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Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)

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Continuing education meetings and workshops: effects on professional practice and health care outcomes

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ABSTRACT

Background
Educational meetings are widely used for continuing medical education. Previous reviews found that interactive workshops resulted in moderately large improvements in professional practice, whereas didactic sessions did not.

Objectives
To assess the effects of educational meetings on professional practice and healthcare outcomes.

Search strategy
We updated previous searches by searching the Cochrane Effective Practice and Organisation of Care Group Trials Register and pending file, from 1999 to March 2006.

Selection criteria
Randomised controlled trials of educational meetings that reported an objective measure of professional practice or healthcare outcomes.

Data collection and analysis
Two authors independently extracted data and assessed study quality. Studies with a low or moderate risk of bias and that reported baseline data were included in the primary analysis. They were weighted according to the number of health professionals participating. For each comparison, we calculated the risk difference (RD) for dichotomous outcomes, adjusted for baseline compliance; and for continuous outcomes the percentage change relative to the control group average after the intervention, adjusted for baseline performance. Professional and patient outcomes were analysed separately. We considered 10 factors to explain heterogeneity of effect estimates using weighted meta-regression supplemented by visual analysis of bubble and box plots.

Main results

[Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)]
In updating the review, 49 new studies were identified for inclusion. A total of 81 trials involving more than 11,000 health professionals are now included in the review. Based on 30 trials (36 comparisons), the median adjusted RD in compliance with desired practice was 6% (interquartile range 1.8 to 15.9) when any intervention in which educational meetings were a component was compared to no intervention. Educational meetings alone had similar effects (median adjusted RD 6%, interquartile range 2.9 to 15.3; based on 21 comparisons in 19 trials). For continuous outcomes the median adjusted percentage change relative to control was 10% (interquartile range 8 to 32%; 5 trials). For patient outcomes the median adjusted RD in achievement of treatment goals was 3.0 (interquartile range 0.1 to 4.0; 5 trials). Based on univariate meta-regression analyses of the 36 comparisons with dichotomous outcomes for professional practice, higher attendance at the educational meetings was associated with larger adjusted RDs (P < 0.01); mixed interactive and didactic education meetings (median adjusted RD 13.6) were more effective than either didactic meetings (RD 6.9) or interactive meetings (RD 3.0). Educational meetings did not appear to be effective for complex behaviours (adjusted RD -0.3) compared to less complex behaviours; they appeared to be less effective for less serious outcomes (RD 2.9) than for more serious outcomes.

**Authors’ conclusions**

Educational meetings alone or combined with other interventions, can improve professional practice and healthcare outcomes for the patients. The effect is most likely to be small and similar to other types of continuing medical education, such as audit and feedback, and educational outreach visits. Strategies to increase attendance at educational meetings, using mixed interactive and didactic formats, and focusing on outcomes that are likely to be perceived as serious may increase the effectiveness of educational meetings. Educational meetings alone are not likely to be effective for changing complex behaviours.

**PLAIN LANGUAGE SUMMARY**

**Continuing education meetings and workshops for health professionals**

Educational meetings are commonly used for continuing medical education with the aim of improving professional practice and, thereby, patient outcomes. Educational meetings include courses, conferences, lectures, workshops, seminars, and symposia.

Eighty-one trials that evaluated the effects of educational meetings were included in this review. Based on these studies, we concluded that educational meetings alone or combined with other interventions can improve professional practice and the achievement of treatment goals by patients. The effect on professional practice tended to be small but varied between studies, and the effect on patient outcomes was generally less. It is not possible to explain the observed differences in effect with confidence but it appeared that higher attendance at the meetings was associated with greater effects, that mixed interactive and didactic education was more effective than either alone, and that the effects were less for more complex behaviours and less serious outcomes.
### Educational meetings with or without other interventions compared to no intervention

**Patient or population:** Health care professionals  
**Settings:** Primary and secondary care  
**Intervention:** Educational meetings with or without other interventions*  
**Comparison:** No intervention

