Discrepancies in drug histories at admission to gastrointestinal surgery, internal medicine and geriatric hospital wards in Central Norway: a cross-sectional study

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ABSTRACT

Objectives To compare discrepancies in drug histories among patients acutely admitted to different hospital wards, classify the discrepancies according to their potential clinical impact and identify appropriate selection criteria for patients that should be subject to a detailed drug history at admission.

Design Cross-sectional study.

Setting Two gastrointestinal surgery wards and one geriatric ward at St Olav’s University Hospital in Trondheim and two general internal medicine wards at Ålesund Hospital in Ålesund, Norway.

Participants All patients acutely admitted to these wards during a period of three months were asked to participate in the study. A total of 168 patients were included. For each patient, drug information available at admission was compared with information from drug lists obtained from the general practitioner and (if applicable) the home care services/the nursing home.

Primary and secondary outcome measures Number of patients with one or more discrepancies in their drug history. Type and clinical impact of the discrepancies found. Selection criteria for patients that should be subject to a detailed drug history.

Results In total, 83% had at least one discrepancy in their drug history. Omission of a drug accounted for 72% of the discrepancies, whereas a difference in dosing was the cause of the remaining 28%. 9% of the discrepancies had the potential to cause severe harm or discomfort. We found no significant differences in the number of discrepancies between hospital wards, genders, ages or levels of care.

Conclusions This study demonstrates the importance of collecting drug information from all available sources when a patient is admitted to hospital. As we found no significant differences in discrepancies between subgroups of patients, we suggest that medication reconciliation should be performed for all patients.

INTRODUCTION

An accurate drug history is an essential part of patient assessment at admission to hospital.
authorities and patient safety organisations in several countries.8 9

An American study from 2008 has shown that pharmacists appear to be better suited and more effective than physicians in obtaining drug histories,10 and a Belgian study published in 2010 demonstrates that pharmacists are especially suited to acquire and supervise accurate medication histories.11 Pharmacists in many countries have developed standardised procedures for obtaining complete drug histories, including systematic medication reconciliation.3 6 7 11 12

Various factors have been associated with an increased risk of discrepancies in the drug history at admission to hospital including age, level of healthcare, number of drugs and diagnoses.3 6 13 At present, results from studies aiming to identify factors predicting errors or discrepancies in drug histories are inconclusive and contradictory.2 3 6 7 13-15 Owing to the limited resources usually available for performing quality controls of drug histories, developing appropriate selection criteria is of crucial importance.

Most previous studies in the area have included elderly patients in internal medicine wards in a single hospital.2 3 6 7 15 16 The results from these studies might not be representative for other patient populations. One study by Unroe et al from 2010 showed a significant difference in the proportion of patients with discrepancies on admission to hospital between patients admitted to cardiology service, general medicine and general surgery, 15%, 22% and 35%, respectively.12

The aim of this study was to identify and compare discrepancies in drug histories among patients acutely admitted to various types of hospital wards, classify the discrepancies according to their potential clinical impact and identify appropriate selection criteria for patients that should be subject to detailed medication reconciliation.

**METHODS**

The study was conducted as a cross-sectional study in five different hospital wards—two gastrointestinal surgery wards at St Olav’s University Hospital in Trondheim, Norway, two general internal medicine wards at Ålesund Hospital in Ålesund, Norway and one geriatric ward at St Olav’s University Hospital. St Olav’s is a large university hospital in the city of Trondheim, Central Norway, with 983 beds of which 48 are located in the gastrointestinal surgery wards and 15 in the geriatric ward. Ålesund Hospital is a secondary hospital covering the northern part of western Norway with approximately 270 beds, of which 48 are located in the two internal medicine wards. All patients acutely admitted to these wards during a period of 3 months were asked to participate in the study. The patients were informed both orally and in writing about the study and those giving their written informed consent were included in the study. The Regional Ethics Committee of Central Norway considered the study to be a quality-control study, and as such, the committee stated that no further approval was necessary.
In Norway, there are several patient record systems both within primary care and within secondary care. These systems do not communicate with each other, and health professionals from primary and secondary care or even from different hospitals do not have access to each other’s patient records. Therefore, the patients’ medication lists were collected manually, and medication reconciliation was performed by a clinical pharmacist after admission to the wards. The patients’ drug lists, as obtained by the doctor in charge at admission to hospital, were collected from the hospital records taken at admission. These lists are seen as the main source of information on which the doctor bases his prescribing in the drug chart. Information from the drug chart was not collected in this study. As soon as practically feasible after admission, the following efforts were taken to obtain supplementary drug information:

For patients living in their own homes, the patients’ general practitioners (GP) were contacted by telephone to provide the latest medication records.

