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## The effects of on-screen, point of care computer reminders on processes and outcomes of care

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### Abstract

**Background**—The opportunity to improve care by delivering decision support to clinicians at the point of care represents one of the main incentives for implementing sophisticated clinical information systems. Previous reviews of computer reminder and decision support systems have reported mixed effects, possibly because they did not distinguish point of care computer reminders from e-mail alerts, computer-generated paper reminders, and other modes of delivering ‘computer reminders’.

**Objectives**—To evaluate the effects on processes and outcomes of care attributable to on-screen computer reminders delivered to clinicians at the point of care.

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AJ participated in screening, data extraction and data analysis.

AM participated in data extraction and screening.

CRR provided support for the analysis.

MPE provided input into overall structure of the review.

JG was involved in the protocol publication and also provided input into the overall structure of the review.

**DECLARATIONS OF INTEREST** ME and JG are authors on one included study and three excluded studies in this review. Four authors (AM, ME, CRR, JG) are editors or staff within the Cochrane EPOC Review Group. Editors and staff are required to conduct at least one Cochrane review. This requirement ensures that editors are aware of the processes and commitment needed to conduct reviews. This involvement does not seem to be a source of conflict of interest in the Cochrane EPOC Review Group. Any editor or staff who is a review author is excluded from editorial decisions on the review in which they are contributors.

**Search methods**—We searched the Cochrane EPOC Group Trials register, MEDLINE, EMBASE and CINAHL and CENTRAL to July 2008, and scanned bibliographies from key articles.

**Selection criteria**—Studies of a reminder delivered via a computer system routinely used by clinicians, with a randomised or quasi-randomised design and reporting at least one outcome involving a clinical endpoint or adherence to a recommended process of care.

**Data collection and analysis**—Two authors independently screened studies for eligibility and abstracted data. For each study, we calculated the median improvement in adherence to target processes of care and also identified the outcome with the largest such improvement. We then calculated the median absolute improvement in process adherence across all studies using both the median outcome from each study and the best outcome.

**Main results**—Twenty-eight studies (reporting a total of thirty-two comparisons) were included. Computer reminders achieved a median improvement in process adherence of 4.2% (interquartile range (IQR): 0.8% to 18.8%) across all reported process outcomes, 3.3% (IQR: 0.5% to 10.6%) for medication ordering, 3.8% (IQR: 0.5% to 6.6%) for vaccinations, and 3.8% (IQR: 0.4% to 16.3%) for test ordering. In a sensitivity analysis using the best outcome from each study, the median improvement was 5.6% (IQR: 2.0% to 19.2%) across all process measures and 6.2% (IQR: 3.0% to 28.0%) across measures of medication ordering.

In the eight comparisons that reported dichotomous clinical endpoints, intervention patients experienced a median absolute improvement of 2.5% (IQR: 1.3% to 4.2%). Blood pressure was the most commonly reported clinical endpoint, with intervention patients experiencing a median reduction in their systolic blood pressure of 1.0 mmHg (IQR: 2.3 mmHg reduction to 2.0 mmHg increase).

**Authors' conclusions**—Point of care computer reminders generally achieve small to modest improvements in provider behaviour. A minority of interventions showed larger effects, but no specific reminder or contextual features were significantly associated with effect magnitude. Further research must identify design features and contextual factors consistently associated with larger improvements in provider behaviour if computer reminders are to succeed on more than a trial and error basis.

### Medical Subject Headings (MeSH)

\*Decision Support Systems, Clinical; \*Outcome and Process Assessment (Health Care); \*Point-of-Care Systems; \*Reminder Systems; Decision Making, Computer-Assisted

### MeSH check words

Humans

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## BACKGROUND

### Description of the condition

Gaps between recommended practice and routine care are widely known (McGlynn 2003; Quality of Health Care 2001; Schuster 1998). Interventions designed to close these gaps fall

into a number of different categories: educational interventions (directed at clinicians or at patients), reminders (again, directed at clinicians or patients), audit and feedback of performance data, case management, and financial incentives to name a few (Shojania 2005). However, none of these categories of interventions confers large improvements in care, especially when evaluated rigorously. In fact, they often produce quite small benefits (Grimshaw 2004; Oxman 1995; Shojania 2006; Walsh 2006) and these benefits tend to involve process measures only, not patient outcomes.

### **Description of the intervention**

Given the difficulty of changing the behaviour of healthcare providers and the resources required by many of the interventions that aim to do so, provider reminders offer a promising strategy, especially given their low marginal cost. Reminders delivered at the point of care prompt healthcare professionals to recall information that they may already know but could easily forget in the midst of performing other activities of care, or, in the case of decision support, provide information or guidance in an accessible format at a particularly relevant time. Paper-based reminders have existed for many years and have ranged from simple notes attached to the fronts of charts (for example reminding providers of the need to administer an influenza vaccine) to more sophisticated pre-printed order forms that include decision support (for example protocols for ordering and monitoring anti-coagulants). Computer-based reminders have the potential to address multiple topics and are automatic; therefore they represent a subset of reminders of great interest to those involved in quality improvement efforts.

### **How the intervention might work**

A number of systematic reviews over the years have evaluated computerised reminders and decision support systems (Dexheimer 2008; Garg 2005; Hunt 1998; Kawamoto 2005). However, these reviews have tended to lump all forms of computerised reminders and decision support together, including, for instance, computer-generated paper reminders and e-mail alerts sent to providers, along with reminders generated at the point of care. It is this last category, computer reminders that prompt providers at the point of care, which represents the most promising form of computerised reminders. Such reminders, embedded into computerised provider order entry systems or electronic medical records, alert providers to important clinical information relevant to a targeted clinical task at the time the provider is engaged in performing the task.

### **Why it is important to do this review**

While point of care computerised reminders have produced some well-known successes (Dexter 2001; Kucher 2005; Overhage 1997), other trials have shown no improvements in care (Ansari 2003; Eccles 2002a; Montgomery 2000), including studies from institutions with well-established computerised order entry systems (Dexter 2004; Sequist 2005; Tierney 2003). Therefore, we sought to quantify the expected magnitudes of improvements in processes and outcomes of care through the use of computerised reminders and decision support delivered at the point of care, and identify any features consistently associated with larger effects.

## OBJECTIVES

In this review, we address the following questions:

1. Do on-screen computer reminders effectively improve processes or outcomes of care?
2. Do any readily identifiable elements of on-screen reminders influence their effectiveness (e.g. inclusion of patient-specific information as opposed to generic reminders for a given condition, requiring a response from users)?
3. Do any readily identifiable elements of the targeted activity (e.g. chart documentation, test ordering, medication prescribing) influence the effectiveness of on-screen reminders?

## METHODS

### Criteria for considering studies for this review

**Types of studies**—We included randomised controlled trials (with randomisation at the level of the patient or the provider) and quasi-randomised trials, where allocation to intervention or control occurred on the basis of an arbitrary but not truly random process (for example even or odd patient identification numbers).

**Types of participants**—Any study in which the majority of participants (> 50%) consisted of physicians or physician trainees; we excluded studies that primarily targeted dentists, pharmacists, nurses, or other health professionals.

**Types of interventions**—The original protocol for this review defined ‘on-screen computer reminders’ as follows:

*Patient or encounter specific information that is provided via a computer console (either visually or audibly) and intended to prompt a healthcare professional to recall information usually encountered through their general medical education, in the medical records or through interaction with peers, and so remind them to perform or avoid some action to aid individual patient care (Gordon 1998).*

This original definition served primarily to distinguish computer reminders that were literally presented to users on a computer screen (hence ‘on-screen reminders’) from computer-generated reminders that were simply printed out and placed in a paper chart. While this distinction remains germane (i.e. some studies still involve ‘computer reminders’ that are really paper-based reminders that happen to have been generated by a computer), the use of computers in healthcare is now sufficiently widespread that the more important concept has become ‘at the point of care’, rather than merely ‘on-screen’. A reminder that is ‘on screen’ but not noticeable to clinicians during the target activities of interest is no more useful than a paper reminder placed in such a manner that clinicians must deviate from their usual charting activities in order to find it.

Thus, from an operational point of view, the focus of this review should be regarded as evaluating ‘point-of-care computer reminders’. By ‘point of care’ we refer to delivery of the

computer reminder to clinicians at the time they are engaged in the target activity of interest, such as prescribing medications, documenting clinical encounters in the medical record, and ordering investigations.

Operationally, we considered a reminder to qualify as delivered at the point of care if the following three criteria applied.

1. The reminder was delivered via the computer system routinely used by the providers targeted by the intervention - typically an electronic medical record or computerised order entry program. For instance, a dedicated computer used solely for performing dose calculations for anticoagulants would not count as 'on-screen/point of care', since it requires clinicians to depart from their usual workflow in order to avail themselves of the reminder or decision support provided by this separate system. We excluded such systems because they in effect require providers to remember to use the reminder system, thus undermining the fundamental purpose of a reminder.
2. The reminder was accessible from within the routinely used clinical information system (typically via a pop-up screen or an icon that indicates the availability of the reminder or decision support feature). A decision support module that could only be accessed by remembering to call up a separate program or website would not count as a point of care reminder (again, because depending on clinicians' remembering to call up the program without any prompting violates the notion of a 'reminder').
3. The reminder targeted the person responsible for the relevant clinical activity. For instance, if handwritten physician orders were entered by a clerk or pharmacist into a computer order entry system, any alert or decision support delivered via the computer system would not qualify as 'point of care' since, for the physician, it was the handwritten order that occurred at the point of care.

For settings without general computer order entry or electronic medical record systems, we allowed the possibility that some specific activities might still routinely occur using a computer system. For instance, an ambulatory clinic might have developed a computer-based system for supporting preventive care activities, even if the rest of the ambulatory record remained paper-based. Or, a hospital might have developed a computer program for ordering certain high-risk drugs (for example chemotherapy or anticoagulants). If a study documented that over 90% of the target activity occurred using the computer system, we regarded such a system as delivering a *de-facto* point of care computer reminder (since the documentation of > 90% use of the computer system for that activity implies that providers would generally not have to remember to use the reminder).

