A systematic review of the outcomes reported in trials of medication review in older patients: the need for a core outcome set

Running head: Outcomes reported in trials of medication review in older patients

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Summary

**Aim:** Medication review has been advocated as one of the measures to tackle the challenge of polypharmacy in older patients, yet there is no consensus on how best to evaluate its efficacy. This study aimed to assess outcome reporting in trials of medication review in older patients.

**Methods:** Randomized controlled trials (RCTs), prospective studies and RCT protocols involving medication review performed in patients aged 65 years or older in any setting of care were identified from: (1) a recent systematic review; (2) RCT registries of ongoing studies; (3) the Cochrane library. The type, definition, and frequency of all outcomes reported were extracted independently by two researchers.

**Results:** 47 RCTs or prospective published studies and 32 RCT protocols were identified. A total of 327 distinct outcomes were identified in the 47 published studies. Merely one fifth (21%) of the studies evaluated the impact of medication reviews on adverse events like adverse drug reactions or drug-related hospital admissions. Most of the outcomes were related to medication use (n=114, 35%) and healthcare use (n=74, 23%). Very few outcomes were patient related outcomes (n=24, 7%). A total of 248 distinct outcomes were identified in the 32 RCT protocols. Overall, the number of outcomes and the number and type of health domains covered by the outcomes varied largely.

**Conclusion:** Outcome reporting from RCTs concerning medication review in older patients is heterogeneous. This review highlights the need for a standardised core outcome set for medication review in older patients, to improve outcome reporting and evidence synthesis.
**Introduction**

Older patients are often exposed to polypharmacy [1,2]. This increases the risk of adverse drug reactions (ADR) and the cost of medications [3–6]. Performing a medication review has been shown to be an efficient approach to optimise the quality of prescriptions in older patients [7,8]. Medication review has been defined by the NICE guidelines as “a structured, critical examination of patient’s medicines with the objective of reaching an agreement with the patient about treatment, optimising the impact of medicines, minimising the number of medication related problems and reducing waste” [9].

Numerous randomized controlled trials (RCT) have been performed to evaluate the impact of medication review on clinical, patient reported, and economic outcomes. Several systematic reviews and meta-analyses have therefore been conducted to summarize the effectiveness of medication review in various settings [7,8,10–20]. However, the heterogeneity of outcomes reported in the RCTs has limited the quality of the conclusions of these systematic reviews. Robust meta-analyses could be performed for only a few outcomes, including hospitalisation and death [15–17,21]. For other outcomes, results were essentially summarized in a descriptive way because of heterogeneity in the choice and definition of the outcomes [17–20,22].

Outcome reporting bias is an under-recognised problem that affects the conclusions in a substantial proportion of systematic and Cochrane reviews [23–25]. This bias has been defined as selection (on the basis of the results) of a subset of the original variables recorded for inclusion in a study publication [26]. In a large unselected cohort of Cochrane systematic reviews, more than half of the reviews did not include full data for the review primary outcome of interest from all eligible trials [23]. Consequently, the updated Consolidated Standards of Reporting Trials (CONSORT) statement has recently recommended the use of identified and well defined outcomes in RCTs. In other fields of medicine, authors have conducted systematic reviews to reveal and quantify the heterogeneity and inconsistency of outcome reporting in a given research area [27–30]. According to the OMERACT
(Outcome Measures in Rheumatology) and the COMET (Core Outcome Measures in Effectiveness Trials) guidelines, some of these works have led to the development of core outcomes sets designed to increase the quality of outcome reporting and evidence synthesis in future research [31,32].

The purpose of this review was to undertake an in-depth analysis of outcome reporting to identify potential heterogeneity in outcomes reported from RCTs and prospective studies of medication review in older patients.