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Adjusted absolute improvement (risk difference)†</th>
<th>Number of studies</th>
<th>Quality of the evidence (GRADE)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compliance with desired practice</td>
<td>Median 6% (1.8 to 15.9)</td>
<td>30</td>
<td>+++O Moderate†</td>
<td>The effect appears to be larger with higher attendance at the educational meetings and with mixed interactive and didactic educational meetings. Educational meetings did not appear to be effective for complex behaviours and they appeared to be less effective for less serious outcomes.</td>
</tr>
<tr>
<td>Patient outcomes</td>
<td>Median 3.0% (0.1% to 4.0%)</td>
<td>5</td>
<td>+++O Moderate†</td>
<td></td>
</tr>
</tbody>
</table>

* The effect of educational meetings alone on professional practice was the same as for multifaceted interventions that included educational meetings.  
† The post intervention risk differences are adjusted for pre-intervention differences between the comparison groups.  
‡ We have downgraded the evidence from high to moderate because of inconsistency in the results that could not be fully explained.

---

**GRADE Working Group grades of evidence**

- **High quality (++++):** Further research is very unlikely to change our confidence in the estimate of effect.
- **Moderate quality (+++O):** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
- **Low quality (++OO):** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
- **Very low quality (++OOO):** We are very uncertain about the estimate.
BACKGROUND

Each year, billions of dollars are spent worldwide on continuing medical education activities (Brown 2002; Vaughn 2006). Continuing professional development is another related but somewhat more comprehensive concept that emphasises a more self-directed approach to education (Peck 2000). In many countries, a demonstration of continuing medical education is mandated by professional or regulatory bodies or it is stimulated by incentives (Peck 2000), which contribute greatly to the increase in these activities.

An underlying assumption is that continuing medical education improves healthcare practice and, thereby, health outcomes for patients. Two overviews of reviews on continuing medical education in general concluded that continuing medical education can be effective (Bloom 2005; Umble 1996) but the effect varied. The first overview was based on 16 reviews conducted between 1984 and 1994 and the other on 26 reviews from the period 1984 to 2004.

There has long been an awareness that the effectiveness of continuing medical education can be measured at three levels: competence, performance, and patient health status (Lloyd 1979) and that the impact declines in that order (Beaudrey 1989). Early studies in this field focused on establishing a causal relationship between continuing medical education and one or all of those outcome levels but, as this relationship was perceived to have been established, the focus shifted (Umble 1996). From the late 1980s, questions of how and why some programs worked better than others were raised and investigators looked for potential explanatory factors. Their focus also shifted from measuring knowledge, attitudes, or skills to measuring physicians’ performance or patients’ health. Commonly reported findings from explanatory analyses were that interventions using an interactive educational format had greater effects than those using a didactic format, and that multifaceted interventions had greater effects than single interventions (Mansouri 2007; Marinopoulos 2007).

Educational meetings are one of the most common continuing medical education activities (Brown 2002; Lloyd 1979). Educational meetings include courses and workshops in various formats. The nature of educational meetings is highly variable in terms of content, the number of participants, the degree and type of interaction, length, frequency, and the targeted practices. Other common continuing medical education activities are audit and feedback (Jamtvedt 2006) and educational outreach (O’Brien 2007), both of which are frequently combined with educational meetings. Quality improvement activities, which are closely related to continuing education (Boonyasai 2007), also commonly use small interactive meetings to facilitate learning and improvements in practice.

Previous versions of this review (Davis 1999; O’Brien 2001) assessed the effects of educational meetings and examined factors that could explain variations in effectiveness. These concluded that interactive workshops can result in moderately large changes in professional practice, while didactic sessions alone are unlikely to change professional practice. Another review of a wide range of guideline implementation strategies (Grimshaw 2004) concluded that educational meetings, with or without educational material, resulted in small to modest improvements when compared to no intervention, which is similar to other strategies.

In this update, we examined the effects of continuing educational meetings on professional practice and patient outcomes. We also investigated factors that might influence the effectiveness of educational meetings. We used methods that have been developed by the Cochrane Effective Practice and Organisation of Care (EPOC) Group (Grimshaw 2003) since the previous review (O’Brien 2001). These methods were used in other recent EPOC reviews (Doumit 2007; Jamtvedt 2006; O’Brien 2007). The provision of printed educational materials has been reported to have little or no effect, in two reviews ( Freemantle 1997; Grimshaw 2001), but this finding has been questioned in a more recent review (Grimshaw 2004). Because printed materials are usually an integral part of educational meetings, we chose to consider printed educational materials as a component of educational meetings and not as an additional independent intervention. Few studies have tested educational meetings without any printed educational materials (Grimshaw 2004).