### Types of outcome measures

**Eligible outcomes:** In order to enhance the interpretability of the results, we categorised eligible outcomes as follows.

- *Dichotomous process adherence outcomes:* the percentage of patients receiving a target process of care (e.g. prescription of a specific medication, documentation of

performance of a specific task, such as referral to a consultant) or whose care was in compliance with an overall guideline.

- *Dichotomous clinical outcomes*: true clinical endpoints (such as death or development of a pulmonary embolism), as well as surrogate or intermediate endpoints, such as achievement of a target blood pressure or serum cholesterol level.
- *Continuous clinical outcomes*: various markers of disease or health status (e.g. mean blood pressure or cholesterol level).
- *Continuous process outcomes*: any continuous measure of how providers delivered care (e.g. duration of antibiotic therapy, time to respond to a critical lab value).

We planned to include studies in the analysis only if they reported at least one clinical or process outcome (i.e. we excluded articles that reported only costs, lengths of stay, and other measures of resource use). As it turned out, meaningful analyses were possible only with the measures of process adherence. For these measures, in order to permit pooling across studies, we required that studies present data as the absolute percentage of patients who received the target process care in each study group (or in a manner that allowed us to calculate these percentages). For instance, we would not include a study that only reported the odds of patients receiving the process of care in the intervention group compared with the control. We made this decision partly because initial review revealed that the vast majority of studies reported their data as percentages of patients who received the process of interest, and partly because this format is most conducive to conveying the expected impacts of computer reminders, namely absolute improvements in adherence to a target process of care or clinical behaviour.

**Primary outcomes:** Although we planned to include any otherwise eligible study that reported the effect of computerised reminders on clinical outcomes, evaluating the impact of reminders on adherence to target processes of care represented the primary goal of our analysis. We recognise that improving patient outcomes represents the ultimate goal of any quality improvement activity. However, we focused on process improvements for this review because we wanted to capture the degree to which computer reminders achieve their main goal, namely changing provider behaviour (Mason 1999). The degree to which such behaviour changes ultimately improve patient outcomes will vary depending on the strength of the relationship between the targeted process of interest and patient level outcomes. In some cases, no such relationship may exist. For instance, the incentive to improve appropriate antibiotic use is usually the population level goal of reducing emergence of resistant microorganisms, not improving the outcomes of care for individual patients. In other cases, a presumed relationship between a given process of care and patient outcomes may be incorrect (for example we would no longer expect a reminder that encourages the use of hormone replacement therapy to improve cardiovascular outcomes in post-menopausal women). Consequently, if we had focused on improvements in clinical endpoints and found that reminders achieved negligible improvements in such outcomes, we would not know if this reflected consistent failure of computer reminders to achieve their

intended goal (changes in provider behaviour) or the fact that reminders had targeted processes with limited connections to patient outcomes.

**Direction of improvements:** Some studies target quality problems that involve ‘underuse,’ so that improvements in quality correspond to increases in the percentage of patients who receive a target process of care (for example increasing the percentage of patients who receive the influenza vaccine). However, other studies target ‘overuse,’ so that improvements correspond to reductions in the percentage of patients receiving inappropriate or unnecessary processes of care (for example reducing the percentage of patients who receive antibiotics for viral upper respiratory tract infections). In order to standardise the direction of effects, all process outcomes were defined so that higher values represented an improvement. For example, data from a study aimed at reducing the percentage of patients receiving inappropriate medications would be captured as the complementary percentage of patients who did not receive inappropriate medications. Increasing this percentage of patients for whom providers did not prescribe the medications would thus represent an improvement.

### Search methods for identification of studies

**Electronic searches**—We searched the MEDLINE database up to July 2008 using Medical Subject Headings for relevant forms of clinical information systems (for example Medical Order Entry Systems, Point-of-Care Systems, Ambulatory Care Information Systems) and combinations of text words such as ‘computer’ or ‘electronic’ with terms such as ‘reminder’, ‘prompt’, ‘alert’, ‘cue’, and ‘support’ (Appendix 1 to Appendix 4). We applied a methodological filter for any type of clinical trial. We also searched the EMBASE, CINAHL and CENTRAL databases using modified search strategies up to July 2008. In addition, we retrieved all articles related to computers and reminder systems or decision support from the Cochrane Effective Practice and Organisation of Care Group (EPOC) database (EPOC 2008) Finally, we scanned bibliographies from key articles. For non-English language articles, we screened English translations of titles and abstracts and pursued full-text translation where possible ( i.e. either to include or confirm exclusion).

### Data collection and analysis

**Study selection and data abstraction**—Two investigators (from KS, AJ, AM) independently screened citations and abstracted included articles using a structured data entry form. In the initial screening, authors based their judgments about inclusion and exclusion solely on the titles and abstracts, but promoted articles to the next stage of the screening process whenever a decision could not be made with confidence. For the second stage of screening, we obtained full text for all references, with each article again judged independently by two authors.

Two authors independently abstracted the following information from articles that met all the inclusion criteria after the second stage of screening: clinical setting, participants, methodological details, characteristics of the reminders design and content, the presence of co-interventions (for example educational materials or performance report cards distributed to clinicians in both study groups), and outcomes. The data abstraction form (available upon

request) was based on the checklist developed by the Cochrane EPOC Group (EPOC 2008). The form was pilot tested and revised iteratively prior to its use for final data abstraction. We resolved discrepancies between authors during either the screening or abstraction stages by discussion between the two authors to achieve consensus. When a conflict could not be resolved, a third author was consulted to achieve consensus or generate a majority decision.

**Quality assessment**—As part of the data abstraction process, authors assessed the following quality criteria based on the Cochrane EPOC Group Data Collection Checklist: concealment of allocation, blinded assessment of primary outcomes, proportion of patients/providers followed up, baseline disparities in process adherence or outcomes in the study groups, protection against contamination, and unit of analysis errors (EPOC 2008).

**Data analysis:** We anticipated that the eligible studies would exhibit significant heterogeneity, due to variations in target clinical behaviours, patient and provider populations, methodological features, characteristics of the interventions, and the contexts in which they were delivered. One approach for addressing these sources of variation would involve meta-regression. Given the number of potentially relevant covariates, however, meta-regression would require many more studies than we anticipated finding. We also expected that many eligible studies would assign intervention status to the provider, rather than the patient, but would not take into account ‘cluster effects’ in the analysis (i.e. they would exhibit ‘unit of analysis errors’). Performing either a conventional meta-analysis or meta-regression using studies with unit of analysis errors would require us to make a number of assumptions about the magnitude of unreported parameters, such as the intra-class correlation coefficients and the distributions of patients across clusters, in order to avoid spurious precision in 95% confidence intervals.

To preserve the goal of providing a quantitative assessment of the effects associated with computerised reminders, without resorting to numerous assumptions or conveying a misleading degree of confidence in the results, we chose to report the median improvement in process adherence (and inter-quartile range) among studies that shared specific features of interest. This approach was first developed in a large review of strategies to foster the implementation of clinical practice guidelines (Grimshaw 2004) and subsequently applied to reviews of quality improvement strategies in a series of reports for the US Agency for Healthcare Research and Quality (Shojania 2004a; Shojania 2004b; Steinman 2006; Walsh 2006).

This method of reporting the median effect sizes across groups of studies involves two distinct uses of the term ‘median’. First, in order to handle multiple outcomes within individual studies, we calculated for each study the median improvement in process adherence across the various outcomes reported by that study. For example, if a study reported 10 process adherence outcomes, we would calculate the absolute difference between intervention and control values for each outcome in order to obtain the median improvement (and interquartile range) across all 10 such differences. This median would then contribute the single effect size for that study. We also captured whenever a study identified a primary outcome and separately analysed those studies. Further, we performed a sensitivity analysis in which, instead of the median outcome, we used the best outcome from



each study. With each study then represented by a single, median outcome, we then calculated the median effect size and interquartile range across all included studies. It is this second use of the ‘median’ that is crucial to the method. Instead of providing a conventional meta-analytic mean (an average weighted on the basis of the precision of the results from each study), we highlight the median effect achieved by included studies, along with an interquartile range for these effects.

The main potential drawback of this method of reporting the median effects of an intervention across a group of studies lies in the equal weight given to all studies (for example no weighting occurs on the basis of study precision). Note, however, that by using the median rather than the mean, the summary estimate is less likely to be driven by a handful of outlying results (such as large effects from small or methodologically poor studies). Moreover, we included an analysis of the impact of study size and various other methodological features on reported effect size. For instance, we compared the median effects across large and small studies (where large was defined as greater than or equal to the median sample size across all included studies). We performed the analysis of potential associations between study size and effect magnitude using various measures of sample size, including the numbers of patients (or episodes of care) without any adjustment for clustering, the effective sample size taking into account cluster effects (using values for intra-class correlation coefficients available in the published literature (Campbell 2000)) and, finally, using the numbers of providers (or other cluster units) as the sample size.

We also compared the median effects across studies with and without various methodological markers of study quality, as well as certain features of the study context (for example ambulatory versus inpatient setting) and characteristics of the reminders (for example inclusion of patient-specific information versus a generic alert, provision of an explanation for the reminder, requiring users to enter a response to the reminder before continuing with their work, requiring users to navigate through more than one reminder screen). We made all such comparisons using a non-parametric rank-sum test (Mann-Whitney). We performed all statistical analyses using SAS version 9.1 (SAS Institute, Inc, Cary, NC).

## RESULTS

### Description of studies

See: Characteristics of included studies; Characteristics of excluded studies.