Methods

Study selection
We considered RCT, quasi-RCT, and other prospective interventional studies which investigated the effect of medication review performed in patients aged 65 years or older [33]. This age limit was widely used in trials of medication review during the last decades and corresponds to the WHO age limit to define older people. The following studies were excluded: studies published before 2000; studies predominantly including patients younger than 65 years old; retrospective studies; no outcome reported; sample size lower than 50 participants; medication reviews for a specific disease or condition (e.g. chronic heart failure) or as part of a multifaceted approach. By multifaceted approach we meant a complex intervention which contained additional interventions to the medication review (e.g. physiotherapy, nutritional advice, occupational therapy).

Search strategy
We performed a systematic review of published studies. Our starting point was the systematic review published in 2014 by Lehnbom et al. [10], which purpose was to examine the evidence regarding the effectiveness of medication review to improve clinical outcomes. Lehnbom et al. identified 43 studies published before March 2014 [10]. As the purpose of the present systematic review was different in nature, we only used the result of the search strategy, i.e. the list of the published studies included in the systematic review. Only the studies complying with our inclusion...
and exclusion criteria were included. In addition, an update of the literature search using the same strategy/queries was performed for the period of March 2014 and July 2015 (the search strategy from [10] is given in supplementary data). Two reviewers (LP and JBB) independently assessed the title and abstracts of studies resulting from the searches. Full texts were investigated for all eligible published studies by two independent reviewers (LP and JBB). Any disagreement on the inclusion of a study was resolved by discussion and consensus. A third reviewer (AS) was involved if needed.

In addition, we used the search terms from the systematic review to identify RCTs protocols related to medication review on the following RCT websites: WHO international clinical trials registry platform: http://apps.who.int/trialsearch/; EU Clinical Trials Register: https://www.clinicaltrialsregister.eu/ctr-search/search; US Clinical Trials register: https://clinicaltrials.gov. Two independent reviewers (BB and JBB) first assessed the titles and subsequently the summaries of the RCTs protocols identified by the queries. Any disagreement on the inclusion of a study was resolved by discussion and consensus. A third reviewer (AS) was involved if needed. It was verified if protocols had matured to full report publications. If this was the case, the paper was evaluated as mentioned above and added to the set of published studies.

Finally, the search terms from the systematic review were also used to identify relevant Cochrane reviews. A reviewer (JBB) identified the relevant Cochrane systematic reviews and then extracted the eligible original studies in the selected Cochrane systematic reviews. The selection process was checked by a second reviewer (AS).

**Data extraction**

An electronic data extraction form was developed with Epidata® software and pilot tested to increase the reliability of the data extractions. All data extractions on outcomes and outcome measurement instruments were performed by two independent reviewers (ST and JBB for RCT protocols; LP and JBB for published studies). Any disagreement was resolved by discussion and consensus. A third reviewer (AS) was involved if needed.

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The characteristics of the RCT protocols and the published studies were extracted by one reviewer (JBB) and included: setting (hospital setting, primary care, nursing home); number of patients intended to be recruited (RCT protocols) or actually included (published studies); mean age (published studies only).

Outcomes used to compare the two groups under investigation (in RCTs and prospective before/after studies) or to evaluate the medication review process were extracted: name of the outcome in free text (what was measured in published studies and what was planned to be measured in protocols); primary or secondary outcome. For each outcome, the following data about measurement instrument were extracted from published studies: which instrument was used to measure the outcome (free text); was the method of measurement clearly defined (the reviewer answered “Yes” if he/she believed that another researcher could reproduce the procedure and its measurement with the explanations provided in the methods section). As the data provided in RCT protocols are often less detailed than in published studies, the data on measurement instrument were extracted when available.

