td>
<td>29 (11.1)</td>
<td>6 (9.4)</td>
</tr>
<tr>
<td>55–64</td>
<td>21 (7.7)</td>
<td>23 (8.8)</td>
<td>6 (9.4)</td>
</tr>
<tr>
<td>65+</td>
<td>29 (10.6)</td>
<td>21 (8.0)</td>
<td>5 (7.8)</td>
</tr>
<tr>
<td>In employment/full time education</td>
<td>210 (77.8)*</td>
<td>205 (79.2)†</td>
<td>51 (81)‡</td>
</tr>
</tbody>
</table>

* n = 270; † n = 259; ‡ n = 63. Percentages are shown in parentheses.
DISCUSSION

These results show that the safety of minor injuries telemedicine is similar to conventional practice. It is inevitable that errors will occur during any large series of consultations, and this research quantified and compared these errors for the three alternatives studied, finding no significant differences between them.

Only one randomised trial of telemedicine in a similar context has previously been published. This was a much smaller study, conducted in North America with just 104 participants, but showed similar results.

Over-treatments, radiographs, and follow up

We studied the over-treatment rate because this might be increased for telemedicine, particularly if medical staff were unsure of the diagnosis, but found no evidence that this was the case.

While the radiography rate was nearly identical for telemedicine and onsite specialist consultation (59.2% compared with 60.5%), a follow up appointment was arranged for a significantly greater proportion of telemedicine patients (35.8% compared with 27.5%; p<0.0001 using McNemar’s χ² test), perhaps reflecting a lack of confidence in telemedicine. This did not reduce during the course of the study. The highest rate of planned follow up was seen in GP consultations (65%). This may reflect the way in which GPs normally practice, as we observed that follow up appointments arranged by a GP were frequently an option that the patient could pursue only if problems persisted. This is supported by the finding that, at the seven day study review, there were no significant differences between the randomisation groups in planned or unplanned use of healthcare services.

Under-treatments

Under-treatments are of greater importance as they may lead to long term effects and possible litigation. Several studies have attempted to quantify the incidence of error in emergency medicine, often concentrating on radiological interpretation. In the United Kingdom, a recent study quoted an error rate of 1.5%, falling to the more acceptable figure of 0.7% when a picture archiving and communications system (PACS) was introduced. In emergency care it has been necessary to establish a risk below which a patient can be reasonably discharged, and this threshold is commonly set at 1%. In this study only one consultation, by the onsite specialist, received a mean under-treatment score worse than −2, and for telemedicine the risk of error leading to severe consequences was less than 1% (95% CI 0% to 0.52%).

Process measures

Patients attending the peripheral hospital were usually seen by a nurse very promptly, and had to wait a mean of only 7.1 minutes for a telemedicine consultation, excepting those cases where the main hospital was too busy to respond. This suggests that patients attending the peripheral site were likely to be assessed by a senior doctor much more rapidly than if they had attended the main hospital in person. Indeed, medical staff commented that there was a possibility

<table>
<thead>
<tr>
<th>Gold standard</th>
<th>Error made</th>
<th>Mean error score</th>
<th>Error source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conjunctival laceration</td>
<td>Laceration missed</td>
<td>−2.5</td>
<td>Onsite specialist</td>
</tr>
<tr>
<td>Undisplaced fracture of the distal fibula</td>
<td>Fracture missed</td>
<td>−1.8</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Compound finger fracture</td>
<td>Fracture missed</td>
<td>−1.6</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Greenstick fracture of the distal radius</td>
<td>Fracture missed</td>
<td>−1.6</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Infected hand laceration</td>
<td>Infection missed</td>
<td>−1.6</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Infected shoulder abrasion</td>
<td>Infection missed</td>
<td>−1.4</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Fractured clavicle</td>
<td>Fracture missed</td>
<td>−1.2</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Cellulitis</td>
<td>Infection missed</td>
<td>−1.2</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Fractured clavicle</td>
<td>Fracture misdiagnosed</td>
<td>−1.2</td>
<td>Onsite specialist</td>
</tr>
<tr>
<td>Avulsion fracture from the tip of the olecranon</td>
<td>Fracture missed</td>
<td>−1.2</td>
<td>Telemedicine</td>
</tr>
<tr>
<td>Knee laceration with an underlying patella fracture</td>
<td>Fracture missed</td>
<td>−1.1</td>
<td>Onsite specialist</td>
</tr>
</tbody>
</table>
that they would leave a patient with more urgent problems to respond to a request for minor injuries telemedicine.

If telemedicine acted to preferentially bypass the queue in the main emergency department, patients might choose to travel to the peripheral hospital for treatment. One approach to this problem would be to “triage” telemedicine cases as if they had come to the main site in person. This would cause practical difficulties at the peripheral site, and might also prove unacceptable to patients who are used to the prompt service that the peripheral hospital usually provides.