**Results of the search**—Our search identified 2036 citations, of which 1662 were excluded at the initial stage of screening and an additional 374 on full-text review, yielding a total of 28 articles that met all inclusion criteria (Figure 1)(Bates 1999; Christakis 2001; Dexter 2001; Eccles 2002a; Filippi 2003; Flottorp 2002; Frank 2004; Hicks 2008; Judge 2006; Kenealy 2005; Kralj 2003; Krall 2004; Kucher 2005; McCowan 2001; Meigs 2003; Overhage 1996; Overhage 1997; Peterson 2007; Rothschild 2007; Roumie 2006; Safran 1995a; Sequist 2005; Tamblyn 2003; Tape 1993; Tierney 2003; Tierney 2005; van Wyk 2008; Zanetti 2003). Four studies contained two comparisons (Eccles 2002a; Flottorp 2002; Kenealy 2005; van Wyk 2008), resulting in 32 included comparisons.

Of the 32 included comparisons, 19 came from US centers and 24 took place in outpatient settings (see ‘Characteristics of included studies’). Most (26) trials used a true randomised design, with only six comparisons involving a quasi-random design (typically allocating intervention status on the basis of even or odd provider identification numbers). Twenty-six of the 32 included comparisons allocated intervention status at the level of providers or provider groups, rather than allocating patients (i.e. they were cluster trials).

### Risk of bias in included studies

**Allocation**—Of the 32 comparisons in the review, concealed allocation definitely occurred in 14 comparisons (Christakis 2001; Dexter 2001; Flottorp 2002; Frank 2004; Kenealy 2005; McCowan 2001; Meigs 2003; Rothschild 2007; Roumie 2006; Safran 1995a; van Wyk 2008). The process of allocation concealment was unclear in 14 comparisons (Bates 1999; Eccles 2002a; Filippi 2003; Hicks 2008; Judge 2006; Krall 2004; Overhage 1996; Overhage 1997; Peterson 2007; Sequist 2005; Tierney 2003; Tierney 2005; Tamblyn 2003) and not done in four comparisons (Kralj 2003; Kucher 2005; Tape 1993; Zanetti 2003).

**Incomplete outcome data**—The proportion of eligible practices or providers with complete follow up was reported in 14 comparisons (Christakis 2001; Flottorp 2002; Kenealy 2005; Krall 2004; McCowan 2001; Meigs 2003; Overhage 1997; Rothschild 2007; Roumie 2006; Tamblyn 2003; van Wyk 2008). The proportion of eligible patients with complete follow up was reported in 12 comparisons (Filippi 2003; Hicks 2008; Kucher 2005; Meigs 2003; Overhage 1997; Rothschild 2007; Roumie 2006; Safran 1995a; Tamblyn 2003; Tierney 2003; Tierney 2005; Zanetti 2003). The number of subjects (professionals, practices or patients) lost to follow up was not clear in 11 comparisons (Bates 1999; Dexter 2001; Eccles 2002a; Frank 2004; Judge 2006; Kralj 2003; Overhage 1996; Peterson 2007; Sequist 2005; Tape 1993).

**Baseline disparities between study groups**—Only seven comparisons reported data in a format that permitted calculation of baseline disparities between study groups. Across these studies, the median difference between adherence in the intervention and control groups was 0.00% (interquartile range (IQR): 2.0% greater adherence in the control to 0.0%).

**Unit of analysis errors**—Of the 26 comparisons with a clustered design, only 12 analysed their results in a manner that took clustering effects into account. Thus, the remaining 14 clustered comparisons exhibited unit of analysis errors.

**Other quality criteria**—Blinded assessment of study outcomes was generally not relevant, as data were typically derived from electronic systems that documented delivery of the target processes of care. Though not the focus of the review, many of the clinical outcomes were also objective ones, such as laboratory data, and so also did not require blinded assessment.

## Effects of interventions

Of the 32 comparisons that provided analysable results for improvements in process adherence, (Bates 1999; Christakis 2001; Dexter 2001; Eccles 2002a; Filippi 2003; Flottorp 2002; Frank 2004; Hicks 2008; Judge 2006; Kenealy 2005; Kralj 2003; Krall 2004; Kucher 2005; McCowan 2001; Meigs 2003; Overhage 1996; Overhage 1997; Peterson 2007; Rothschild 2007; Roumie 2006; Safran 1995a; Sequist 2005; Tamblyn 2003; Tape 1993; Tierney 2003; Tierney 2005; van Wyk 2008; Zanetti 2003), 21 reported outcomes involving prescribing practices, six specifically targeted adherence to recommended vaccinations, 13 reported outcomes related to test ordering, three captured documentation, and seven reported adherence to miscellaneous other processes (for example composite compliance with a guideline).

Only nine comparisons reported pre-intervention process adherence for intervention and control groups. For these comparisons, the marginal improvement in the intervention (i.e. the median improvement in the intervention group minus the improvement in the control group) was 3.8% (IQR: 0.4% to 7.9%).

Given the small number of studies that reported baseline adherence, improvements attributable to interventions were calculated as the absolute difference in post-intervention adherence (i.e. the post-intervention improvement in the target process of care observed in the intervention group minus that observed in the control group). Using this post-intervention difference between study groups, the median improvements in process adherence associated with computer reminders were: 4.2% (IQR: 0.8% to 18.8%) across all process outcomes, 3.3% (IQR: 0.5% to 10.6%) for improvements in prescribing behaviours, 3.8% (IQR: 0.5% to 6.6%) for improvements in vaccination, and 3.8% (IQR: 0.4% to 16.3%) for test ordering behaviours (Table 1). Table 1 also shows the results obtained when we used the outcome with the largest improvement from each study instead of the outcome with the median improvement.

Eight comparisons reported dichotomous clinical endpoints; intervention patients experienced a median absolute improvement of 2.5% (IQR: 1.3% to 4.2%). These endpoints included intermediate endpoints, such as blood pressure and cholesterol targets, as well as clinical outcomes, such as development of pulmonary embolism and mortality. Blood pressure represented the most commonly reported outcome. Patients in intervention groups experienced a median reduction in their systolic blood pressure of 1.0 mmHg (IQR: 2.3 mmHg reduction to 2.0 mmHg increase). For diastolic blood pressure, the median reduction was 0.2 mmHg (IQR: 0.8 mm reduction to 1.0 mm increase).

## Impacts of study features on effect sizes

There were sufficient comparisons involving process adherence to permit various analyses of potential associations between various study features and the magnitude of effects (Figure 2). The six quasi-randomised controlled trials reported larger improvements in process adherence than the 26 truly randomised comparisons (7.0%, IQR: 1.2% to 28.0% versus 3.4%: IQR 0.6% to 16.3%), but this difference was not statistically significant ( $P = 0.53$ ).

Sample size did not correlate with effect size, whether calculated on the basis of numbers of patients or providers (Figure 2).

One might expect studies with low adherence in control groups to report larger improvements in care, but in fact studies with control adherence rates higher than the median across all studies had a non-significant trend towards larger effect sizes (Figure 2). We analysed the potential impact of baseline adherence in several other ways (for example studies with baseline adherence in top quartile versus all others to look for a 'ceiling effect', and studies with baseline adherence in bottom quartile versus all others to look for a floor effect) but found no indication that baseline adherence significantly affected the magnitude of effect in the intervention group.

Interventions that targeted inpatient settings showed a trend towards larger improvements in processes of care than did those that occurred in outpatient settings: 8.7% (IQR: 2.7% to 22.7%) versus 3.0% (0.6% to 11.5%) for outpatient settings ( $P = 0.34$ ). However, all interventions delivered in inpatient settings occurred at Brigham and Women's Hospital in Boston or the Regenstreif Institute at the University of Indiana. Both of these institutions have mature 'homegrown' computerized provider order entry systems, and the recipients of computer reminders from these institutions consisted primarily of physician trainees, either of which factors may be more relevant than the fact of the inpatient setting.

Studies from the US reported slightly larger improvements in process adherence: 5.0% (IQR: 2.0% to 23.2%) versus 1.2% (IQR: 0.4% to 6.2%) for non-US studies), but this difference was not significant ( $P = 0.12$ ). Moreover, this trend at least partly reflected the results of studies from US institutions with long track records with clinical information systems (for example the Regenstreif Institute and Brigham and Women's Hospital in Boston).

Grouping studies on the basis of track records in clinical informatics (for example analysing studies from Brigham and Women's Hospital, the Regenstreif Institute and Vanderbilt University versus all others) did not result in significant differences, except in the case of Brigham and Women's Hospital. The four studies from Brigham and Women's Hospital by themselves reported significantly higher improvements in process adherence than all other studies: 16.8% (IQR: 8.7% to 26.0%) versus 3.0% (IQR: 0.5% to 11.5%;  $P = 0.04$ ).

Lastly, the magnitude of effects attributable to computer reminders appeared to vary with the presence of co-interventions (delivered to intervention and control groups). The 32 comparisons that reported process adherence outcomes included 18 that evaluated a computer reminder versus usual care and 14 that evaluated a computer reminder plus at least one other quality improvement intervention (for example educational materials) versus this same co-intervention in the control group. Comparisons involving no co-interventions (that is computer reminder alone versus usual care) showed a median improvement in process adherence of 5.7% (IQR: 2.0% to 24.0%), whereas studies of multifaceted interventions (that is computer reminders plus additional interventions versus those additional interventions alone) showed a median improvement in adherence of only 1.9% (IQR: 0.0% to 6.2%;  $P = 0.04$  for this difference).

This apparent difference might reflect a ceiling effect, with co-interventions delivered to the intervention and control groups leaving little room for computer reminders to demonstrate additional improvements. If this were the case, one would expect higher post-intervention adherence rates in the control groups of studies that combined computer reminders with other interventions. However, the opposite proved true: post-intervention values for process adherence (in both intervention and control groups) were in fact slightly higher in the studies involving comparisons of computer reminders by themselves, not in the studies involving additional interventions.