Classification of outcomes into health domains and subdomains

The classification of the outcomes extracted from the included studies was achieved in several steps. Firstly, a list of 19 pre-defined subdomains corresponding to the most frequently reported outcomes (e.g. hospitalisation, all-cause death) were identified from the systematic review updated in this study [10]. These pre-defined subdomains were used by the two reviewers to associate each extracted outcome to a subdomain. The reviewers were free to propose a new subdomain if an outcome did not fit with the proposed subdomains. Secondly, a new list of subdomains was drafted on the basis of the pre-defined and the newly proposed subdomains. A consultation exercise was then performed with experts in clinical pharmacy and geriatricians. The objective was to get consensus on subdomain terms, to avoid major overlaps between subdomains, and to aggregate subdomains into health domains. The OMERACT filter 2.0 was used to organise this classification. A
total of 57 subdomains were identified and grouped into eight health domains (supplementary data; eTable 1).

Analysis
Quantitative variables were described by median, first and third quartiles (Q1, Q3), minimum and maximum because of skewed distributions. Qualitative variables were expressed as frequencies and percentages.

Results

Selection of studies and RCT protocols
In total, 47 published studies [34–80] and 32 RCT protocols were included in this systematic review. Details on the sources, reasons for exclusion, and selection process are presented in Figure 1.

Characteristics of the studies and RCT protocols included for data abstraction
The characteristics of the published studies and the RCT protocols included for data abstraction are provided in Table 1. The median number of participants included in published studies and intended to be included in RCTs described in protocols were similar. The studies were performed mainly in Europe, followed by Australia and USA. More recent RCT protocols have mainly been developed in European countries. The median number of outcomes per study or per RCT protocol was seven. This number varied importantly, from a single outcome in a published study [69] to 19 distinct outcomes in a RCT protocol [81].

Outcomes identified in the published studies and RCT protocols
The published studies and RCT protocols covered a median of four of the eight health domains identified (range: 1 to 6 for published studies; 2 to 7 for RCT protocols). The results are summarised in Figure 2 and detailed per sub-domains in eTable 2. More than 50% of all outcomes reported in published studies concerned medication use (e.g. the number of drugs, the number of potentially inappropriate medications, or overuse) or healthcare use (e.g. hospitalisations or GP visits). Other
health domains like patient reported outcomes, geriatric syndromes, or costs evaluation were far less frequently investigated. Only one fifth (21%) of the 47 published studies have evaluated the impact of medication reviews on adverse events such as ADR or drug-related hospital admissions. Consequently, less than one in ten published studies (n=4; 9%) had a primary outcome in the adverse event domain.

Outcomes related to medication use were reported by 89% of published studies and 94% of RCT protocols, suggesting an informal consensus on reporting this health domain. However, there was substantial variation in the subdomains used to explore this health domain between studies. Twelve different subdomains were identified and variously measured in the studies. The number of studies using each subdomains varied from only one time (“anticholinergic drug use” in 1 RCT protocols) to 26 times (“number of drugs” in 17 published studies and 9 RCT protocols). Eight of these 12 subdomains were used at least once as a primary outcome in a published study or in a RCT protocol. This between-subdomain heterogeneity was also observed for the seven other health domains, as detailed in eTable 2. For example, seven, 10, and 12 different subdomains were variously used to explore the healthcare use, patient-reported outcome, and geriatric syndromes domains, respectively. In total, nearly half of the subdomains (n=28; 49%) were used by less than 10% of the 79 selected studies (published studies and RCT protocols). Conversely, only four subdomains were used by a third or more of the 79 selected studies, namely “number of drugs”, “hospitalisation”, “all-cause death”, and “health-related quality of life”.

Clarity of measurement instrument reported

The total number of outcomes and the specific number of outcomes for which an instrument measurement was clearly described in the methods are presented per health domain in Table 2 and per subdomain in supplementary data (eTable 3). In total, the measurement instruments were considered clearly reported in 52% to 83% of all domains, except for cost outcomes. In the costs domain, the methods most often insufficiently described the source of drug costs or costs related to
healthcare use, or the way in which calculations were performed. For medication use, the methods were considered clear in two thirds (65%) of the outcomes. The outcomes most often considered poorly defined were the outcomes “Number of drugs”, and “Drug-drug interactions”.