For patients living in their own homes receiving home care services and patients living in nursing homes, we contacted, in addition to the GP, also the caregivers by telephone, asking them to provide a copy of the patients’ drug list. Drug information from the home care services and the nursing homes was classified together, since the patients in both cases did not handle their own drugs, but received their drugs from a nurse.

Age, gender, level of care before admission and data from all available drug lists were registered for each patient. Trade name, generic name, administration form, strength and dosing were registered for all drugs. Over-the-counter drugs were only registered if they were prescribed by a doctor.

Finally, the drugs were classified according to the Anatomical Therapeutic Chemical system.17

For each patient, drug lists from the GP and/or the home care services/the nursing home were compared with the drug information available at admission. Discrepancies revealed were linked to the drug involved and registered in a separate form. Since the hospital record list could be compared with information both from the GP and from the home care services or the nursing home, there could be up to two discrepancies per drug. None of the patients had information both from the home care services and from the nursing home. Discrepancies were either classified as ‘omission of drug’ (ie, the drug was lacking in one of the lists) or ‘difference in dosing’ (ie, differences with regard to administration form, strength or dosage).

An expert panel consisting of four persons (clinical pharmacist, geriatrician, physician from a specific ward and clinical pharmacist) rated the discrepancies for their potential clinical impact. Assessment of the discrepancies was performed using a previously published method where the discrepancies were rated in three classes according to whether they had potential to cause minimal, moderate or severe discomfort or harm to the patient.2 This rating scale developed by Cornish et al, has also been used in several other studies.5 18–21

In addition, a fourth class, denoted non-classifiable, was added. When evaluating the effect of each discrepancy, the expert panel took into account that the error was carried forward for an average hospital stay in the same department, which was 2–3 days for gastrointestinal surgery and internal medicine wards and 4–7 days for the geriatric ward. The evaluation also took the patients’ diagnoses and clinical status, as described in the electronic patient record into account. Each member of the panel first rated the discrepancies alone before the cases were discussed in a meeting. Disagreements between the panel members were solved by discussion, and consensus was reached in all cases. The expert group started its work by evaluation of 10 pilot cases collected in the same way as in the main study. This was done to give the expert panel a common understanding of the classification system and the working model.

Statistical comparisons of number of drugs and total number of discrepancies between different groups of patients were performed using Mann-Whitney U test. Comparisons of number of discrepancies from different sources of information were performed using an independent sample Kruskal-Wallis test (SPSS V.19; SPSS). p Values <0.05 were considered statistically significant.

RESULTS

A total of 168 patients were included in the study, 56 from the gastrointestinal surgery wards, 51 from the general internal medicine wards and 61 from the geriatric ward. According to information available at admission, there were a total of 901 prescriptions, that is, a mean of 5.4 (range 0–19) prescriptions per patient. However, when information from all sources was combined, the total number of prescriptions was 1176, that is, a mean of 7.0 (range 0–24) prescriptions per patient. Twelve patients (7%) did not use any drugs at all, and the mean number of prescriptions was significantly lower than the mean number of 7.7 drugs (range 0–24; p=0.956).

Patients below the age of 80 years had a mean number of 6.5 drugs prescribed (range 0–24) which was significantly lower than the mean number of 7.7 drugs (range 0–19) prescribed to patients 80 years or older (p=0.034). Patients who administered their own drugs had a mean number of 5.9 drugs prescribed (range 0–17), which was significantly lower than the mean number of 8.9 drugs (range 0–24) prescribed to patients with help from the home care services or a nursing home (p<0.001).

Of the 168 patients in the study, 139 (83%) had at least one discrepancy when comparing the information
available at admission with the other available sources of information. In the gastrointestinal surgery wards, the number of patients with discrepancies was 43 (77%); in the internal medicine wards, the number was 47 (92%) and in the geriatric ward, the number was 49 (80%). Among these patients, there was a range from 1 to 32 discrepancies.

The number of discrepancies when comparing the drug information available at admission to the different sources of information is presented in Table 2.