OBJECTIVES

This review addressed the following questions:

1. Are educational meetings and workshops, alone or in combination with other interventions, effective in improving professional practice or healthcare outcomes?

Comparisons to answer the first question:

- Any intervention in which educational meetings is a component compared to no intervention (Comparison 1).

The primary aim of this analysis was to explore heterogeneity, including potential differences between the effects of educational meetings alone and educational meetings as a component of multifaceted interventions. The main explanatory factors that we considered were the:

- type of intervention (educational meetings alone, with or without educational material, or multifaceted interventions that included educational meetings);
- contribution of educational meetings as a component of the intervention for multifaceted interventions;
2. How does the effectiveness of education meetings compare with that of other interventions?

Comparisons to answer the second question:

- Educational meetings compared to other interventions (Comparison 3).

3. Can educational meetings be made more effective by modifying how they are done?

Comparisons to answer the third question:

- Any intervention in which educational meetings are a component compared to educational meetings alone (Comparison 4).
- Interactive educational meetings compared to didactic (lecture-based) educational meetings (Comparison 5).
- Any other comparison of different types of educational meetings (Comparison 6).

**METHODS**

Criteria for considering studies for this review

Types of studies
Randomised controlled trials (RCTs) were included. Studies using quasi-randomisation or other methods were excluded.

Types of participants
We included studies involving qualified health professionals or health professionals in postgraduate training (for example resident physicians). Studies involving only undergraduate students were excluded.

Types of interventions
We included the following types of educational meetings: conferences, lectures, workshops, seminars, symposia, and courses.

Types of outcome measures
We included studies that reported objectively measured health professional practice behaviours or patient outcomes in a healthcare setting. Studies that measured knowledge or performance in a test situation only were excluded. Studies using patients’ (or simulated patients’) subjective ratings of health professionals’ performance were included.

Search methods for identification of studies

The review was updated using the Cochrane Effective Practice and Organisation of Care Group (EPOC) Trials Register and pending file. We identified all potentially relevant articles in the Register (see EPOC, Specialised Register http://www.mrw.interscience.wiley.com/cochrane/clabout/articles/EPOC/frame.html). We screened studies (1999 to March 2006) that were coded as an RCT or clinical controlled trial (CCT) and with the EPOC-controlled vocabulary term ‘educational meeting’. The EPOC pending file (studies identified using the EPOC search strategy and awaiting assessment) was also searched for the same period, by the EPOC Trials Search Coordinator. We included studies from the previous version of this review and did not undertake any additional searches for studies before 1999. The search history for the previous review is presented in Appendix 1. The reference lists of related systematic reviews and all obtained articles were screened.

An updated search was done in EMBASE (Appendix 1), Scopus, and the EPOC Trials Register (2006 to December 2007). Potentially relevant studies are listed under Studies awaiting classification.

Data collection and analysis

Two review authors (AB and LF) independently screened the titles and abstracts identified from the search process and eliminated any obviously irrelevant studies. The remaining studies were retrieved in full text. Two review authors (LF and AB, AR, GJ, MAOB, FW, or DA) independently applied inclusion criteria. Differences in opinion were resolved by discussions and the involvement of a third author. Studies included in the previous review were reassessed because of changes in the data extraction form and the methods used in this updated review.

Risk of bias

The risk of bias for all included studies was independently assessed by two authors (LF and AB, AR, GJ, MAOB, FW or DD) using seven criteria suggested by EPOC for assessing the risk
of bias of RCTs (see EPOC Data collection checklist http:/
www.epoc.cochrane.org/Files/Website/Reviewer%20Resources/
Data%20Collection%20Checklist%20-%20EPOC%20-%202007- Feb- 27.doc): concealment of allocation, follow up of professionals, follow up of patients or episodes of care, blinded assessment of primary outcome(s), baseline measurement, reliable primary outcome measure(s), and protection against contamination. An overall rating (low, moderate, or high risk of bias) was assigned based on these criteria. As a rule of thumb, studies were assigned a rating of low risk of bias if the first three criteria were scored as done, and there were no important concerns related to the last three criteria; moderate if one or two criteria were scored as not clear or not done; and high if more than two criteria were scored as not clear or not done. For cluster randomised trials protection against contamination was rated as done or not. We also rated concealment of allocation as done if all clusters were randomised at one time and there was no reason to suspect that the allocation process had been influenced by the investigators or participants. We rated completeness of follow up as done if data for at least 80% of the clusters in a cluster randomised trial were collected. However, if many practitioners or their patients had been lost to follow up, we assigned a rating of not done. Any discrepancies in ratings were resolved by discussion and the involvement of a third author.