**Discussion**

This systematic review provides an analysis on which health domains have been and are currently evaluated in published and planned/ongoing studies (RCT protocols) of medication review in older patients. It reveals that there is substantial heterogeneity, as the number of outcomes and the number and type of health domains covered by the outcomes vary from one study to another. Furthermore, it shows that some essential health domains including adverse events and patient related outcomes have been poorly evaluated in published studies. These results highlight the need for developing a core outcome set for medication review in older patients. Having strong outcome data would help clarifying if and how medication review can be effective, among which population, in which context, and what are the important contextual factors which support positive outcomes.

Not surprisingly, our results have shown that most studies of medication review in older patients have reported, or planned to report, outcomes related to the medication use domain. These outcomes were often the primary outcomes. However, the 13 distinct subdomains identified in the medication use domain were variously used by authors (eTable2). Despite this heterogeneity between subdomains, several systematic reviews of RCTs have tried to summarize the effect of medication review on such outcomes. Their results showed that medication review is effective in improving drug appropriateness, drug underuse, or potentially inappropriate prescribing, but there were more conflicting results for drug-related problems [7,8,18,20]. Evaluating the effects of medication review on more specific outcomes such as inappropriate use of antipsychotic drugs was limited due to inconsistency in the various definitions of the outcome used across the studies [22]. We identified that clarity in the reporting of the measurement instruments in some domains varied
substantially and this impacts upon the extent to which the outcomes from such studies can be accurately interpreted, and may limit the validity of comparison of results across studies.

Our results also show that many published studies have reported outcomes related to the use of healthcare resources domain, mainly hospitalisations. Meta-analyses however have failed to identify a significant effect of medication review on hospital admissions, length of hospital stays, or emergency department visits [12,15–17,21,82]. These negative results may have discouraged new studies to use these outcomes as a primary outcome. One hypothesis is that these outcomes are influenced by many other factors, which could explain why this outcome is less often used as primary outcome in RCT protocols.

Ten (21%) published studies reported outcomes related to the adverse events domain. This may be due to the difficulty in detecting and adjudicating adverse events prospectively during an RCT. Hence, the effects of medication review on ADR or drug-related hospital admissions remain unclear and have not been assessed by any meta-analysis. Reporting bias of harm outcomes in RCTs and systematic reviews has been identified as an important problem [83]. It seems particularly detrimental that harm outcomes have been so poorly reported in RCTs of medication review, as the purpose of this intervention is actually to manage the risk of prescribed drugs and to reduce the number of drug-related problems. A higher proportion of on-going studies planned to report adverse events according to their protocols. However, it is uncertain whether this greater focus on these outcomes in protocols will be released in the final studies’ results. Outcomes related to costs were also less extensively investigated. The effect of medication review on costs is therefore most often analysed through a narrative approach in systematic reviews [13,20].
Patient reported outcomes have been investigated in 43% of published studies and are planned to be investigated in 69% of RCT protocols. Most of these outcomes concern health-related quality of life (eTable 2). Very few studies have investigated or will investigate subdomains like drug-regimen complexity or the alignment of drug regimen with patient preference, although these aspects are included in the definition of a medication review [9].

Several systematic reviews of medication review identified that inconsistency and imprecision in outcome reporting limited their analyses and conclusions [13,14,17,18,20,22]. Our present review has confirmed and extended these findings. While other systematic reviews aimed at pooling and summarizing results on a limited number of outcomes, we have extracted and reported all kinds of outcomes used in this field. Several authors have used hospitalisation or mortality to assess the clinical effectiveness of a medication review [12,15,16,21]. However, our results suggest that the choice to perform meta-analyses on these outcomes are most likely related to the high proportion of studies reporting these outcomes as opposed to their relevance. In fact, hospitalisation or death in older patients with polypharmacy and multimorbidity can be related to many factors other than medications [84,85]. Incidence of ADR or drug-related hospital admissions may be more relevant to assess the effectiveness of a medication review, but it is indeed much more difficult and time-consuming to detect and adjudicate properly all ADEs or drug-related hospital admissions during a RCT than to record all-causes hospitalisations or mortality.