Only 17 (2.8%) of patients would have been referred directly to the main hospital if they had not been recruited into the trial, so the potential for telemedicine to reduce unnecessary patient transfers in this context is very limited. The nurse’s assessment of what would have happened had the patient not been recruited could have been biased by the knowledge that the patient was about to enter the trial, but this transfer rate of 2.8% is very similar to the 3% which existed before the start of the study. The low transfer rate is attributable to the effective service already provided, and the fact that most of the injuries were minor. If the telemedicine plan had been followed for these 17 patients then eight would still have been transferred, with the remaining nine treated locally.

Telemedicine weaknesses

In 16 cases (2.8%) the doctor providing a telemedical opinion felt that telemedicine was inadequate in some way. This related either to problems in visualising and interpreting a radiograph (particularly chest radiographs) or to difficulties in assessing a wound (for example, burn depth). Nevertheless, no significant discrepancy resulted in any of these cases, suggesting that the problem was recognised and appropriately resolved.

Problems with telemedicine are often attributed to technical deficiencies that can be overcome by improving the performance of equipment.22 In this study, however, we were conforming (with the exception of using a television rather than a computer monitor) to a technical standard that achieves satisfactory results in a laboratory setting.23 This suggests that further improvements in technology may not provide complete solutions, and that some difficulties relate to the medium itself.

Study weaknesses

We assumed that treatment discrepancies were attributable to telemedicine rather than variations in clinician performance. As most participants had comparatively straightforward injuries, and the doctors involved were of comparable experience, this assumption appears reasonable. Furthermore, the use of a large expert panel and removal of discrepancies scoring between +1 and −1 should have prevented individual treatment variation from influencing the results.

It was not possible to achieve consecutive recruitment because of the requirement for the onsite specialist to be available. Nevertheless, minor injuries rarely presented overnight, when recruitment would have been inefficient, and there is no reason to suspect that our sample is not representative of the wider population of minor injuries patients.

The methodology of this study was unusual in that rather than being randomised to receive only one type of consultation, we elected to have each recruited patient undergo more than one consultation type, with strict blinding between the consultations. This allowed us to collect a large sample in which telemedicine and conventional consultation could be directly compared with high sensitivity. A disadvantage was that the sample size was not specifically calculated for the randomised trial of clinical effectiveness. Therefore the clinical outcomes, although supportive of our safety data, should be viewed with caution, particularly as only a comparatively small number of patients followed a GP treatment plan.

Each patient underwent two or three consecutive consultations, and the onsite assessment always preceded telemedicine. This could lead to an order effect, with refinement and shortening of the history given by the patient. If such an effect existed, however, it would tend to reduce the length of telemedicine consultations, in contrast with the results obtained. The GP consultations might also have been shortened by this effect, although some were performed before and some after telemedicine. The doctors involved were also careful to avoid

### Table 4

<table>
<thead>
<tr>
<th>Action</th>
<th>Number (n=600)</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Telephone general practitioner and ask them to come to the peripheral emergency department to see patient</td>
<td>60</td>
<td>10.0</td>
</tr>
<tr>
<td>Arrange appointment for patient with a general practitioner or on a following day</td>
<td>50</td>
<td>8.3</td>
</tr>
<tr>
<td>Ring general practitioner for advice</td>
<td>282</td>
<td>47.0</td>
</tr>
<tr>
<td>Ask patient to wait until a general practitioner comes to the peripheral emergency department</td>
<td>147</td>
<td>24.5</td>
</tr>
<tr>
<td>Arrange for patient to be seen at the peripheral emergency department specialist clinic</td>
<td>11</td>
<td>1.8</td>
</tr>
<tr>
<td>Refer directly to the main emergency department</td>
<td>14</td>
<td>2.3</td>
</tr>
<tr>
<td>Refer to an inpatient team at the main hospital</td>
<td>3</td>
<td>0.5</td>
</tr>
<tr>
<td>Ring general practitioner for advice</td>
<td>31</td>
<td>5.2</td>
</tr>
<tr>
<td>Ring the main emergency department for advice</td>
<td>2</td>
<td>0.3</td>
</tr>
<tr>
<td>Total</td>
<td>600</td>
<td>100.0</td>
</tr>
</tbody>
</table>

### Table 5

Comparison of the radiography and follow up rates for each type of consultation

<table>
<thead>
<tr>
<th>Consultation type</th>
<th>Number of patients radiographed</th>
<th>Number of patients asked to attend for follow up</th>
</tr>
</thead>
<tbody>
<tr>
<td>Telemicine (n=579)</td>
<td>343 (59.2) (55.1% to 63.3%)</td>
<td>207 [35.8] (31.8% to 39.8%)</td>
</tr>
<tr>
<td>Onsite specialist (n=600)</td>
<td>363 [60.5] (56.5% to 64.4%)</td>
<td>165 [27.5] (24.0% to 31.3%)</td>
</tr>
<tr>
<td>General practitioner (n=177)</td>
<td>94 [53.1] (45.5% to 60.6%)</td>
<td>115 [65.0] (57.5% to 72.0%)</td>
</tr>
</tbody>
</table>

Data shown as numbers, percentages, and 95% CI.
giving information to the patient during each consultation, minimising the effect of consecutive consultations.