This relationship between comparison type and effect size at least partially reflected confounding by other studies features. For instance, dropping the four studies from Brigham and Women's Hospital from the analysis substantially decreased the magnitude of the difference between studies with and without co-interventions (median improvement of 0.9%, IQR: 0.0% to 5.0% versus 3.8%, IQR: 1.2% to 23.2%), and the difference was no longer statistically significant ( $P = 0.08$ ). Also, of note, none of the  $P$  values reported in the analysis adjusted for multiple comparisons nor was stratification by the presence of co-interventions a prespecified hypothesis for our analysis, further adding to the possibility that the observed difference reflects a chance association.

### Features of computer reminders

We analysed a number of characteristics of the computer reminders (or the larger clinical information system) to look for associations with the magnitude of impact (Figure 3). The degree of improvement did not differ significantly between studies based on the type of quality problem targeted (underuse versus overuse of a given process of care), the conveyance of patient specific information versus a more generic alert, provision of an explanation for the alert, whether or not the reminder conveyed a specific recommendation, whether or not the authors of the study had developed the reminder, or the type of system used to deliver the reminder (CPOE versus electronic medical record).

There was a trend towards larger effects with reminders that required users to enter a response of some kind (12.9%, IQR 2.7% to 22.7%) versus those that did not (2.7%, IQR: 0.6% to 5.6%;  $P = 0.09$ ). However, this trend was confounded by the fact that all four comparisons from Brigham and Women's Hospital involved reminders that required responses from users. Dropping these four studies decreased the median effect of reminders that required user responses to 10.6% (IQR: 0.3% to 21.4%) and removed any appearance of statistical significance ( $P = 0.48$ ). Of note, though, the magnitude of the difference remains substantial (10.6% versus 2.7%); it is possible that the lack of significance reflects lack of power.

We also analysed whether effect sizes differed between reminders that were 'pushed' onto users (that is users automatically received the reminder) versus reminders that required users to perform some action to receive it (that is users had to 'pull' the reminders). Only four comparisons involved 'pull' reminders and these showed comparable effects to 'push' reminders. Of note, however, one trial (van Wyk 2008) directly compared these two modes of reminder delivery. In this three-armed cluster-RCT of reminders for screening and treatment of hyperlipidemia, patients cared for at practices randomised to automatic alerts

were more likely to undergo testing for hyperlipidemia and receive treatment than were patients seen at clinics where reminders were delivered to clinicians only ‘on-demand.’

### Sensitivity analysis

We reanalysed the potential predictors of effect size (study features and characteristics of the reminders) using a variety of alternate choices for the representative outcome from each study, including the outcome with the middle value (rather than a calculated median) and the best outcome (that is the outcome associated with the largest improvement in process adherence). None of these analyses substantially altered the main findings, including the lack of any significant association between study or reminder features and the magnitude of effects achieved by computer reminders. Of note, using the best outcome from each study rather than the median outcome, improvements attributable to reminders in studies at Brigham and Womens Hospital were no longer significantly larger than those achieved in studies from other centers (16.8%, IQR: 8.7% to 26.0% versus 4.6%, IQR: 2.0% to 13.4%;  $P = 0.09$  for the comparison). However, the difference still appears large, so loss of significance may simply reflect the lack of power.

## DISCUSSION

Across 32 comparisons, computer reminders achieved small to modest improvements in care. The absolute improvement in process adherence was less than 4% for half of the included comparisons. Even when we included the best outcome from each comparison, the median improvement was only 5.6%. For improvements in prescribing, perhaps the behaviours of greatest general interest, improvements were even smaller.

With the upper quartile of reported improvements beginning at a 15% increase in process adherence, some studies clearly did show larger effects. However, we were unable to identify any study or reminder features that predicted larger effect sizes, except for a statistically significant (albeit unadjusted for multiple comparisons) difference in effects seen in studies involving the computer order entry system at Brigham and Women’s Hospital. A trend towards larger effects was seen for reminders that required users to enter a response in order to proceed, but this finding may have been confounded by the uneven distribution of studies from Brigham and Women’s Hospital. Thus, we do not know if the success of computer reminders at the Brigham partially reflects the design of reminders requiring user responses or if other features of the computer system or institutional culture of Brigham play the dominant role.

The finding that comparisons of computer reminders alone versus usual reported larger effect sizes than comparisons involving computer reminders and other co-interventions represented an unexpected finding. Exploratory analyses did not reveal a plausible explanation for this result except that it may have reflected uneven distribution of confounders. One additional explanation might be that investigators chose to incorporate computer reminders in multifaceted interventions when attempting to change more complex (and therefore difficult to change) behaviours than those addressed by reminders alone. However, this unexpected finding may also constitute a chance association, especially as none of the  $P$  values reported in the analysis adjust for multiple comparisons.

A major potential limitation of our analysis was the heterogeneity of the interventions and the variable degree with which they were reported, including limited descriptions of key intervention features of the reminders and the systems through which they were delivered. We attempted to overcome this problem by abstracting basic attributes, such as whether user responses were required and whether or not the reminder contained patient-specific information, but heterogeneity within even these apparently straightforward categories could mask important differences in effects. Also, other characteristics which we found difficult to operationalise for example the 'complexity' of the reminder), or which were inadequately reported, may also correlate with important differences in impact. This problem of limited descriptive detail of complex interventions and the resulting potential for substantial heterogeneity among included interventions in systematic reviews has been consistently encountered in the literature (Grimshaw 2003; Ranji 2008; Shojania 2005; Walsh 2006).

Our focus on the median effects across studies represents another potential limitation. However, as outlined in the 'Methods' section, we chose this approach precisely to avoid spurious precision due to heterogeneity and clustering effects that could not be taken into account in many studies. This approach is becoming increasingly common in Cochrane Reviews of interventions to change practice (Grimshaw 2004; Jamtvedt 2006; O'Brien 2007) and has also been used in other evidence syntheses (Grimshaw 2004; Shojania 2004b; Steinman 2006; Walsh 2006). This method conveys the range of effects associated with the intervention of interest and also allows for analysis of factors associated with effect size.

Additional studies continue to appear and we plan to assess eligible new studies formally for inclusion in six months. At that time we will also include a study that had previously been excluded as a time series, but which we have since decided merits inclusion as a controlled clinical trial (Durieux 2000).

In summary, computer reminders delivered at the point of care have achieved variable improvements in target behaviours and processes of care. The small to modest median effects shown in our analysis may hide larger effects. However, the current literature does not suggest which features of the reminder systems, the systems with which they are delivered, or which target problems might consistently predict larger improvements.

## **AUTHORS' CONCLUSIONS**

### **Implications for practice**

On-screen computer reminders may become more prevalent as healthcare institutions advance in the use of computer technology. There appears to be a wide range of effects of the intervention, making it difficult to provide specific suggestions about how to maximize the benefits.

### **Implications for research**

Although some studies have clearly shown substantial improvements in care from point of care computer reminders it is concerning that the majority of studies have shown fairly small improvements across a range of process types. This finding of small to modest improvements is not unique to computer reminders. As had been said before, there are no

‘magic bullets’ when it comes to changing provider behavior and improving care (Shojania 2005; Oxman 1995). However, given that the opportunity to deliver computer reminders at the point of care represents one of the major incentives to implementing sophisticated clinical information systems, future research will need to identify key factors (related to the target quality problem or the design of the reminder) that reliably predict larger improvements in care from these expensive technologies.

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## CHARACTERISTICS OF STUDIES

### Characteristics of included studies *[ordered by study ID]*

#### Bates 1999

Methods	RCT
Participants	Medical & surgical inpatient services, teaching hospital, Boston, USA (Brigham and Women’s Hospital) 11,586 episodes of care
Interventions	Alerting providers to potentially redundant orders for laboratory tests



Outcomes	Process adherence (test ordering), resource use	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	Quote "randomization was performed using an internal identification number not available to providers." Unclear if allocation was concealed from investigators
Incomplete outcome data addressed? All outcomes	Unclear	Not specified in the text

### Christakis 2001

Methods	Cluster-RCT	
Participants	Outpatient pediatric teaching clinic, Seattle, USA (University of Washington) 1339 episodes of care, 38 providers	
Interventions	Displaying evidence regarding the use and duration of antibiotics for otitis media in children	
Outcomes	Process adherence (prescribing)	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Yes	Quote: "stratified randomization using an electronic random number generator"
Incomplete outcome data addressed? All outcomes	Yes	Authors report 92% and 74% complete follow-up for providers for 2 analyses

### Dexter 2001

Methods	Cluster-RCT	
Participants	General medicine inpatient service, teaching hospital, Indianapolis, USA (Indiana University, Regenstrief Institute) 6371 patients, 8 provider teams	
Interventions	Notifying providers of hospitalised patients' eligibility for 4 preventive care measures	
Outcomes	Process adherence (prescribing, vaccination)	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Yes	Quote "a blinded system of coin randomization"
Incomplete outcome data addressed? All outcomes	Unclear	Difficult to assess if all subjects were included

**Eccles 2002a**

Methods	Cluster-RCT	
Participants	Ambulatory general practices, United Kingdom	
	<ul style="list-style-type: none"> <li>a. 2335 patients, 60 practices</li> <li>b. 2363 patients, 60 practices</li> </ul>	
Interventions	Decision support for management of outpatients with (a) angina or (b) asthma	
Outcomes	Process adherence (documentation)	
Notes	System for delivery of reminder: EMR Additional interventions delivered to intervention and control group: distribution of educational materials to providers	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	Not reported in text of study
Incomplete outcome data addressed? All outcomes	Unclear	Not reported

**Filippi 2003**

Methods	Cluster-RCT	
Participants	Ambulatory general practices in Italy 15,343 patients, 300 providers	
Interventions	Identifying diabetic patients who would benefit from antiplatelet agents for cardiovascular risk reduction	
Outcomes	Process adherence (prescribing)	
Notes	System for delivery of reminder: EMR Additional interventions delivered to intervention and control group: distribution of educational materials to providers	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	Not stated in text
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 74.4% calculated from paper for patients

