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Deprescribing medicines in the acute setting to reduce the risk of falls

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ABSTRACT
Background Falls are a common cause of morbidity and hospitalisation in older people. Inappropriate prescribing and polypharmacy contribute to falls risk in elderly patients. This study’s aim was to quantify the problem and find out if medication review in the hospital setting led to deprescribing of medicines associated with falls risk.

Methods Admissions records for elderly patients were examined to identify those whose presenting complaint included a fall. Inpatient medication charts, pharmaceutical care notes, medical notes and discharge summaries were examined to identify any falls-risk medicines from admission histories and to determine if any medication review took place, and whether or not changes were made as a result. In particular deprescribing and dose reduction details were analysed.

Results 100 patients over 70 years old were admitted following a fall during the 2 months study period. The mean number of medicines on admission was 6.8 per patient with polypharmacy found in 62/100 (62%). One or more falls-risk medicine was found in 65/100 (65%) patients. Medicines review was carried out in 86/100 (86%) of patients, and 59/697 (8.5%) medicines were deprescribed. Pharmacist involvement in medication review led to a significant reduction in the number of falls-risk medicines per patient (p=0.002).

Conclusions Inappropriate prescribing and polypharmacy are found frequently in elderly patients at admission following a fall. Comprehensive medicines reviews should be carried out in all such patients with the objective of deprescribing or reducing doses to minimise risk of harm. Involvement of a pharmacist improves the rate of reduction of falls-risk medicines.

INTRODUCTION
Improving patient outcomes through medicines optimisation is a national priority and a key part of the work being undertaken by the National Institute for Health and Care Excellence (NICE) 2015 Quality Standard 86 includes the statement: ‘older people who present for medical attention because of a fall should have a multifactorial falls-risk assessment including cardiovascular examination and a medication review’.1

Polypharmacy is independently associated with negative outcomes in frailty particularly where there is decline in nutritional status, functional ability and cognitive capacity.4 Specifically, it has been shown that it increases the risk of falls in elderly populations and problematic polypharmacy is therefore recognised as a potentially modifiable contributing factor in falls and frailty.5

Freeland et al found that among patients aged 65 years and older who have experienced a fall in the past year, the addition of each medication above four increases their fall risk by 14%.6 Similarly, Damian et al found that each added medicine increased physician-reported falls in the preceding 30 days by 7%.7 These findings corroborate earlier prospective cohort studies which have shown that increasing medication use is associated with increased falls-risk scores in elderly patients.8 9

Further to this, the simultaneous discontinuation of many drugs was found to be ‘safe’ and appears to improve quality of life in elderly community dwellers.10

Other studies demonstrate that multiple medication lists in elderly patients are more likely to include falls-risk medicines. A systematic review found that benzodiazepines, antidepressants, antihypertensives and antipsychotics are among the most common medicines to be prescribed to older people with polypharmacy.11

A prospective cohort study of older adults coming into hospital following a fall, found that compared with robust elderly, those with frailty were prescribed a significantly higher number of falls-risk medicines. The number prescribed on discharge was additionally significantly associated with recurrent falls.12

The National Institute for Health and Care Excellence (NICE) 2015 Quality Standard 86 includes the statement: ‘older people who present for medical attention because of a fall should have a multifactorial falls-risk assessment including cardiovascular examination and a medication review’.1

Medication review has been defined as ‘any systematic assessment of the pharmacotherapy of an individual patient that aims to evaluate and optimise patient medication by a change in prescription either by a recommendation or by a direct change’.13 As part of the wider CLAHRC project on medicines optimisation in frailty, we defined levels of medication review in hospital14 (box 1).

Interim reviews can be doctor or pharmacist led (especially as part of medicines reconciliation on admission); the timing and breadth of the changes are adjusted according to patient need with their

immediate safety and well-being central to the decision but not necessarily with their full knowledge or participation (in contrast to comprehensive review). Some reviews may lie between the two categories so for ‘comprehensive’ we require the patient to be involved (or their carer if capacity is impaired) and a senior clinician (doctor or pharmacist) to lead.

It is imperative, however, to acknowledge that a condition for medication review is reliable medicines reconciliation at admission and discharge from hospital. Any review process typically begins with confirmation of a patient’s current medication; if inaccurate, decisions may be made based on the wrong information. Therefore, prescribing and deprescribing is safe only in the context of a full understanding of the patient’s drug history.