In summary, our results highlight the need for a core outcome set (COS) for future studies of medication review in older patients. A COS is an agreed standardised collection of outcome variables which should be measured and reported in all trials for a specific condition or clinical area. The development of a COS would tackle the challenge of obtaining a consensus on which outcomes are deemed essential for all stakeholders involved in the management of a given condition, including patients. It has the potential to reduce heterogeneity between trials, lead to research that is more likely to measure relevant outcomes, and enhance the value of evidence synthesis by reducing the risk of outcome reporting bias and ensure that all trials report usable information [86–88].

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Moreover, in a second step, the measurement instrument recommended to be used to evaluate each outcome included in the COS can be determined [89]. This aspect may be important as our study has shown that the methods were not clearly detailed in a significant proportion of studies for all categories of domains and subdomains. A consensus on the measurement instrument has also the potential to increase the quality of value of evidence from subsequent systematic reviews and allowing meaningful meta-analyses to be performed. As part of the European Commission-funded OPERAM project, this systematic review will serve as a starting point to developing a COS for clinical trials of medication review in older patients. The OPERAM project will perform a multicentre randomised controlled trial to assess the impact of an intervention to optimise pharmacotherapy and to enhance compliance in 1900 multimorbid patients aged 70 years and older. In order to tackle the challenge of measuring relevant outcomes for these patients, the OPERAM project included the development a COS suitable for all settings.

Some limitations of this review should be mentioned. As the aim of this project was of descriptive nature, we did not perform analytical analyses to formally compare results from published studies and protocol, and had a too low number of studies to evaluate if the between study variation could be partially explained by the setting (hospitalised versus nursery home versus community dwelling). This systematic review stopped in July 2015, more than one year before its submission. However, the variation in outcome reporting between published studies should not have changed considerably and has been captured indirectly through the analysis of RCT protocols. It seems therefore unlikely that an updated version would have impacted on the results. The selection criteria were those from the initial systematic review by Lehnbom et al. [10], which could have influenced the outcomes identified. For example, smaller studies with less than 50 subjects, which were excluded from the review, may have reported outcomes that require detailed review of patients or records, such as ADRs or drug-related admissions, more often. Variation between studies was investigated descriptively at the subdomain level but we didn’t report variation between outcomes in a given subdomain. The outcomes may vary in a subdomain in the way they were reported (e.g. number of
potentially inappropriate medications per patient or number of patients with at least one potentially inappropriate medication) or in the way they were measured (e.g. Beers criteria, STOPP criteria, PRISCUS list). Another limitation is that the categorization into domains and subdomains may be somehow arbitrary. We were also reliant upon the descriptions of domains and measures in the published studies and, as we identified, at times these were not clearly described. We used a consultation exercise as other studies or study protocols have [30,90], as we could not identify an existing framework to guide this process.

**Conclusion**

This systematic review revealed a significant heterogeneity in outcome reporting in studies of medication review in older patients. Some essential health areas including adverse events and patient reported outcomes have been insufficiently evaluated. We advocate the development of a COS to improve the quality of outcome reporting and evidence synthesis in future research in this important field.

**Competing interest**

All authors have completed the Unified Competing Interest form and declare no support from any organization for the submitted work.

**Authors contribution**

JBB conceptualized and designed the study, performed the selection of studies and data extraction, carried out analyses and interpretation of data, drafted the initial manuscript, and approved the final manuscript as submitted.
LGP designed the study, performed the selection of studies and data extraction, critically reviewed the manuscript and approved the final manuscript as submitted.

ST performed data extraction, critically reviewed the manuscript and approved the final manuscript as submitted.