Data extraction
Two authors independently completed data extraction for all studies. A revised version of the EPOC data collection checklist (see EPOC Data collection checklist http:/
www.epoc.cochrane.org/Files/Website/Reviewer%20Resources/
Data%20Collection%20Checklist%20-%20EPOC%20-%202007- Feb- 27.doc) was used to collect information on study design, type of intervention, presence of controls, type of targeted behaviour, participants, setting, methods (unit of allocation, unit of analysis, methodological quality), outcomes, and results. In addition data, as noted below, were registered. For studies with data that could not be extracted or that lacked baseline information, and that were not older than six to eight years, we contacted the investigators. Discrepancies between authors were resolved through discussion.

Description of explanatory factors

Type of intervention
We categorised interventions as educational meetings alone (with or without educational material) or as multifaceted interventions that included educational meetings. We defined multifaceted interventions as including two or more interventions, such as educational meetings and reminders. In some instances it was difficult to decide whether an intervention was primarily educational outreach, audit and feedback, or continuous quality improvement. In such cases, we used the investigators' objective, research question, or description of the focus of the study to categorise the intervention. We used the following EPOC definitions (http:/
www.epoc.cochrane.org/Files/Website/Reviewer%20Resources/
Data%20Collection%20Checklist%20-%20EPOC%20-%202007- Feb-27 doc) of interventions that might be combined with educational meetings:

- Reminders: any intervention, manual or computerised, that prompts the healthcare provider to perform some action
- Educational outreach: a personal visit by a trained person to health professionals in their own settings
- Audit and feedback: any summary of clinical performance of health care over a specified period of time, given in a written, electronic, or verbal format

Contribution of educational meetings
For multifaceted interventions, two of us independently and subjectively categorised the contribution of educational meetings as a component of the intervention: as a major, moderate, or minor component.

Intensity
We categorised the overall intensity of the educational meetings based on the following characteristics (with the categories listed from 'more intensive' to 'less intensive' for each characteristic):

- number of participants (small, moderate, or large group);
- format (interactive versus didactic);
- source (representatives coming from the local organisation versus a 'professionals' standards review organisation (internal versus external organisation) or the researchers);
- frequency of the educational intervention, categorised as frequent (> 10), moderate (five to 10), infrequent (two to four), and once only;
- total length of education, categorised as prolonged (five days or more), moderate (two to four days), brief (one day), and very brief (less than one day).

Overall intensity was assessed by combining the above characteristics, as:

- intensive (small group AND interactive format AND a supervisor or senior colleague or representative from the local organisation as the source AND (frequent OR prolonged education));
- moderately intensive (any other combination of characteristics than those described in intensive or non-intensive groups, such as small or moderate AND interactive or both interactive + didactic AND local or external organisation AND moderate frequency or moderate length of meeting);
- non-intensive (small or moderate or large group AND (didactic format OR a 'professionals' standards review
If reported by the authors, we recorded the proportion of study participants that attended the educational session(s); if not, we estimated attendance on the basis of information in the text. If this was not possible, attendance was recorded as unknown.

**Setting of care**

We recorded the setting of care as general (family) practice, community-based, hospital (inpatient or outpatient), mixed or other.

**Format**

We categorised educational meetings as interactive, didactic, or mixed. We defined didactic sessions as those that were predominantly lectures or presentations but which may have included question and answer periods. Interactive workshops and seminars were defined as sessions that involved some type of interaction amongst participants in small (< 10 participants), moderate (10 to 19 participants), or large (> 19 participants) groups. The interaction could include role play, case discussion, or the opportunity to practise skills. Mixed sessions included both didactic and interactive components. When in doubt, we categorised educational meetings as mixed.