In total there were 697 discrepancies among all patients. Difference in dosing counted for 195 discrepancies (28%), whereas omission of drug was the cause of the remaining 502 discrepancies (72%). Of the omitted drugs, 404 (80%) were missing in the drug list recorded at admission. In the remaining 98 cases (20%), it had been stated in the hospital record that

**Table 2** Characteristics of the 168 patients included in the study and number of drugs prescribed according to the various sources of information

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Gastrointestinal surgery</th>
<th>Internal medicine</th>
<th>Geriatrics</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients</td>
<td>56</td>
<td>51</td>
<td>61</td>
<td>168</td>
</tr>
<tr>
<td>Mean age, years (range)</td>
<td>62 (22–91)</td>
<td>78 (45–92)</td>
<td>83 (65–96)</td>
<td>74 (22–96)</td>
</tr>
<tr>
<td>Female gender, n (%)</td>
<td>28 (50.0)</td>
<td>29 (56.9)</td>
<td>44 (72.1)</td>
<td>101 (60.1)</td>
</tr>
<tr>
<td>Living in their own home without home care services, n (%)*</td>
<td>47 (83.9)</td>
<td>32 (62.7)</td>
<td>22 (36.1)</td>
<td>101 (60.1)</td>
</tr>
<tr>
<td>Living in their own home with home care services, n (%)*</td>
<td>7 (12.5)</td>
<td>15 (29.4)</td>
<td>31 (50.8)</td>
<td>53 (31.5)</td>
</tr>
<tr>
<td>Living in a nursing home, n (%)*</td>
<td>2 (3.6)</td>
<td>4 (7.8)</td>
<td>8 (13.1)</td>
<td>14 (8.3)</td>
</tr>
<tr>
<td>Number of patients with no drugs according to all sources, n (%)</td>
<td>9 (16.1)</td>
<td>0 (0)</td>
<td>3 (4.9)</td>
<td>12 (7.1)</td>
</tr>
<tr>
<td>Mean number of drugs per patient (range) according to information from the various sources†</td>
<td>Information available at admission to hospital</td>
<td>3.1 (0–11)</td>
<td>8.7 (3–19)</td>
<td>4.7 (0–17)</td>
</tr>
<tr>
<td>Information from the general practitioner‡</td>
<td>3.9 (0–12)</td>
<td>8.4 (3–16)</td>
<td>4.6 (0–15)</td>
<td>5.2 (0–16)</td>
</tr>
<tr>
<td>Information from the home care services/nursing home§</td>
<td>8.5 (5–14)</td>
<td>8.5 (4–16)</td>
<td>6.0 (0–15)</td>
<td>7.2 (0–16)</td>
</tr>
<tr>
<td>All available sources combined</td>
<td>4.8 (0–14)</td>
<td>10.4 (5–24)</td>
<td>6.2 (0–20)</td>
<td>7.0 (0–24)</td>
</tr>
</tbody>
</table>

*Percentages may not total 100 due to rounding.
†All patients had information from at least one supplementary source (ie, general practitioner and/or home care services/nursing home).
‡Information available for 54 gastrointestinal surgery patients, 49 internal medicine patients and 53 geriatric patients.
§Information available for 7 gastrointestinal surgery patients, 19 internal medicine patients and 31 geriatric patients.
the patient used a drug at admission, but this drug was not listed in any other source of information.

The distribution of patients according to their total number of discrepancies for each of the three departments is shown in figure 1.

Among the 17 drugs prescribed for 20 patients or more, paracetamol and lactulose had the highest percentage of prescriptions with discrepancies, with 74% and 70%, respectively. The percentage of discrepancies related to all the prescriptions of the same drug is presented in figure 2.

There were no significant differences in the number of discrepancies between male and female patients (mean 4.1; range 0–32 vs mean 4.2; range 0–22; p=0.772) or between patients below and above the age of 80 years (mean 3.7; range 0–32 vs mean 4.7; range 0–22; p=0.056). Patients who administered their own drugs had a mean number of 2.8 discrepancies (range 0–11), as compared with 6.2 discrepancies (range 0–32) in patients whose drugs were administered by the home care services or lived in a nursing home (p<0.001).