**Application to practice**

We have shown that telemedicine is safe and effective in minor injuries practice. However, there is very little evidence regarding the effectiveness of telemedicine in other aspects of emergency medicine, such as acute illness or more severe injuries.

If telemedicine is used to support nursing staff at a high consultation volume can be expected, particularly if it replaces an existing medical service. This is time consuming for the hospital providing the consultations, and also runs the risk that the transfer rate will be increased because of system failures or diagnostic doubt. Direct referral to the main hospital was a comparatively rare event (2.8%), but where it occurred transfer was avoided in 53% of cases. Therefore, telemedicine used to support doctors or as a supplement to an existing service may approximately halve the patient transfer rate, and is in keeping with the one previous paper that studied telemedicine to support remote general practitioners, in which transfer was avoided in 58% of cases.11

While a comparatively low cost system is capable of delivering satisfactory clinical results in terms of safety and patient outcome, there are a number of potential problems that may be overlooked by enthusiasts. Many of these problems will not be influenced by further technological advances, and must be tackled in other ways: some are inherent to telemedicine itself, but many are related to service delivery and organisation. It is important that decision makers are not overly distracted by technical concerns but focus on the managerial and economic issues that underpin telemedicine as a change in the way that care is delivered.

**Conclusion**

In the management of minor injuries, telemedicine is capable of providing a satisfactory standard of care. Significant errors are rare and similar in frequency to conventional practice and accepted risks. There is no evidence that telemedicine is superior to the routine practice of GP consultation. Telemedicine requires about twice as long as an onsite consultation, in addition to the time entailed in responding to a request and breaking away from other activities. While telemedicine may be effective for minor injuries work, emergency medicine encompasses a variety of other conditions that may be less amenable to this technology, and where research is still required.

**ACKNOWLEDGEMENTS**

Our thanks go to the patients and staff of Tewkesbury and Cheltenham Hospitals and all the general practitioners who participated in this study, and without whom it would not have been possible. Also our grateful acknowledgements go to Professor Gordon Wilcock for his academic supervision, Professor Richard Wootton for his advice and support, Mr Christopher Foy and Professor Tim Peters for their statistical advice, and Mr Iain Grant, Mr Paul Howarth, Dr Tom Llewellyn, Mr Christian Oakland, Mr Sapal Tachakra, and Mr Paul Wilson for their assistance in the development of this study.

**Table 6** Summary of the main measures of clinical effectiveness

<table>
<thead>
<tr>
<th>Outcome at review</th>
<th>Telemedicine (n = 270)</th>
<th>Onsite specialist (n = 259)</th>
<th>General practitioner (n = 63)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Completely better</td>
<td>67 (24.8) (19.8% to 30.4%)</td>
<td>61 (23.6) (18.5% to 29.2%)</td>
<td>10 (15.9) (7.9% to 27.3%)</td>
<td>0.32</td>
</tr>
<tr>
<td>Returned to normal activity</td>
<td>127 (47.0) (41.0% to 53.2%)</td>
<td>106 (40.9) (34.9% to 47.2%)</td>
<td>30 (47.6) (34.9% to 60.6%)</td>
<td>0.32</td>
</tr>
<tr>
<td>Taken time off work</td>
<td>91 (33.7) (28.1% to 39.7%)</td>
<td>95 (36.7) (30.8% to 42.9%)</td>
<td>18 (28.6) (17.9% to 41.3%)</td>
<td>0.45</td>
</tr>
<tr>
<td>Seen in an emergency department</td>
<td>89 (33.0) (27.4% to 38.9%)</td>
<td>74 (28.6) (23.2% to 34.5%)</td>
<td>25 (39.7) (27.6% to 52.8%)</td>
<td>0.20</td>
</tr>
<tr>
<td>Seen in an outpatient clinic</td>
<td>37 (13.7) (9.8% to 18.4%)</td>
<td>41 (15.8) (11.6% to 20.9%)</td>
<td>3 (4.8) (1.0% to 13.3%)</td>
<td>0.07</td>
</tr>
<tr>
<td>Seen in a general practitioner’s surgery or at home</td>
<td>6 (2.2) (0.8% to 4.8%)</td>
<td>10 (3.9) (1.9% to 7.0%)</td>
<td>2 (3.2) (0.4% to 11.0%)</td>
<td>0.55</td>
</tr>
<tr>
<td>Sought further unscheduled care</td>
<td>14 (5.2) (2.9% to 8.5%)</td>
<td>15 (5.8) (3.3% to 9.4%)</td>
<td>4 (6.3) (1.8% to 15.5%)</td>
<td>0.91</td>
</tr>
<tr>
<td>Required a change in treatment</td>
<td>26 (9.6) (6.4% to 13.8%)</td>
<td>17 (6.6) (3.9% to 10.5%)</td>
<td>4 (6.3) (1.8% to 15.5%)</td>
<td>0.37</td>
</tr>
</tbody>
</table>

Data shown as numbers, percentages, and 95% CI.

**REFERENCES**


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