**Complexity**

The complexity of the targeted behaviour was independently and subjectively categorised by two of us as: high, moderate, or low. The categories depended upon the number of behaviours required, the extent to which complex judgments or skills were necessary, and whether other factors such as organisational change were required for the behaviour to be improved; they also depended on whether there was a need for change by the individual or professional only (one person), communication change, or change in systems. If an intervention was targeted at relatively simple behaviours but there were a number of different behaviours, for example compliance with multiple recommendations for prevention, the complexity was assessed as moderate.

**Seriousness**

The seriousness (importance) of the outcome was independently and subjectively categorised by two of us as: high, moderate, or low. Acute problems with serious consequences were considered as high. Primary prevention was considered moderate. Numbers of unspecified tests or prescriptions were considered as low.

**Baseline compliance**

Baseline compliance with the targeted behaviours was treated as a continuous variable, ranging from zero to 100%, based on the pre-intervention level of compliance given as a mean for both or all groups before the intervention.

**Risk of bias**

We categorised studies as having a low, moderate, or high risk of bias, as described above.

**Data analysis**

We only included in the primary analyses studies with a low or moderate risk of bias and that reported baseline data. For the first main comparison, we carried out a sensitivity analysis by including studies with a high risk of bias and baseline data. All outcomes were expressed as ‘compliance with desired practice’. Professional and patient outcomes were analysed separately. Studies were weighted according to the number of health professionals participating.

To avoid the effect of potentially important baseline differences in compliance between the intervention and control groups of trials, the analyses were based on adjusted estimates of effect, where we adjusted for baseline differences in compliance. For dichotomous outcomes we calculated the adjusted risk difference.

The adjusted risk difference (RD) is the difference in compliance between intervention and control group means after the intervention minus the difference between groups before the intervention. A positive risk difference indicates that compliance improved more in the educational intervention group than in the control group, for example an adjusted risk difference of 0.09 indicates an absolute improvement in care (improvement in compliance) of 9%.

For continuous outcomes we calculated the percentage change relative to the control mean after the intervention: adjusted difference between the post-intervention experimental and control group means divided by the post-intervention control group mean x 100.

Comparisons that allocate clusters but do not account for clustering in the analysis have potential unit of analysis errors resulting in artificially low P values and overly narrow confidence intervals. For such comparisons, we extracted the point estimate and not the P value or confidence interval. For studies with no unit of analysis error and with low or moderate risk of bias and reported baseline data, we recorded adjusted odds ratios (or other measure of effect) and the P values or confidence intervals reported by the authors. We compared these results with our analyses to assess the robustness of our analyses.

When several outcomes were reported in a trial, we only extracted results for the primary outcome. If there was more than one primary outcome, or if the primary outcome was not specified, we calculated effect sizes for each outcome and extracted the median value across the outcomes. In the results tables, we tabulated the median adjusted risk difference (RD) in compliance for the primary outcomes for studies that reported an odd number of primary outcomes. For studies that reported an even number of primary outcomes, we chose the higher of the two middlemost adjusted RDs in compliance for the primary outcomes. In trials that reported summary as well as individual measures of performance, we used the summary measures.

Heterogeneity was explored visually by preparing tables, bubble plots (where the size of the bubble corresponds to the number of healthcare professionals who participated) and box plots (display-
ing medians, interquartile ranges, and ranges) to explore the size of the observed effects in relation to each of these variables. We considered each potential explanatory factor one at a time by looking for patterns in the distribution of the RDs. We hypothesised that greater effects would be associated with:

- multifaceted interventions (versus educational meetings alone),
- more intensive education meetings,
- higher attendance at educational meetings,
- interactive (versus didactic) educational meetings
- less complexity of the targeted behaviour,
- more serious outcomes,
- lower baseline compliance, and
- high risk of bias (versus moderate).

The visual analyses were supplemented with univariate statistical analyses. We used weighted meta-regression to examine how the size of the effect was related to each of the 10 potential explanatory factors listed above, weighted according to the number of healthcare professionals. These analyses were conducted using generalised linear modelling in SAS (Version 9.1. SAS Institute Inc, Cary, NC, USA). We conducted the main analysis for the first comparison using the adjusted RD as the measure of effect.