**Flottorp 2002**

Methods	Cluster-RCT	
Participants	Ambulatory general practices in Norway	
	<ul style="list-style-type: none"> <li>a. 9887 episodes of care, 57 practices</li> <li>b. 16,939 episodes of care, 56 practices</li> </ul>	

Interventions	Display of guidelines for appropriate use of antibiotics and laboratory tests in (a) women with suspected urinary tract infection or (b) patients with sore throat
Outcomes	Process adherence (prescribing, test ordering, other)
Notes	System for delivery of care: EMR Additional interventions delivered to intervention and control groups: educational materials for providers and patients, educational workshops for providers, financial incentives for providers

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "142 practices were randomised by computer"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 79.6% for practices calculated from paper

**Frank 2004**

Methods	CCT
Participants	Urban ambulatory practice (academic status not reported), Australia 10,507 patients
Interventions	Notifying providers of outpatients' eligibility for various processes of care (mostly preventive measures)
Outcomes	Process adherence (test ordering, documentation, vaccination)
Notes	System for delivery of care: EMR

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	No	Quote "All patients who attended were randomised by the terminal digit of their record number"
Incomplete outcome data addressed? All outcomes	Unclear	Data not presented for number of patients followed up

**Hicks 2008**

Methods	Cluster-RCT
Participants	Community and hospital-based academically-affiliated primary clinics, Boston, USA (Brigham and Women's Hospital) 1834 patients, 12 clinics
Interventions	Presenting guideline-based suggestions for management of patients with hypertension
Outcomes	Process adherence (guideline-adherent prescribing for hypertension treatment)
Notes	System for delivery of reminder: EMR with e-prescribing

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Unclear	No description of allocation concealment

Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 50% calculated from paper for patients
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### Judge 2006

Methods	Cluster-RCT	
Participants	Academically-affiliated, long-term care facility, Toronto, Canada 3843 episodes of care, 7 wards	
Interventions	Alerting providers to various potential adverse drug events (e.g. severe drug interactions, out-of-recommended-range doses for elderly patients)	
Outcomes	Process adherence (prescribing)	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	No description of allocation concealment
Incomplete outcome data addressed? All outcomes	Unclear	Unable to assess follow-up data

### Kenealy 2005

Methods	Cluster-RCT	
Participants	General practices, New Zealand outpatient 2814 patients, 33 practices	
Interventions	Suggesting to providers that they screen patients over 50 years of age	
Outcomes	Process adherence (test ordering)	
Notes	System for delivery of reminder: EMR Additional interventions delivered to some participants in the intervention and control groups: distribution of educational materials to providers, educational outreach	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Yes	Quote "An independent person used Excel to generate random numbers in blocks of 8 and place the names of intervention groups in sealed and consecutively numbered envelopes"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 94.4% calculated from paper for professionals

### Kralj 2003

Methods	Cluster-CCT	
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Participants	Two community oncology outpatient practices, USA 2170 episodes of care, 2 practices	
Interventions	Prompting providers to order erythropoietin for patients with haemoglobin < 120 g/dL	
Outcomes	Process adherence (prescribing)	
Notes	System for delivery of reminder: EMR with link to CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	No	No description, cluster trial
Incomplete outcome data addressed? All outcomes	Unclear	No description provided

**Krall 2004**

Methods	Cluster-RCT	
Participants	Ambulatory Family and Internal Medicine practices, Oregon, USA (Kaiser Permanente Northwest) 1076 patients, 100 providers	
Interventions	Notification of patients' eligibility to receive aspirin for cardiovascular risk reduction	
Outcomes	Process adherence (prescribing)	
Notes	System for delivery of reminder: EMR and CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	No description of allocation
Incomplete outcome data addressed? All outcomes	Yes	Authors report 100% follow up for professionals

**Kucher 2005**

Methods	CCT	
Participants	Major teaching hospital, Boston, USA (Brigham and Women's Hospital) 2506 patients	
Interventions	Identifying hospitalized patients at increased risk for developing venous thromboembolism	
Outcomes	Process adherence (prescribing, other), clinical outcomes	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	No	CCT, allocation based on medical record number
Incomplete outcome data addressed?	Yes	Follow-up rate of 94.2% calculated from paper for patients

All outcomes

**McCowan 2001**

Methods	Cluster-RCT
Participants	Outpatient general practices, United Kingdom 477 patients, 46 practices
Interventions	Decision support for management of outpatients with asthma
Outcomes	Process adherence (other), clinical outcomes
Notes	System for delivery of reminder: EMR

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "random number sequence"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 37.9% calculated for providers

**Meigs 2003**

Methods	Cluster-RCT
Participants	Internal medicine ambulatory clinic, teaching hospital, Boston, USA (Massachusetts General Hospital) 598 patients, 2 provider teams
Interventions	Display of recommended target goals of care, last known values of relevant lab tests (e. g. HbA1c, creatinine, lipids)
Outcomes	Process adherence (test ordering), clinical outcomes, resource use
Notes	System for delivery of reminder: EMR

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "A coin was tossed to select an intervention group and a control group"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 46.2% for providers and 37.5% for patients calculated from paper

**Overhage 1996**

Methods	Cluster-RCT
Participants	Inpatient internal medicine service, academic medical center, Indianapolis, USA (Regenstrief Institute) 1622 patients, 24 providers
Interventions	Suggesting orders for various preventive care measures in eligible patients
Outcomes	Process adherence (prescribing, test ordering, vaccination, other)

Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	No description of allocation concealment
Incomplete outcome data addressed? All outcomes	Unclear	Unclear reporting of total number of patients

### Overhage 1997

Methods	Cluster-RCT	
Participants	Medicine service, teaching hospital, Indianapolis, USA (Regenstrief Institute) 2181 patients, 6 services	
Interventions	Prompting providers about "corollary orders"	
Outcomes	Process adherence (test ordering), clinical outcomes, resource use	
Notes	System for delivery of reminder: CPOE Additional interventions delivered to intervention and control groups: drug utilization review program	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	No description of allocation concealment
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 93.5% for professionals and 100% for patients calculated from paper

### Peterson 2007

Methods	CCT	
Participants	Academic medical center, Nashville, USA (Vanderbilt University) 2981 patients	
Interventions	Providing medication decision support for drugs in the hospitalized elderly (avoiding certain drugs and modified dosing for others)	
Outcomes	Process adherence (medication dosage and criteria)	
Notes	System for delivery of reminder: CPOE	
<b>Risk of bias</b>		
<b>Item</b>	<b>Authors' judgement</b>	<b>Description</b>
Allocation concealment?	Unclear	No description of allocation concealment
Incomplete outcome data addressed? All outcomes	Unclear	Information not provided

### Rothschild 2007

Methods	Cluster-RCT
Participants	Academic medical center, Boston, USA (Brigham and Women's Hospital) 350 episodes of care, 453 providers
Interventions	Presentation of guidelines regarding indications for transfusion of red cells, platelets, and frozen plasma
Outcomes	Process adherence (prescribing), clinical outcomes
Notes	System for delivery of reminder: CPOE Additional interventions delivered to intervention and control groups: provider education (printed materials, workshops)

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "A computerized program generated the randomization scheme for each block"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 100% for both patients and providers calculated from paper

**Roumie 2006**

Methods	Cluster-RCT
Participants	2 hospitals, 8 ambulatory clinics, Nashville, USA (Vanderbilt University) 871 patients, 116 providers
Interventions	Alert in electronic medical record displaying recent blood pressure value and outlining national recommendations for hypertension treatment and blood pressure goals
Outcomes	Process adherence (prescribing), clinical outcomes
Notes	System for delivery of reminder: EMR Additional interventions delivered to intervention and control groups: provider education (printed materials delivered via e-mail)

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "randomization sequence was computer-generated"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 88% for providers and 77% for patients calculated from paper

**Safran 1995a**

Methods	Cluster-RCT
Participants	Academic primary care clinic, Boston, USA (Beth Israel Hospital) 349 patients, 2 teams
Interventions	Alerting providers to eligibility of HIV-positive patients for various recommended processes of care
Outcomes	Process adherence (other), process outcomes, clinical outcomes
Notes	System for delivery of reminder: EMR



*Risk of bias*

Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "by flipping a coin"
Incomplete outcome data addressed? All outcomes	Unclear	Follow-up rate of 51.3% for patients calculated from paper

**Sequist 2005**

Methods	Cluster-RCT
Participants	Outpatient primary clinics (academic and community) Boston area, USA (Brigham and Women's Hospital) 6243 patients, 20 clinics
Interventions	Displaying guidelines for recommended aspects of care for patients with diabetes and coronary artery disease
Outcomes	Process adherence (prescribing, other)
Notes	System for delivery of reminder: EMR Additional interventions delivered to intervention and control groups: paper reminders to providers

*Risk of bias*

Item	Authors' judgement	Description
Allocation concealment?	Unclear	Not described in paper
Incomplete outcome data addressed? All outcomes	Unclear	Insufficient information to assess in paper

**Tamblyn 2003**

Methods	Cluster-RCT
Participants	Primary care practices in Quebec, Canada 12,560 encounters, 107 providers
Interventions	Alerting providers to various potential adverse drug events (e.g. based on drug-drug interactions, and drug-disease or drug-age contraindications)
Outcomes	Process adherence (prescribing)
Notes	System for delivery of reminder: CPOE

*Risk of bias*

Item	Authors' judgement	Description
Allocation concealment?	Unclear	Not described in paper
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 100% for providers and 100% for patients calculated from paper

**Tape 1993**

Methods	Cluster-CCT
Participants	Internal medicine teaching clinic, Omaha, USA (University of Nebraska) 1809 patients, 2 clinics
Interventions	Drawing attention to deficiencies in preventive care measures for a given patient
Outcomes	Process adherence (test ordering, vaccination)
Notes	System for delivery of reminder: EMR Additional interventions delivered to intervention and control groups: provider education (conferences), paper reminders to providers