In order to support clinicians in reducing potentially inappropriate prescriptions (PIP) and optimise therapy, a PIP list for older patients was developed locally directly from the STOPP criteria of O’Mahony et al (see online supplementary appendix A). From this we further developed a deprescribing support tool ‘STOPIT’, used previously in an earlier version. The tool includes a section on medicines known to contribute to falls, commonly through orthostatic hypotension and sedation. These falls-risk medicines are listed in box 2.

Opioids are included as they are linked with falls through sedation and confusion, particularly in the vulnerable elderly. Other medicines may indirectly induce falls through mechanisms including vasovagal syndrome, carotid sinus hypersensitivity, bradycardia, tachycardia, periods of asystole and hypoglycaemia. These medicines are not on our definitive list for falls risk but their effects, risks and benefits in vulnerable patients are considered in a comprehensive review.

Aims and objectives

Our aim was to identify patients affected by falls, and find whether medication review in the acute setting led to deprescribing of falls-risk medicines.

Specific objectives were to:

- Identify all admissions aged over 70 years following a fall, on polypharmacy or problematic medicines.
- Examine hospital clinician involvement in medication review.
- Quantify nature and extent of deprescribing.

Ethical approval

Ethics approval was not required for this work as it is part of a service evaluation and improvement project. An ethics waiver was granted by Chelsea and Westminster Hospital NHS Foundation Trust Research and Development lead and National Research Ethics Service (NRES).

Setting

The study was conducted at Chelsea and Westminster Hospital, London (CWH) supported by CLAHRC NWL. Across all specialties, CWH admits an average of 740 adults a month, approximately 500 of whom are 70 years and older.

METHODS

Admissions coding data and emergency department discharge summaries (DSUMs) for the period 02 February 2015 to 31 March 2015 were examined to identify patients who were aged 70 years or older who were known to have had a fall and were in hospital as a result overnight. Recruited patients were followed-up prospectively, and data obtained from medical records and electronic prescribing records. All patients were seen by their usual pharmacist during their regular visits to the ward. Screening of prescriptions and documentation of pharmaceutical care contributions were undertaken as per ward pharmacy inpatient standard procedures. Weekly consultant ward rounds were attended by pharmacists as part of routine practice. No specific research interventions were made, however, all pharmacists were aware of the project through departmental teaching sessions aimed at improving communications regarding medication changes.

Final DSUMs were checked retrospectively for documentation of medication reviews and any changes made to regular medicines.

All data was collated by research student (AR) supervised by a specialist pharmacist (EW). A database was created for this cohort including details of the following:

- Medicines reconciliation on admission verified as ‘reliable’ by a pharmacist according to local policy.
- Medicines review documented as having taken place and by whom.
- Pharmacist role in any medication review or change.

Medicines purchased over the counter for acute use ‘as required’ were not included as regular medication. Neither were acute courses of, for example, antibiotics and simple (non-opiate) analgesia.

We define polypharmacy as the prescription of six or more regular medicines as used by other researchers. Appropriate polypharmacy requires that all medicines have a clear indication and are not PIPs.

Descriptive statistics were performed and falls-risk medicines prescribed before and after review were compared using the Kruskal-Wallis test for significant difference in rank.

RESULTS
Identification of patients with a history of falls
The total number of admissions to CWH for patients aged 70 or over during the study period was 1020, of whom 126 (12%) were identified having presented following a fall. Of these patients, 26/126 (21%) were ‘lost to follow-up by pharmacists’ as they were discharged from the Emergency Observation Unit without a prescription or other identifiable documentation about medication.

The remaining 100 were included in the analysis: the mean age was 85 years, the oldest was 101 years old; 61/100 (61%) were female. One patient was recorded as having an additional admission. In the remaining 21/86 documented ‘loss to follow-up’ patients, one (5%) case there is documentation of patients with a history of falls.

Prevalence of polypharmacy
In total, 679 medicines were taken by 100 patients before any medication review took place, a mean of 6.8 per patient (range 0–18, including unconfirmed medications).

Polypharmacy was present in 62/100 (62%) patients, and problematic polypharmacy (the patient was taking at least one medication identified as potentially inappropriate) was found in 57/62 (92%) of those with polypharmacy; or 57/100 (57%) of all patients. Fifty-five out of 62 (89%) polypharmacy admissions had a documented medication review.

Polypharmacy was found in more patients at discharge than at admission in the cohort of 100: while 5/62 (8%) polypharmacy patients were no longer on six or more medicines after review, 9/38 (24%) non-polypharmacy patients returned home with six or more medicines: thus 66/100 (66%) patients were discharged from hospital with polypharmacy. Bisphosphonates and vitamin D accounted for 49/54 (91%) of the newly prescribed medicines.