We planned to supplement these univariate analyses with a multivariate meta-regression. In order to minimise the risk of spurious estimates of effect from the meta-regression, due to a high number of independent variables compared to the number of studies in the analysis, we planned to perform the meta-regression in a stepwise manner with three steps:

1. Each of the potential explanatory factors were analysed as the only independent variable in a meta-regression in order to assess an unadjusted baseline effect, variables with a P value > 0.3 were excluded as explanatory variables in step 3;
2. We examined interactions between the following factors and the type of intervention - the intensity of educational meetings and interactive versus didactic educational meetings, interactions with a P value > 0.3 were excluded from further analysis;
3. Explanatory variables from step 1 (P value ≤ 0.3) and interactions from step 2 were evaluated for potential combination into a final meta-regression model.

**Publication bias**

We used a funnel plot to visually explore the risk of publication bias, using the number of health professionals as a proxy for the precision of the estimate and the adjusted RD as the treatment effect.

## Results

**Description of studies**

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

The search of the EPOC Trials Register and pending files (1999 to March 2006) yielded 768 references. The update of the EMBASE search retrieved references to 2355 studies, while the new search in the EPOC Trials Register and pending files returned 246 references. We identified 77 references which are now awaiting further assessment (see Characteristics of studies awaiting classification).

The table Characteristics of excluded studies lists 33 references, including 14 references that were excluded from the original review.

In this update, 49 new studies have been added to the 32 studies from the previous review, making a total of 81 included studies. Of the 49 new studies, we contacted 20 investigators for further information regarding baseline data or extraction of data. Although 14 of these replied, we were only able to include one of those 14 studies in our analysis.

**Characteristics of the providers and settings**

Thirty-two trials were based in North America (28 in the USA, four in Canada); 34 in Europe (14 in United Kingdom; 10 in the Netherlands; three in Norway; two in France; and one each in Sweden, Denmark, Belgium, Spain, and Scotland); three in Australia; two in Indonesia and South-Africa; and one each in Mali, Thailand, Peru, Mexico, Zambia, Sri Lanka, New Zealand, and Brazil. In most trials the health professionals were physicians. In two studies the providers were nurses (Mazzuca 1987; Simons 2001), in three studies they were pharmacists (Kimberlin 1993; Chalker 2005; de Almeida Neto 2000) or non-physician prescribers (Santoso 1996), and 18 studies involved mixed providers. The setting in 43 of the studies was categorised as general practice, 16 as community-based care, 17 as hospital-based care and five as other types of settings.

**Targeted behaviours**

In 11 trials the behaviours were preventive care, including: identifying and managing problems in marital relationships (Simons 2001; Thompson RS 2000), smoking cessation (Kotke 1989; Strecher 1991; Ward 1996), breastfeeding promotion activities (Westphal 1995), exercise and health behaviours counselling (Kerse 1999; Wilson 1992), screening sigmoidoscopy (Perera 1983), nutrition counselling (Ockene 1996), and follow up of patients with coronary artery disease (Kessling 2002).

Three studies focused on test ordering behaviour change: improved quality of cholesterol testing (Van der Weijden 1999), and decreasing the number of tests requested (Verspagen 2003; Verspagen 2004). Six studies targeted screening behaviours for: cancer (Boisiel 1995; Dietrich 1992; Dolan 1997), cancer and hypertension (Jennett 1988), arthritis (Mazzuca 1987), and presentation of screening tests (Smith 1995).

Thirteen trials targeted prescribing: reducing antibiotic use (Angunawela 1991; Welschen 2004) or both antibiotics and steroids (Chalker 2005), identification of drug misuse (de Almeida 1995).
Neto 2000), improved prescription of non-steroidal anti-inflammatory drugs (Figueiras 2001), reducing the use of injections (Hadiyono 1996), prescription of ACE inhibitors (Kasje 2004), prescribing indicators for upper respiratory tract infection (Meyer 2001), appropriate use of drugs for acute diarrhoea (Santoso 1996) or for asthma (Venenga 1999), adequate informing behaviour regarding prescribed drugs (Kimberlin 1993; Maiman 1988), and prescribing for osteoporosis (Solomon 2004).