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	No	Alternate week randomisation
Incomplete outcome data addressed? All outcomes	Unclear	Insufficient information to assess in paper

**Tierney 2003**

Methods	Cluster-RCT
Participants	Academic primary care group practice, Indianapolis, USA (Regenstrief Institute) 378 encounters, 32 practice sessions
Interventions	Presenting guideline-based suggestions for management of heart failure and coronary artery disease
Outcomes	Process adherence (prescribing, vaccination), clinical outcome, resource use
Notes	System for delivery of reminder: CPOE Additional interventions delivered to intervention and control groups: provider (printed materials, workshops, outreach visits), use of local opinion leaders

**Risk of bias**

Item	Authors' judgement	Description
Allocation concealment?	Unclear	Quote "We used the random number generator...", but unclear if procedure was followed for the whole sample
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 81% for patients calculated from paper

**Tierney 2005**

Methods	RCT
Participants	General medicine practice, teaching hospital, Indianapolis, USA (Regenstrief Institute) 363 episodes of care
Interventions	Presenting guideline-based suggestions for management of asthma and chronic obstructive pulmonary disease
Outcomes	Process adherence (prescribing, test ordering, vaccination), clinical outcome, resource use
Notes	System for delivery of reminder: CPOE

Additional interventions delivered to intervention and control groups: provider education (printed materials and workshops)

<i>Risk of bias</i>		
Item	Authors' judgement	Description
Allocation concealment?	Unclear	Quote "We randomized all half-day sessions to intervention or control status via a coin flip" but adjustment to allocation performed and whether the adjustment was concealed
Incomplete outcome data addressed? All outcomes	Unclear	Follow-up rate of 89.9% for patients calculated from paper

### van Wyk 2008

Methods	Cluster-RCT
Participants	General practice clinics in the Netherlands 3955 patients, 24 clinics
Interventions	Automatic presentation of patient-specific recommendations related to screening and treatment of patients with dyslipidemia
Outcomes	Process adherence (screening, treating)
Notes	System for delivery of reminder: EMR

<i>Risk of bias</i>		
Item	Authors' judgement	Description
Allocation concealment?	Yes	Quote "Randomization was performed with a table of random numbers"
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 95% of practices calculated from paper.

### Zanetti 2003

Methods	CCT
Participants	Cardiac surgery service at academic medical center, Boston, USA (Brigham and Women's Hospital) 449 operations
Interventions	Audible alarm on operating room computer with accompanying visual alert that patient should receive a second dose of antibiotic prophylaxis due to prolonged operative time
Outcomes	Process adherence (prescribing), clinical outcomes
Notes	System for delivery of reminder: CPOE

<i>Risk of bias</i>		
Item	Authors' judgement	Description
Allocation concealment?	No	Assignment based on case number
Incomplete outcome data addressed? All outcomes	Yes	Follow-up rate of 82% for patients calculated from paper

CPOE: Computerised provider order entry; EMR: electronic medical record; RCT: randomised controlled trial; CCT: Controlled clinical trial.

## Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Aase 1999	Excluded topic
Ageno 1998	Not a provider reminder
Anaya 2004	Not a provider reminder
Andersen 2002	Not a provider reminder
Anderson 1976	Excluded topic
Anderson 2003	Not a provider reminder
Anonymous 2002	Not an on-screen computer reminder
Ansari 2003	Ineligible comparison or inappropriate control
Apkon 2005	Excluded topic
Aronsky 2001	Excluded topic
Ashe 2004	Not an on-screen computer reminder
Atherton-Naji 2001	Not a provider reminder
Bailey 2007	Not an on-screen computer reminder
Baird 1984	Not a provider reminder
Barnett 1983	Not an on-screen computer reminder
Baskerville 2001	Not a provider reminder
Bates 1997	No eligible outcomes
Beaulieu 2004	Not an on-screen computer reminder
Becker 1989	Not an on-screen computer reminder
Bekkering 2005	Excluded topic
Bennett 2003	Study design
Berkenstadt 2006	Excluded topic
Bindels 2000	Study design
Bindels 2003	Not an on-screen computer reminder
Bindels 2004	Study design
Bird 1990	Not an on-screen computer reminder
Bjorsness 2003	Not an on-screen computer reminder
Black 2002	Not a provider reminder
Boekeloo 1990	Not an on-screen computer reminder
Bogucki 2004	Study design
Bomba 2006	Not a provider reminder
Bonevski 1999	Not a provider reminder
Borlawsky 2005	Study design
Bosworth 2005	Data not interpretable
Bovbjerg 2000	Not a provider reminder
Brown 2003	Excluded topic
Brownbridge 1986	Study design
Buchsbaum 1993	Not an on-screen computer reminder

<b>Study</b>	<b>Reason for exclusion</b>
Bullard 2004	Not a provider reminder
Burack 1989	Not an on-screen computer reminder
Burack 2003	Not an on-screen computer reminder
Burton 1991	Not a provider reminder
Cabrero-Canosa 2004	Excluded topic
Calabrisi 2002	Study design
Cannon 2000	Not a provider reminder
Care Management 2002	Not a provider reminder
Casebeer 2003	Not a provider reminder
Cho 2003	Not a provider reminder
Christakis 2004	No eligible outcomes
Christensen 2004	Not an on-screen computer reminder
Clayton 2007	Not a provider reminder
Coe 1977	Not an on-screen computer reminder
Collins 2003	Not an on-screen computer reminder
Colombet 2004	Study design
Cooke 2001	Not a provider reminder
Cornia 2003	Not a provider reminder
Costanza 2000	Not a provider reminder
Coyle 2004	Not an on-screen computer reminder
Cramer 1999	Not a provider reminder
Crane 2006	Not an on-screen computer reminder
Creedon 1998	Not a provider reminder
Curry 2003	Data not interpretable
Daley 2004a	Not a provider reminder
Daley 2004b	Not an on-screen computer reminder
Daugaard 2002	Not a provider reminder
Davis 2005	Study design
Day 2006	Excluded topic
de Dombal 1994	Study design
de Fine Olivarius 2001	Not a provider reminder
Degia 2005	Not a provider reminder
Degli 1999	Not a provider reminder
del Mar 2006	Duplicate publication
Demakis 2000	Ineligible comparison or inappropriate control
Derose 2005	Not an on-screen computer reminder
Desbiens 2000	Not an on-screen computer reminder
Dexter 1998	Not an on-screen computer reminder
Dexter 2004	Ineligible comparison or inappropriate control

<b>Study</b>	<b>Reason for exclusion</b>
Dowie 2002	Not a provider reminder
East 1999	Not a provider reminder
Ebell 1995	Study design
Eccles 2000	Data not interpretable
Eccles 2001	Not an on-screen computer reminder
Eccles 2002	Not an on-screen computer reminder
Edmonson 2005	Not a provider reminder
Emery 2007	Excluded topic
Evans 1994	No eligible outcomes
Faresjo 2006	Not a provider reminder
Feeg 2005	Not a provider reminder
Feldstein 2006a	Not an on-screen computer reminder
Feldstein 2006b	Not an on-screen computer reminder
Fihn 1994	Not a provider reminder
Fiks 2007	Study design
Fischer 2003	Study design
Fitzmaurice 1996	Not a provider reminder
Fitzmaurice 2000	Not an on-screen computer reminder
Flanagan 1999a	Not an on-screen computer reminder
Flanagan 1999b	Study design
Flottorp 2003a	Not a provider reminder
Flottorp 2003b	Study design
Fordham 1990	Not a provider reminder
Frame 1994	Not an on-screen computer reminder
Frances 2001	Ineligible comparison or inappropriate control
Fransen 2004	Not a provider reminder
Fretheim 2003	Study design
Fretheim 2006a	Ineligible comparison or inappropriate control
Fretheim 2006b	Duplicate publication
Frolich 1997	Not an on-screen computer reminder
Galanter 2005	Study design
Garrett 2002	Not a provider reminder
German 1987	Not an on-screen computer reminder
Gill 2000	Not an on-screen computer reminder
Glasgow 1995	Not an on-screen computer reminder
Goff 2003	Not an on-screen computer reminder
Goldberg 2000	Study design
Goldberg 2002	Not an on-screen computer reminder
Goodey 2000	Not an on-screen computer reminder

<b>Study</b>	<b>Reason for exclusion</b>
Grimaud 1999	Not a provider reminder
Hales 1995	Not an on-screen computer reminder
Harpole 1997	Ineligible comparison or inappropriate control
Haruna 1998	Not a provider reminder
Hatcher 2005	Study design
Heidenreich 2005	Not an on-screen computer reminder
Heiman 2004	Not an on-screen computer reminder
Hetlevik 1998	No eligible outcomes
Hetlevik 1999	No eligible outcomes
Hetlevik 2000	Ineligible comparison or inappropriate control
Hickling 1989	Not a provider reminder
Hobbs 1996	Not a provider reminder
Hoch 2003	Not an on-screen computer reminder
Hoepfer 1984	Not a provider reminder
Hogg 1990	Not an on-screen computer reminder
Holbrook 2002	Study design
Holbrook 2005	Data not interpretable
Hollingworth 1995	Excluded topic
Hornig 2006	Not a provider reminder
Hsieh 2004	Study design
Javitt 2005	Not an on-screen computer reminder
Jenkins 2000	Study design
John 2008	Study design
Johnson 2004	Excluded topic
Johnson 2006	Not a provider reminder
Johnston 2004	Not a provider reminder
Junghans 2007	Not a provider reminder
Kailajärvi 2000	Study design
Kaplan 2000	Not a provider reminder
Karikoski 2003	Not a provider reminder
Katz 2004	Not an on-screen computer reminder
Keefe 2005	No eligible outcomes
Kellett 1997	Not an on-screen computer reminder
Koop 2002	Not a provider reminder
Kuilboer 2006	Data not interpretable
Kuperman 1999	Not an on-screen computer reminder
Kuperman 2001	Study design
Lafata 2007	Ineligible comparison or inappropriate control
Laflamme 2005	Not a provider reminder