BB performed the selection of studies, critically reviewed the manuscript and approved the final manuscript as submitted.

OD participated in study design and critically reviewed the manuscript and approved the final manuscript as submitted.

AR participated in study design and critically reviewed the manuscript and approved the final manuscript as submitted.

JW designed the study, critically reviewed the manuscript and approved the final manuscript as submitted.

AS conceptualized and designed the study, checked the selection of studies and data extraction, carried out interpretation of data, drafted the initial manuscript, and approved the final manuscript as submitted.

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Figure 1: Flowchart of included studies and RCT protocols.

Abbreviations: RCT: Randomized controlled trials; MRev: Medication Review; SR: Systematic review
Figure 2: Percentage of health domains covered by the 47 published studies and 32 RCT protocols included in the systematic review, according to the outcomes identified. The percentage of studies in which the domain was covered by a primary outcome is given in darker colour.

Abbreviations: HC: healthcare; Pub. St.: published studies; St. Pr.: study protocols

For the full list of subdomains contained in each domain, please refer to eTable 1.
**Table 1:** Summary characteristics and demographics of included studies (published studies included 35 randomized controlled trials and 12 prospective studies).

<table>
<thead>
<tr>
<th></th>
<th>Published studies (n=47)</th>
<th>RCT protocols (n=32)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Number of study participants</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>334</td>
<td>300</td>
</tr>
<tr>
<td>Interquartile range</td>
<td>[170; 568]</td>
<td>[195; 600]</td>
</tr>
<tr>
<td>Range</td>
<td>[50; 7202]</td>
<td>[50; 3685]</td>
</tr>
<tr>
<td><strong>Average age</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>80</td>
<td></td>
</tr>
<tr>
<td>Interquartile range</td>
<td>[75.7; 82.7]</td>
<td></td>
</tr>
<tr>
<td>Range</td>
<td>[73.5; 86.8]</td>
<td></td>
</tr>
<tr>
<td><strong>Setting</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospital</td>
<td>15 (32%)</td>
<td>11 (34%)</td>
</tr>
<tr>
<td>Community</td>
<td>19 (40%)</td>
<td>16 (50%)</td>
</tr>
<tr>
<td>Nursing homes</td>
<td>13 (28%)</td>
<td>5 (16%)</td>
</tr>
<tr>
<td><strong>Country</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EU</td>
<td>28 (Belgium 2; Denmark 3; Germany 1; Ireland 3; The Netherlands 1; Spain 2; Sweden 5; UK = 11)</td>
<td>25 (Belgium 2; France 1; Germany 7; Ireland 2; Italy 1; Netherlands 6; Norway 1; Slovenia 1; Spain 1; Sweden 1; Europe MC 1)</td>
</tr>
<tr>
<td>Australia</td>
<td>8</td>
<td>3</td>
</tr>
<tr>
<td>USA</td>
<td>7</td>
<td>1</td>
</tr>
<tr>
<td>Others</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td><strong>Year of publication</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2000 – 2005</td>
<td>14 (30%)</td>
<td></td>
</tr>
<tr>
<td>2006 – 2010</td>
<td>6 (13%)</td>
<td></td>
</tr>
<tr>
<td>2011 – 2015</td>
<td>27 (57%)</td>
<td></td>
</tr>
<tr>
<td><strong>Number of outcomes</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>Interquartile range</td>
<td>[5; 10]</td>
<td>[6; 9]</td>
</tr>
<tr>
<td>Range</td>
<td>[1; 14]</td>
<td>[2; 19]</td>
</tr>
</tbody>
</table>
Table 2: Number of outcomes per health domain with the clarity of the measurement instrument used to measure them, as identified in the published studies or in RCT protocols.