Falls-risk medications
One hundred and twelve out of 679 (16.5%) admission medicines were falls-risk medicines taken by 65 patients (table 1): 30/112 (26.7%) of falls-risk medicines were reduced or stopped. A Kruskal-Wallis test is used to find a significant reduction (p=0.004) in number of falls-risk medicines on these patients’ prescriptions postreview (91) compared with at admission (112).

Of the 65/100 (65%) patients found to be on a falls-risk medicine, 29/65 (45%) had no reduction in medication including 12/65 (18%) who had no documented medicines reconciliation. Six out of 65 (9%) did have a review and no changes were made to falls-risk medicines; other prescription items were started or doses increased in these cases. Twenty-three out of 65 (35%) patients had a comprehensive review which led to reduction of falls-risk medicines and 5/65 (8%) were no longer on any by discharge.

Medication changes
Overall, 38/100 (38%) patients had their regular prescriptions reduced in some way while in hospital. A total of 75/697 (10.8%) medicines were affected: 59/697 (8.5%) of all medicines were deprescribed, 12/697 (1.7%) were reduced to a lower dose and 4/697 (0.6%) medicines were held pending further review (table 2).

Of these changed medicines, 29/75 (39%) were falls-risk medicines in 21 patients (table 2). However, while there was a decrease in falls-risk medicines following review, after changes were made to regular medicines in hospital there was a net increase to 733 in the total number of medicines prescribed for the cohort. A list of all medicines deprescribed is in online supplementary appendix B.

Pharmacist involvement
Pharmacists were involved in 45/86 (52%) cases where it was documented that a medication review had taken place (with or without changes made). The number of falls-risk medicines decreased by a mean of 0.53 per patient before and after review in these 45 (not including the dose reductions). A Kruskal-Wallis test finds a significant reduction between the numbers before and after review (p=0.002). Of note, in all cases where medication review led to a reduction in falls-risk medicines for a patient a pharmacist was involved and the changes were documented.

Table 1 Medicines taken by elderly falls patients

<table>
<thead>
<tr>
<th>n=100 patients</th>
<th>On admission</th>
<th>At discharge</th>
<th>Difference</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of regular medicines</td>
<td>679</td>
<td>733</td>
<td>+54</td>
</tr>
<tr>
<td>No. of FRM</td>
<td>112</td>
<td>91</td>
<td>−21 (p=0.004)</td>
</tr>
<tr>
<td>Patients on ≥1 FRM</td>
<td>65</td>
<td>60</td>
<td>−5</td>
</tr>
<tr>
<td>% of all medicines that are FRM</td>
<td>16.4%</td>
<td>12.4%</td>
<td>−4%</td>
</tr>
<tr>
<td>Mean FRM/patient reviewed</td>
<td>1.19</td>
<td>0.939</td>
<td>−0.26</td>
</tr>
<tr>
<td>Mean FRM/patient reviewed with pharmacist involved</td>
<td>1.44</td>
<td>0.91</td>
<td>−0.53 (p=0.002)</td>
</tr>
<tr>
<td>FRM per patient not reviewed</td>
<td>0.77</td>
<td>0.77</td>
<td>0</td>
</tr>
</tbody>
</table>

*p=0.001; †Includes three patients on no medicines from admission through to discharge therefore not reviewed in this context.

FRM, falls-risk medicine.
Table 2  Falls-risk medicines deprescribed, held or reduced

<table>
<thead>
<tr>
<th>Patient</th>
<th>Medicine</th>
<th>Deprescribed (D)</th>
<th>Held (H)</th>
<th>Reduced dose (R)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Tramadol</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Fentanyl</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Lercanidipine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Losartan</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Ramipril</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>Buprenorphine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>Buprenorphine</td>
<td>R</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>Lercanidipine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>Lisinopril</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>Moxonidine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>Tolterodine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>Losartan</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>13</td>
<td>Tramadol</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>14</td>
<td>Amlodipine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>Oxybutynin</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>Diazepam</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>Ramipril</td>
<td>R</td>
<td></td>
<td></td>
</tr>
<tr>
<td>18</td>
<td>Lisinopril</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>19</td>
<td>Irbesartan</td>
<td>H</td>
<td></td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>Cetirizine</td>
<td>D</td>
<td></td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>Zopiclone</td>
<td>D</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

DISCUSSION

A fall, whether or not an injury is sustained, is a common reason for patients aged 70 or older to present to hospital. We identified 126 patients in a period of 2 months who were admitted (for at least one night) from a total of 1020 admissions recorded in this age group during this time (12%). We followed-up a cohort of 100, 97 of whom were on regular medication from admission. Polypharmacy, defined as six or more medicines was prevalent (62%) with an average of 6.8 per patient. In just five polypharmacy patients, all medicines were considered appropriate in the context of their age and history of falls.