<b>Study</b>	<b>Reason for exclusion</b>
Landis 1992	Not an on-screen computer reminder
Lattimer 1998	Not a provider reminder
Lattimer 2000	Not a provider reminder
Lau 1998	Study design
LeBaron 2004	Not a provider reminder
Lemelin 2001	Not a provider reminder
Lesourd 2002	Not an on-screen computer reminder
Leung 2003	Not a provider reminder
Lewis 1996	Not an on-screen computer reminder
Lilford 1992	Not a provider reminder
Lin 2004	Not a provider reminder
Lipkus 1999	Not an on-screen computer reminder
Litzelman 1993	Not an on-screen computer reminder
Lobach 1997	Not an on-screen computer reminder
Lobach 1994	Not an on-screen computer reminder
Lobach 1996	Not a provider reminder
Love 2008	Data not interpretable
Lowensteyn 1998	Not an on-screen computer reminder
Lubitz 1995	Not a provider reminder
MacIntyre 2003	Not an on-screen computer reminder
MacLean 2004	Not an on-screen computer reminder
Margolis 1992	Data not interpretable
Margolis 2004	Not a provider reminder
Martens 2006	Data not interpretable
Martens 2007	Data not interpretable
Martin 2007	Not an on-screen computer reminder
Matheny 2007	Not an on-screen computer reminder
McAlister 2006	Not an on-screen computer reminder
McCartney 1997	Not an on-screen computer reminder
McDonald 1976	Not an on-screen computer reminder
McDonald 1980	Not an on-screen computer reminder
McDonald 1984	Not an on-screen computer reminder
McDonald 1992	Not an on-screen computer reminder
McDowell 1986	Not a provider reminder
McDowell 1989a	Not a provider reminder
McDowell 1989b	Not an on-screen computer reminder
McEwen 2006	Not an on-screen computer reminder
McGowan 1999	Excluded topic
McGowan 2000	Not a provider reminder



<b>Study</b>	<b>Reason for exclusion</b>
McGregor 2006	Not an on-screen computer reminder
McIsaac 2002	Not an on-screen computer reminder
McKinley 2001	Not an on-screen computer reminder
McPhee 1989	Not an on-screen computer reminder
McPhee 1991	Not an on-screen computer reminder
Medow 2001	Excluded topic
Mills 2006	Not a provider reminder
Montani 2000	Not a provider reminder
Montgomery 2000	Ineligible comparison or inappropriate control
Montori 2002	Not a provider reminder
Morisky 2002	Not a provider reminder
Murray 1999	Not an on-screen computer reminder
Murray 2004	Duplicate publication
Musser 2001	Not a provider reminder
Myers 2004	Not an on-screen computer reminder
Nguyen 2000	Not an on-screen computer reminder
Nilasena 1995	Not an on-screen computer reminder
Norman 2004	Not a provider reminder
Olivarius 2001	Not an on-screen computer reminder
Oppenheim 2002	Study design
Ornstein 1991	Not an on-screen computer reminder
Ornstein 2001	Not an on-screen computer reminder
Otero-Sabogal 2006	Not an on-screen computer reminder
Overhage 1995	Study design
Overhage 2001	Not a provider reminder
Oyen 2005	Study design
Palen 2006	Ineligible comparison or inappropriate control
Parry 2000	Duplicate publication
Patterson 1998	Not an on-screen computer reminder
Peck 1973	Not a provider reminder
Persell 2008	No eligible outcomes
Petrucci 1991	Excluded topic
Poller 1993	Not an on-screen computer reminder
Poller 1998	Not a provider reminder
Preston 2000	Not an on-screen computer reminder
Raebel 2005	Excluded topic
Raebel 2007a	Excluded topic
Raebel 2007b	Excluded topic
Ramnarayan 2003	Excluded topic

<b>Study</b>	<b>Reason for exclusion</b>
Rapley 2006	Not a provider reminder
Reeve 2008	Excluded topic
Rhodes 2006	Not an on-screen computer reminder
Rind 1991	Study design
Rind 1995	Study design
Robinson 2007	Not a provider reminder
Rodman 1984	Not a provider reminder
Rogers 1982	Not an on-screen computer reminder
Rogers 1984	Not an on-screen computer reminder
Rollman 2001	Not an on-screen computer reminder
Rollman 2002	Not an on-screen computer reminder
Rood 2005	Excluded topic
Rosenman 2003	Study design
Rosser 1991	Not an on-screen computer reminder
Rosser 1992	Not an on-screen computer reminder
Rotman 1995	Data not interpretable
Rotman 1996	No eligible outcomes
Roukema 2008	Excluded topic
Roumie 2004	Study design
Roumie 2007	Duplicate publication
Rubenstein 1995	Not an on-screen computer reminder
Ryff-de Leche 1919	Not a provider reminder
Sabnis 2003	Not an on-screen computer reminder
Safran 1993	Duplicate publication
Safran 1995b	Duplicate publication
Safran 1996	Duplicate publication
Saitz 2003	Not an on-screen computer reminder
Samore 2005	Not an on-screen computer reminder
Sanders 2000	Not an on-screen computer reminder
Scharer 2002	Not a provider reminder
Scheel 2002	Not an on-screen computer reminder
Scholes 2006	Not an on-screen computer reminder
Schriger 1997	Study design
Schriger 2001	Not an on-screen computer reminder
Schwartz 2005	Not a provider reminder
Schwid 1999	Excluded topic
Shandro 2002	Not a provider reminder
Shannon 2001	Not an on-screen computer reminder
Shapiro 1987	Not a provider reminder

<b>Study</b>	<b>Reason for exclusion</b>
Shea 1995	No eligible outcomes
Shojania 1998	No eligible outcomes
Shu 2001	Not an on-screen computer reminder
Sicotte 1998	Not a provider reminder
Simon 2000	Not an on-screen computer reminder
Simon 2006	Ineligible comparison or inappropriate control
Sittig 1990	Study design
Smith 2008	Not an on-screen computer reminder
Smithuis 1994	Non-English
Solomon 2001	Not a provider reminder
Sommers 1984	Not an on-screen computer reminder
Stair 1995	Not a provider reminder
Stamos 2001	Not an on-screen computer reminder
Steele 2003	Not a provider reminder
Stewart 2003	Study design
Stone 2005	Not a provider reminder
Subramanian 2004	Not an on-screen computer reminder
Séroussi 2004	Study design
Tai 1999	Not a provider reminder
Tamblyn 1997	Duplicate publication
Tang 1999	Study design
Taylor 2008	No eligible outcomes
Thomas 1983	Not an on-screen computer reminder
Thomas 2007	Not a provider reminder
Thompson 1999	Not a provider reminder
Tierney 1986	Not an on-screen computer reminder
Tierney 1987	No eligible outcomes
Tierney 1988	No eligible outcomes
Tierney 1990	No eligible outcomes
Tierney 1993	No eligible outcomes
Topal 2005	Study design
Topol 2007	Study design
Torgerson 1993	Not a provider reminder
Toth-Pal 2004	Study design
Tremaine 2001	Not a provider reminder
Trivedi 2003	Study design
Trivedi 2007	Not a provider reminder
Tung 2003	Study design
Turner 1989	Not an on-screen computer reminder

Study	Reason for exclusion
Vadher 1997a	Not an on-screen computer reminder
Vadher 1997b	Not a provider reminder
van den Besselaar 2007	Not an on-screen computer reminder
van Overbeeke 1996	Not a provider reminder
van Wyk 2003	Not a provider reminder
Verstappen 2007	Not an on-screen computer reminder
Vesely 2006	Excluded topic
Vinker 2002	Not an on-screen computer reminder
Vissers 1995	Not an on-screen computer reminder
Vissers 1996a	Not an on-screen computer reminder
Vissers 1996b	Not an on-screen computer reminder
Walton 1997	Study design
Wanger 1997	Study design
Warren 1999	Excluded topic
Weaver 2004	Study design
Weber 2008	Not an on-screen computer reminder
Weingarten 1989	Not an on-screen computer reminder
Weir 2003	Not an on-screen computer reminder
Wellingham 2003	Study design
White 1987	Not a provider reminder
White 1991	Not a provider reminder
Whitty 2004	Study design
Williams 1998	Not an on-screen computer reminder
Wolfenden 2005	Data not interpretable
Wright 2007	Not an on-screen computer reminder
Yarnall 1998	Not an on-screen computer reminder
Yealy 2005	Not an on-screen computer reminder
Young 1981	Not an on-screen computer reminder
Young 2003	Not a provider reminder
Ziemer 20056	Not an on-screen computer reminder
Zimmerman 2003	Not a provider reminder
Zwar 2002	Not an on-screen computer reminder

### Characteristics of studies awaiting assessment [ordered by study ID]

#### Durieux 2000

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Methods

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Participants
Interventions
Outcomes
Notes

**Plaza 2005**

Methods
Participants
Interventions
Outcomes
Notes

Study published in Spanish - awaiting translation. Expected to be eligible for inclusion.

**DATA AND ANALYSES**

This review has no analyses.

**HISTORY**

Protocol first published: Issue 2, 1998

Review first published: Issue 3, 2009

Date	Event	Description
11 November 2009	Amended	Minor changes to figures

**WHAT'S NEW**

Last assessed as up-to-date: 10 January 2009.