When correcting for the number of drugs prescribed by calculating number of discrepancies per prescribed drug (the number of discrepancies in a patient divided by the number of drugs prescribed to the same patient), there was no statistically significant difference between any of the groups compared. Male and female patients had means of 0.50 and 0.49, respectively (p=0.502), whereas patients below and above the age of 80 years had means of 0.53 and 0.48, respectively (p=0.135). Patients who were prescribed less than five drugs had a mean of 0.53, as compared with 0.48 in patients that were prescribed five drugs or more (p=0.465). Patients who administered their own drugs had a mean of 0.47, as compared with 0.54 in patients that had their drugs administered by the home care services or lived in a nursing home (p=0.235).

More than half of the discrepancies (381/697; 55%) were classified as having the potential to cause only minimal harm or discomfort to the patients. In addition, 33% (231/697) had the potential to cause moderate harm or discomfort, 9% (62/697) had the potential to cause severe harm or discomfort, whereas 3% (23/697) were considered non-classifiable. The number of discrepancies classified according to their potential to cause harm or discomfort to the patient within each group is presented in table 3.

Among the 168 patients included in the study, 28 (17%) had a total of 62 discrepancies with a potential to cause severe harm or discomfort (class 3), with a range from 1 to 13 severe discrepancies per patient. Omission of drug counted for 58 of the severe discrepancies, whereas difference in dosing was the cause of the remaining 4.

Thirty-two different drugs were involved in these severe discrepancies. The drugs/drug groups most often implicated were antithrombotic agents (n=8), insulin (n=7), corticosteroids for systemic use (n=6), oral blood glucose lowering drugs (n=6) and beta blockers (n=5).

Table 4 presents some representative case histories with one or more discrepancies classified as severe.

### Table 2

<table>
<thead>
<tr>
<th>Source of Information</th>
<th>Gastrointestinal surgery</th>
<th>Internal medicine</th>
<th>Geriatrics</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Information from the general practitioner†</td>
<td>3.0 (0–11)</td>
<td>4.1 (0–18)</td>
<td>2.6 (0–17)</td>
<td>3.2 (0–18)</td>
</tr>
<tr>
<td>Information from the home care services/nursing home‡</td>
<td>4.0 (2–8)</td>
<td>4.0 (0–16)</td>
<td>2.9 (0–7)</td>
<td>3.4 (0–16)</td>
</tr>
<tr>
<td>All available sources combined§</td>
<td>3.4 (0–15)</td>
<td>5.4 (0–32)</td>
<td>3.7 (0–24)</td>
<td>4.1 (0–32)</td>
</tr>
</tbody>
</table>

*All patients had information from at least one supplementary source (ie, general practitioner and/or home care services/nursing home).
†Information available for 54 gastrointestinal surgery patients, 49 internal medicine patients and 53 geriatric patients.
‡Information available for 7 gastrointestinal surgery patients, 19 internal medicine patients and 31 geriatric patients.
§Up to two discrepancies per drug.

### Table 3

<table>
<thead>
<tr>
<th>Discrepancies</th>
<th>Gastrointestinal surgery (n=56)</th>
<th>Internal medicine (n=51)</th>
<th>Geriatrics (n=61)</th>
<th>Total (n=168)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N</td>
<td>N per patient</td>
<td>N</td>
<td>N per patient</td>
</tr>
<tr>
<td>Class 1</td>
<td>132</td>
<td>2.4</td>
<td>113</td>
<td>2.2</td>
</tr>
<tr>
<td>Class 2</td>
<td>43</td>
<td>0.8</td>
<td>119</td>
<td>2.3</td>
</tr>
<tr>
<td>Class 3</td>
<td>7</td>
<td>0.1</td>
<td>44</td>
<td>0.9</td>
</tr>
<tr>
<td>Class 0</td>
<td>10</td>
<td>0.2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>192</td>
<td>3.4</td>
<td>276</td>
<td>5.4</td>
</tr>
</tbody>
</table>