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Published studies (317 outcomes)</th>
<th>RCT protocols (240 outcomes)</th>
<th>Published studies</th>
<th>RCT protocols</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%&lt;sup&gt;(a)&lt;/sup&gt;</td>
<td>n</td>
<td>%&lt;sup&gt;(a)&lt;/sup&gt;</td>
</tr>
<tr>
<td>Medication use</td>
<td>99</td>
<td>31%</td>
<td>51</td>
<td>21%</td>
</tr>
<tr>
<td>Use of HC resources</td>
<td>70</td>
<td>22%</td>
<td>44</td>
<td>18%</td>
</tr>
<tr>
<td>Adverse events</td>
<td>11</td>
<td>3%</td>
<td>17</td>
<td>7%</td>
</tr>
<tr>
<td>Death</td>
<td>18</td>
<td>6%</td>
<td>11</td>
<td>5%</td>
</tr>
<tr>
<td>Patient reported outcomes</td>
<td>32</td>
<td>10%</td>
<td>38</td>
<td>16%</td>
</tr>
<tr>
<td>Geriatric syndrome</td>
<td>24</td>
<td>8%</td>
<td>42</td>
<td>18%</td>
</tr>
<tr>
<td>Costs</td>
<td>21</td>
<td>7%</td>
<td>16</td>
<td>7%</td>
</tr>
<tr>
<td>Process evaluation</td>
<td>42</td>
<td>13%</td>
<td>21</td>
<td>9%</td>
</tr>
</tbody>
</table>

Abbreviations: HC: healthcare

a) Based on the total number of identified outcomes (n = 317 for published studies; n = 240 for RCT protocols)

b) Based on the total number of outcomes identified in the given domain (value is “n” of the column “Outcomes”)

* The method of measurement was considered clearly defined if the reviewer believed that another author could reproduce the procedure and its measurement with the explanations given in the method section.
Appendix 1: Search strategy

Medication reconciliation
Embase (2000-2014)
1. medication reconciliation.mp. or exp medication therapy management/
2. medication discrepancy.mp.
3. adverse drug events.mp.
4. 1 or 2 or 3
5. exp hospital/
6. exp patient admission/
7. exp patient discharge/
8. exp patient transfer/
9. exp emergency medicine/
10. exp emergency health service/
11. 5 or 6 or 7 or 8 or 9 or 10
12. 4 and 11
13. limit 12 to (english language and embase and yr="2000 - 2014")
14. exp general practice/
15. exp general practitioner/
16. exp primary health care/
17. 14 or 15 or 16
18. 4 and 17
19. limit 18 to (english language and embase and yr="2000 - 2014")
20. exp home for the aged/
21. exp nursing home/
22. exp elderly care/
23. residential aged care facilities.mp.
24. 20 or 21 or 22 or 23
25. 4 and 24
26. limit 25 to (english language and embase and yr="2000 - 2014")
27. 13 or 19 or 26

Ovid Medline (2000-2014)
1. medication reconciliation.mp. or exp medication therapy management/
2. medication discrepancy.mp.
3. adverse drug events.mp.
4. 1 or 2 or 3
5. exp hospital/
6. exp hospital admission/
7. exp hospital discharge/
8. exp patient transfer.mp.
9. exp emergency medicine/
10. exp emergency health service/
11. 5 or 6 or 7 or 8 or 9 or 10
12. 4 and 11
13. exp general practice/
14. exp general practitioner/
15. exp primary health care/
16. 13 or 14 or 15
17. 4 and 16
18. exp homes for the aged/
19. exp nursing homes/
20. exp health services for the aged/
21. exp elderly care/
22. residential aged care facilities.mp.
23. 18 or 19 or 20 or 21 or 22
PecInfo (2000-2014)
1. medication reconciliation.mp
2. medication discrepancy.mp
3. adverse drug events.mp
4. 1 or 2 or 3
5. exp hospitals/
6. exp hospital admission/
7. exp hospital discharge/
8. exp client transfer/
9. emergency medicine.mp
10. exp emergency services/services/
11. emergency medical services.mp
12. 5 or 6 or 7 or 8 or 9 or 10 or 11
13. 4 and 12
14. exp primary health care/ or exp family medicine/
15. exp general practitioners/ or exp general practice.mp
16. 14 or 15 or 16
17. 4 and 17
18. exp nursing homes/ or exp residential care institutions/ or exp elder care/
19. health services for the aged.mp
20. residential aged care facilities.mp
21. 22 or 23
22. 4 and 22
23. 4 or 12 or 24
24. 13 or 14 or 23
25. limit 24 to (english language and yr="2000-2014")