Date	Event	Description
7 December 2010	Amended	Minor typo change to title

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- \* Indicates the major publication for the study

## PLAIN LANGUAGE SUMMARY

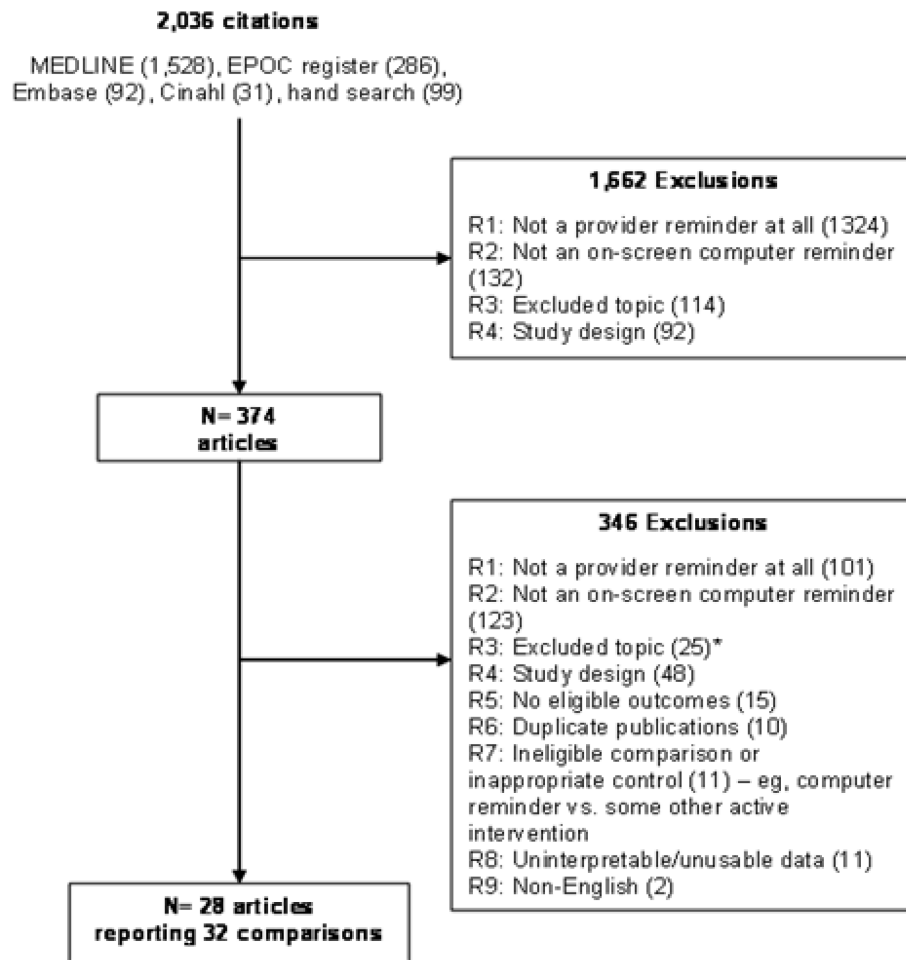
### **On screen point of care computer reminders to improve care and health**

It is known that doctors do not always provide the care that is recommended or according to the latest research. Many strategies have been tried in an attempt to reduce this gap between what is recommended and what is done. A potentially low cost way to do this could be to use computer systems that remind physicians about important information while they make decisions. For example, a doctor could be ordering antibiotics for a child with an ear infection. At that point, the computer the doctor is working on displays a pop up window with a reminder about the evidence for the best dose and length of time the antibiotics should be prescribed.

This review found 28 studies that evaluated the effects of different on-screen computer reminders. The studies tested reminders to prescribe specific medications, to warn about drug interactions, to provide vaccinations, or to order tests. The review found small to moderate benefits. The reminders improved physician practices by a median of 4%. In eight of the studies, patients' health improved by a median of 3%.

Although some studies showed larger benefits than these median effects, no specific reminders or features of how they worked were consistently associated with these larger benefits. More research is needed to identify what types of reminders work and when.

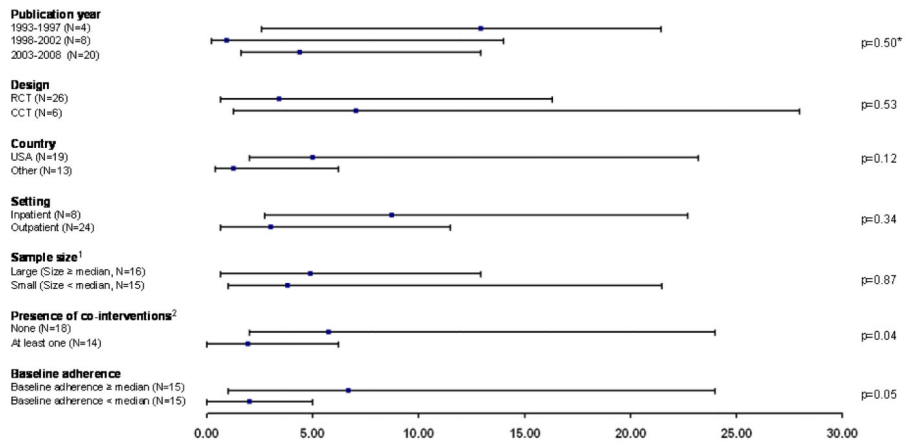




\* Excluded topics included:

- 1) 'expert systems' (e.g., artificial intelligence or neural network applications) for facilitating diagnosis (such as decision support for diagnostic reasoning or interpretation of imaging studies) or for estimating prognosis;
- 2) decision support not directly related to patient care (e.g., coding medical records);
- 3) reminders directed primarily at non-physicians. Only three articles were excluded solely on this basis—two involving pharmacists and one involving nurses.

**Figure 1.**  
Results of literature search and application of eligibility criteria

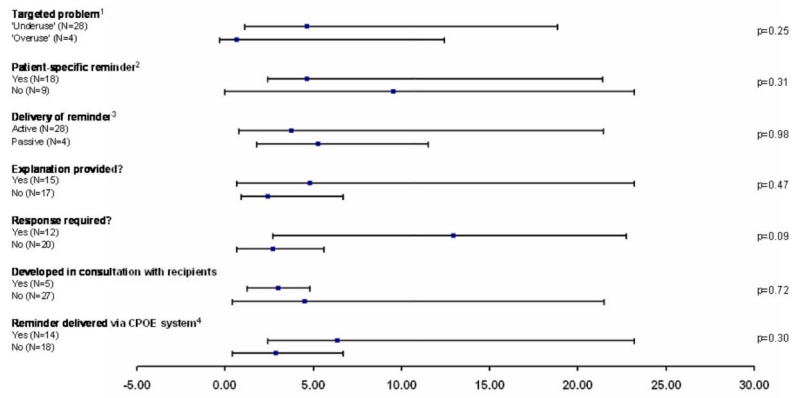


\* Kruskal-Wallis test

<sup>1</sup> Results for sample size used raw numbers of patients, without adjusting for cluster effects. Adjusting for clustering did not alter the results, nor did using the number of clusters instead of the number of patients. The total number of comparisons for the sample size analysis is 31, because 1 study did not report number of patients.

<sup>2</sup> Result compares the effect sizes of comparisons involving computer reminders alone versus usual care (i.e. no co-interventions) and reminders plus one or more other quality improvement interventions versus those other interventions alone.

**Figure 2.**  
Median effects for process adherence by study feature



<sup>1</sup>Underuse refers to targeting quality improvements that correspond to increasing the percentage of patients who receive a target process of care (e.g. increasing the percentage of patients receiving the influenza vaccine). Overuse refers to targeting improvements that correspond to reductions in the percentage of patients receiving inappropriate care (e.g. reducing the percentage of patients who receive antibiotics for viral upper respiratory tract infections).

<sup>2</sup>Reminders with no patient specific information were those triggered on the basis of demographic triggers (e.g. age) or the intent to order a medication or investigation irrespective of any features of the patient involved (e.g. a reminder triggered by any order for heparin or any request for a certain radiologic investigation), as opposed to patient-specific laboratory results (e.g. a reminder related to the patient's serum creatinine) or combinations of medications or laboratory values exhibited by the patient. The sample size is reduced due to inability to accurately assess the presence or absence of the feature.

<sup>3</sup>Active reminders were those that appeared automatically when triggering conditions were met, as opposed to passive reminders, where, for instance, users might be presented with the option to click on a link to receive decision support related to their current task. In some informatics context, this distinction is referred to as "push" vs. "pull".

<sup>4</sup>CPOE – computerized order entry system, reminders systems without CPOE were typically electronic medical record systems.

**Figure 3.**  
Median effects for process adherence by reminder feature

**Table 1**

Median improvements in process adherence across included studies

Dichotomous outcomes (number of intervention vs. control comparisons)	Median absolute improvement (Interquartile range)	
	Using median outcome from each study	Using best outcome from each study
All process outcomes (N = 32)	4.2% (0.8% to 18.8%)	5.6% (2.0% to 19.2%)
Prescription of medications (N = 21)	3.30% (0.5% to 10.6%)	6.2% (3.0% to 28.0%)
Prescription of recommended vaccines (N = 6)	3.8% (0.5% to 6.6%)	4.8% (0.5% to 7.8%)
Test ordering (N = 13)	3.8% (0.4% to 16.30%)	9.6% (0.6% to 24.0%)
Elements of recommended documentation (N = 3)	0.0% (-1.0% to 1.3%)	2.0% (2.0% to 4.0%)
Other process outcomes (N = 7)	1.0% (0.8% to 8.5%)	4.0% (0.8% to 8.5%)

The Table shows average improvements (expressed as the median and interquartile range) across included comparisons for different types of process outcomes. All process outcomes were defined so that higher values always represent an improvement. For example, data from a study aimed at reducing the percentage of patients receiving inappropriate medications would be captured as the complementary percentage of patients receiving appropriate medications, so that an increase in process adherence would represent an improvement.

Most studies reported multiple endpoints but did not specify a primary outcome. For the main analyses, we used the median improvement from each study (that is the median change in adherence to a target guideline or process of care across all such changes reported for the study) as the single representative outcome for that study. We then calculated the median improvements across all included studies for different types of process measures, as shown in the middle column of the table. The column to the far right presents the same results when we used the best improvement from each study as its representative outcome.