Class 1: potential to cause minimal harm or discomfort to the patient, class 2: potential to cause moderate harm or discomfort to the patient, class 3: potential to cause severe harm or discomfort to the patient, class 0: non-classifiable.
Table 4  Examples of patients with discrepancies considered having the potential to cause severe harm or discomfort.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Case</th>
<th>Class 3 discrepancies</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>An 82-year-old man with severe dementia, atrial flutter, diabetes mellitus type II and metastatic prostate cancer. He was living in a nursing home and was admitted to the geriatric ward because of acute delirium. The hospital record at admission did not mention any drugs or the prostate cancer diagnosis. However, according to the nursing home notes, the patient was using methylprednisolone 8mg and morphine slow release 60mg daily in addition to glipizide, paracetamol and lactulose.</td>
<td>Omission of methylprednisolone and of morphine</td>
</tr>
<tr>
<td>B</td>
<td>A 75-year-old woman with heart failure and previously myocardial infarction and stroke. She was living in a nursing home and was admitted to the department of internal medicine because of seizures. Venlafaxine and fluoxetine were listed in the hospital record at admission. However, according to the nursing home notes, she was using ramipril 5mg daily for her heart failure in addition to seven other drugs.</td>
<td>Omission of ramipril for heart failure</td>
</tr>
<tr>
<td>C</td>
<td>A 63-year-old woman with chronic obstructive pulmonary disease (COPD). She was living in her home, administering her own drugs and was admitted to the gastrointestinal surgery ward with acute abdominal pain. The hospital record at admission listed use of salbutamol, ipratropium bromide and acetylcysteine. According to the general practitioner she was currently also prescribed methylprednisolone 15mg and theophylline for her COPD in addition to antiplatelet treatment with acetylsalicylic acid.</td>
<td>Omission of methylprednisolone and of theophylline</td>
</tr>
<tr>
<td>D</td>
<td>An 86-year-old woman with hypothyroidism and several transient ischaemic attacks was admitted to the geriatric ward because of general functional deterioration, headache and dizziness. She was living in her home with help from the home care services. The hospital record at admission did not list any drugs, and the hypothyroidism was not stated. Both the general practitioner and the home care services listed dipyridamole, acetylsalicylic acid, hydrochlorothiazide, fluoxetine and levothyroxine.</td>
<td>Omission of levothyroxine</td>
</tr>
</tbody>
</table>

DISCUSSION

The principal finding in the present study is that 83% of the patients had one or more discrepancies in their drug history at admission to hospital. This number is higher than in most other studies, although it also varies considerably between previous studies. A review from 2005 found errors in 10%–67% of the cases, and newer studies from Sweden, Norway and USA show error rates of 47%, 50% and 51%, respectively. There is no indication that the higher error rates in our study are caused by stricter definitions in other studies. Most other studies compare the different sources of information to the hospital drug chart and not the drug information available at admission to hospital. To the best of our knowledge, only two previous studies has compared the drug information available at admission to other sources of drug information. We believe it is more appropriate to evaluate drug information in the admission record than in the hospital drug chart because changes might take place on the discretion of the physician in charge before the drug list is transferred from the admission record to the drug chart. On the other hand, information from the patients themselves was not included in our analysis, and this might in fact have led to a lower error rate.

Patients admitted to the gastrointestinal surgery wards were younger, had less assistance from the home care services and were less often living in nursing homes than the patients from internal medicine and geriatric wards. One could therefore have expected fewer patients with discrepancies in the gastrointestinal surgery wards than in both the internal medicine and the geriatric wards. However, this was not the case, as 77% of the patients in the gastrointestinal surgery and 80% in the geriatric wards had at least one discrepancy, compared with as much as 92% of the internal medicine patients. This indicates that type of ward (as a rough indicator of the cause of admittance) does not seem to be an important factor for predicting the risk of discrepancies in drug history of a patient.

Potential to harm or cause discomfort to the patient could have been underestimated in this study, since the effect of each discrepancy took into account that the mistake was carried forward for just an average hospital stay in the actual department, which was 2–3 days for gastrointestinal surgery and internal medicine wards and 4–7 days for the geriatric ward. We know, both from the literature and from our own unpublished data, that erroneous drug lists with consequent incorrect use of drugs often follow the patients out of the hospital and will not be corrected for months or even years. For many of the discrepancies, the potential to cause harm or discomfort to the patient will rise substantially with time. The effect of each discrepancy took into account that the potential to cause harm or discomfort to the patient will rise substantially with time. We chose this time perspective after thorough discussion among the members of the expert group to avoid overestimating the potential to cause harm or discomfort.

We found that patients with home care services or living in a nursing home had more discrepancies than patients that were handling their drugs themselves, although this difference disappeared after correction for the number of drugs prescribed. In contrast, a Swedish study from 2012 found that patients living in their own home without any care services had an increased risk of medication history
errors. These differences might be caused by the fact that the healthcare regulation and information systems in Sweden and Norway are somewhat different. In Sweden, healthcare professionals in primary and secondary care in some cases have access to each other’s electronic records, and there is one common record for the primary care sector. In Norway, there are several patient record systems both within primary care and within secondary care. These systems do not communicate, and health professionals from primary and secondary care or even from different hospitals do not have access to each other’s patient records. This non-transparency and the complicated systems for transfer of information might also lead to a higher error rate, especially for patients receiving home care services or living in a nursing home.