We identified problematic polypharmacy in 57% of our cohort (inpatients over 70 years old who had fallen prior to admission). PIPs have previously been estimated to occur in about a third of older patients. We identified PIPs included all anticholinergics, known to contribute to falls but we would argue many medicines are unrecognised as being in this category. We note one small study of anticholinergic burden and falls identified regular anticholinergic prescriptions in 22% of patients, and using their criteria only 10% of patients had no identified risk of falls from their medicines. We used a shorter, evidence-based list of falls-risk medicines, though inclusive of all anticholinergics and found 65% of our patients on one or more. The association of anticholinergics with falls has been reaffirmed in a community study in men over 65 years.

Despite the increase in the number of polypharmacy patients (nine more at discharge than on admission), we have shown that review improved prescribing and optimised medicines overall. The increase was due largely to the prescribing of calcium and vitamin D supplementation in our cohort; these additional medicines are considered appropriate according to the STOPP/START criteria although an alternative view is emerging which may lead to further consideration of local guidelines.

The NICE pathway on falls in older people recommends that those on ‘psychotropic’ medicines should have their medication reviewed, with a specialist input if appropriate, and discontinued if possible to reduce their risk of falling. We did not include all medicines affecting mood; only antipsychotics, benzodiazepines and ‘z-drugs’ which are considered directly linked with falls. We may therefore need to update our list (box 2); if we had included all psychotropics the number of falls-risk medicines on admission in our study would be increased by 16 to a total of 128.

A study by Browne et al and colleagues in 2014 looked at 50 inpatients at risk of falls and found that only 20% of falls-risk medicines were suitable for change after reviews, which were found to be time and resource intensive. However, the authors suggest targeting resources to those on polypharmacy (four or more in their study) including as required (prn) rather than restricting to ‘regular’ medications results in greater deprescribing rates. Another study demonstrated that the risk of subsequent falls is reduced in patients for whom falls-risk medicines are reduced, suggesting that it is an effective use of resource if the medical cost of a fall is taken into account.

We have put in place specific tools to prompt safe, patient-centred deprescribing when falls-risk medicines are identified and although we did not measure their uptake and use in this project, we are further promoting STOPIT and teaching about PIPs to junior staff. We believe pharmacists can and should be more proactive in stopping medicines and such tools encourage and promote good prescribing practice. This is supported by our finding that suggests pharmacist involvement in medication reviews significantly reduces the number of falls risk medicines. However, the mean reduction in number was less than one drug (0.53) per patient. The benefit in terms of reducing falls is unclear but there are other benefits of stopping medicines, such as decreased pill burden and costs. We plan to measure these with follow-up of a larger cohort, also checking for readmissions and further falls over a longer period.

Our recommendation is to involve pharmacists in identifying patients at risk, beginning the process of comprehensive medication review from the point of verifying the drug history (in line with Royal Pharmaceutical Society standards) and prompting the prescriber at all stages of the patient journey through the hospital and at transfer. For this we may need to improve knowledge and understanding of anticholinergic medicines and switch patients’ prescriptions to lower anticholinergic burden when they are at risk of falls (or confusion). In addition, we would explore ways in which the electronic prescribing record system could alert to the need for a medicines review following medicines reconciliation documentation as described by Graabaek et al.

Limitations

The period of study was February and March 2015. These months may not be typical; seasonal variation in falls rates is likely. However, prescribing of falls-risk medicines is not expected to vary with time of year.

It is unclear if the number of reviews documented is a true reflection of the number actually carried out on elderly patients admitted following a fall. Reviews without changes being made are under-represented.
Only regular medicines meant for long-term continuation were included in our analysis. We, therefore, excluded acute use of several contributors to falls risk, for example, sedating antihistamines, if they were as required medicines. Further to this our definitive falls-risk medicine list was limited to those with evidence linking them to actual falls. Other antihypertensive medicines such as β-blockers are implicated, for example, if the patient experiences a postural drop in blood pressure; there is a case for extending our list to include these in line with other published data.17

CONCLUSION

Inappropriate prescribing and polypharmacy are found frequently in elderly patients at admission following a fall. Comprehensive medicines reviews should be carried out in all such patients with the objective of deprescribing or reducing doses to minimise risk of harm. This is significantly more likely to be achieved with a pharmacist input. We suggest involving pharmacists in reviews from the start of medicines reconciliation at admission (earlier where possible), improving recognition of falls-risk medicines by all clinicians and supporting prescribers in deprescribing.

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