CINAHL (Jan 2009 - Jan 2014)
1. (MH "Medication Reconciliation")
2. "medication discrepancy"
3. (MH "Adverse Drug Events")
4. S1 OR S2 OR S3
5. (MH "Hospitals, Urban") OR (MH "Hospitals, Community") OR (MH "Hospitals, Public")
   OR (MH "Hospitals, Rural")
6. "hospital admission"
7. "hospital discharge"
8. "patient transfer"
9. (MH "Emergency Medicine")
10. (MH "Emergency Service")
11. (MH "Emergency Medical Services")
12. S5 OR S6 OR S7 OR S8 OR S9 OR S10 OR S11
14. (MH "Family Practice") OR (MH "Physicians, Family")
15. (MH "Primary Health Care")
16. (MH "Medical Practice")
17. S14 OR S15 OR S16
18. S4 AND S17
19. "home for the aged"
20. (MH "Nursing Homes")
21. (MH "Health Services for the Aged")
22. "elderly care"
23. "residential aged care facilities"
PsyInfo (2000-2014)
1. exp Hospitals/
2. exp Hospital Admission/
3. exp Hospital Discharge/
4. exp Client Transfer/
5. emergency medicine.mp.
6. exp Emergency Services/
7. emergency medical services.mp.
8. 1 or 2 or 3 or 4 or 5 or 6 or 7
9. exp Family Physicians/
10. exp Primary Health Care/ or exp Family Medicine/
11. exp General Practitioners/ or general practice.mp.
12. 3 or 5 or 11
13. exp Nursing homes/ or exp Residential Care institutions/ or exp Elder Care/
14. health services for the aged.mp.
15. residential aged care facilities.mp.
16. drug utilization review.mp.
17. medication review.mp
18. medication therapy management.mp.
19. medication history review.mp.
20. medical history review.mp.
21. 16 or 17 or 18 or 19 or 20
22. 8 and 21
23. 12 and 21
24. 13 or 14 or 15
25. 21 and 24
26. 32 or 23 or 24
27. limit 26 to (english language and yr="2000 - 2014")

CINAHL (Jan 2000 - Jan 2014)
1. (MH "Hospitals, Urban") OR (MH "Hospitals, Community") OR (MH "hospitals, Public") OR (MH "hospitals, Rural")
2. "hospital admission"
3. "hospital discharge"
4. "patient transfer"
5. (MH "Emergency Medicine")
6. (MH "Emergency Services")
7. (MH "Emergency Medical Services")
8. 51 OR 66 OR 27 OR 61 OR 0 OR 51 OR 51 OR 51
9. (MH "Family Practice") OR (MH "Physicians, Family")
10. (MH "Primary Health Care")
11. (MH "Medical Practice")
12. 59 OR 51 OR 511
13. "Home for the aged"
14. (MH "Nursing Homes")
15. (MH "Health Services for the Aged")
16. "Elderly care"
17. "Residential aged care facilities"
18. 513 OR 514 OR 515 OR 516 OR 517
19. (MH "utilization review")
20. "Medication management therapy"
21. "Medical history review"

22. "medication history review"
23. 019 OR 020 OR 022
24. 58 AND 523
25. 58 AND 524
26. 518 AND 524
27. 524 OR 525 OR 526 Limiters — Data published from: 20000101-20140331; English Language

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