The number of discrepancies per drug prescribed was not significantly higher for patients using a high number of drugs than for those using fewer drugs. As opposed to the Swedish study,3 the proportion of drugs associated with a discrepancy was similar for patients with five or more drugs and for those with less than five drugs. In our study, the number of discrepancies per patient seems to increase in parallel with the number of drugs prescribed.

This study has several limitations. First, there is no ‘golden standard’ regarding the information of which drugs the patient actually has been taking. In fact, none of the lists we have consulted are necessarily ‘correct’ (depending on the definition of the word ‘correct’—is it the drugs the patient has taken or the drugs the patient should have been taking). Although we do not know in detail which drugs and dosages the patient has been taking, most probably no one else does either (with the possible exception of the patient himself/herself), as there almost always will be some degree of non-adherence and in some cases also overadherence. Information from the patients themselves was collected in our study, but it was not integrated with the rest of the data because most patients had very incomplete knowledge of which drugs they were using. For example, they could state that they were taking a white tablet ‘for their heart’ twice daily, but without knowing the specific name of the drug. This information could be important in the clinical setting, but for the purpose of our study, it was in most such cases not possible to decide whether it should be classified as a discrepancy or not related to the information in the hospital record. In a Danish study from 2003, Andersen et al5 concluded that second interviews and GP lists reveal extra information in two-thirds of cases, but they did not specify the relative importance of these two sources. However, in the clinical setting, both procedures should be routinely performed to compile a more comprehensive basis for drug prescribing.

Second, the harm assessment was made for each drug discrepancy separately rather than by performing one common harm rating for all discrepancies found for a patient, which it can be argued would have been more clinically relevant. Another factor regarding the harm assessment is that the scale employed did not take into account that a discrepancy potentially could have a positive effect for the patient, such as the omission of an obviously inappropriate drug.

On the other hand, this study also has some strengths. Among these is the fact that patients were included from three different types of wards in two hospitals, thereby including both surgical and medical causes of admittance. Moreover, the patients had a broad age range compared with most other studies.27 Thus, our results could be considered valid for a variety of patient populations and not just for older patients in internal medicine wards, who have been included in most previous studies. Consequently, we consider that medication reconciliation should have high priority regardless of the patient’s age or cause of admittance.

Another strength is the use of a multidisciplinary expert group and the inclusion of pilot cases to enhance the understanding of the severity score classification and increase the conformity of the evaluations in the group. Moreover, all discrepancies were discussed in the group, irrespective of whether the members had made identical classifications or not.

In conclusion, our study shows that a high proportion of patients admitted to hospital have discrepancies in their drug histories, and it supports the importance of collecting drug information from all available sources. Furthermore, we found no significant differences in the number of discrepancies when comparing different hospitals, hospital wards, genders, ages or levels of care. We therefore conclude that medication reconciliation should be considered an important potential quality improvement method that preferably should be performed for all patients admitted to hospital, even for patients receiving assistance from home care services or living in a nursing home.

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**Acknowledgements** We would like to thank the staff at the gastrointestinal surgery wards at St Olav’s University Hospital in Trondheim, the internal medicine wards at Ålesund Hospital in Ålesund and the geriatric ward at St Olav’s University Hospital for their collaboration. We are grateful to the Liaison Committee between
the Central Norway Regional Health Authority and the Norwegian University of Science and Technology and to the Central Norway Hospital Pharmacy Trust for funding the study.

Contributors JKS planned the study and collected and analysed the data, did the calculations and wrote the first version of the manuscript. OSI planned the study, analysed the data and contributed to the final version of the manuscript. TCM planned the study, collected and analysed parts of the data and contributed to the final version of the manuscript. RH collected and analysed parts of the data and contributed to the final version of the manuscript. OSp planned the study, analysed the data, did the calculations and contributed to the final version of the manuscript.

Funding This work was supported by the Liaison Committee between the Central Norway Regional Health Authority and the Norwegian University of Science and Technology.

Competing interests None declared.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement There is no more available data from this study.

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