Forty-one trials focused on the general management of a wide array of problems. Behaviour was focused on in 41 trials with indicators for management of: low-back pain (Bekkerin 2005; Engers 2005), urinary tract infection and sore throat (Flottorp 2002), sexually transmitted diseases (Garcia 2003), depression (Gask 2004; Gerrity 1999; King 2002; Thompson C 2000; Worrall 1999), schizophrenia (Gray 2004), obstetric practices (Gülmezoglu 2006), preterm delivery (Leviton 1999), tuberculosis (Lewin 2005), obesity (Moore 2003b), asthma (Snee 1999), cardiovascular disease (Ornstein 2004), diabetes (Park 1995; Varroud-Vial 2004; Woodcock 1999), acute myocardial infarction (White 1985), epilepsy (Davis 2004), angina (Heller 2001), infertility (Morrison 2001), neonatal care (Wirtschafter 2003), hypercholesterolaemia (Browner 1994), and congestive heart failure (Feldman 2004). Other studies allocated to this category were several studies targeted at improving communication skills: for physicians (Brown 1999; Clark 1998; Delvaux 2005; Fallowfield 2002; Gilroy 2004; Harmsen 2005; Levinson 1993; Roter 1995) or to a related field, improvement of dietary consultatations (Moore 2003a). Single studies sought to increase the use of active sick leave (Scheel 2002), brief therapy training (Pekarik 1994), and research evidence in public health (Forsetlund 2003); and to improve overall quality management (Bexell 1996), referral practice (Rowlands 2003), and guideline-consistent behaviour (Schechtman 2003). Heale 1988 compared the effect of interactive versus didactic education.

The other studies targeted: handling of frequent attenders at an out-of-hours service (Christensen 2004), improvement of skills in spirometry (Eaton 1999), the rate of back surgery (Goldberg 2001), positioning of stroke patients (Jones 2008), patients’ trust building (Thom 1999) and promotion of the use of advance directives for end of life decisions (Sulmasy 1996).

Characteristics of the intervention

Thirty-two trials tested multifaceted interventions. The most commonly used co-interventions were: reminders (five studies), patient education materials (five studies), supportive services (five studies), feedback reports (10 studies), and educational outreach (five studies). In 12 of these studies educational meetings were rated as: the main component of the intervention, moderate in 13 studies, and as a minor component in seven. We categorised one study as having intensive educational meetings (Gilroy 2004), 25 as moderately intensive, and 54 as non-intensive meetings. In addition, one study compared moderate and low-intensity educational meetings (Browner 1994). Out of the 81 studies, 44 (54%) had an attendance of 80% or more. In 21 studies (26%) it was unclear how many had attended the meetings.

Twenty-three studies examined the effectiveness of interactive educational meetings and 10 studies examined the effectiveness of didactic educational meetings, while 43 studies tested a mixed format. In three studies it was not clear how the intervention should be characterised. In addition, two studies had more than two arms. These compared interactive, didactic, and mixed formats (Heale 1988); or an interactive format compared to didactic (Santoso 1996).

Outcome measures

There was large variation in the number of outcome measures, as well as what was being measured: 62% of the studies used dichotomous outcomes measures, 32% used continuous measures, and four studies used both types of measures. Professional practice, patient outcomes, or both, were studied in: 58 (72%), 9 (11%), and 14 (17%) of the studies respectively. The time to follow up varied from 14 days to two years, with a median follow up of six months.

Risk of bias in included studies

We judged 17 studies to have a low risk of bias, 44 a moderate risk, and 20 a high risk. In 47 trials, we assessed that the allocation of participants to experimental and control groups was adequately concealed. For all but one of the remaining trials, adequacy of concealment could not be determined from the published reports. Outcomes were assessed blindly in 50 of the 81 studies, with for all but three of the remaining studies blinding was assessed as not clear. It was often difficult to assess the loss to follow up, of practices or professionals; for example sometimes the number of health personnel was not reported at all, only the number of clusters. We tried to balance these two considerations; that is if data for all clusters were reported we did not rate follow up as done if more than 20% of participants had been lost to follow up. We assessed 52 (64%) of the trials as having over 80% follow up of participants, 17 trials (21%) as not clear, and 12 (15%) as having less than 80% follow up of the units randomised.

Effects of interventions

See: Summary of findings for the main comparison

Comparison 1: any intervention in which educational meetings were a component compared to no intervention
There were 80 trials involving more than 11,000 health professionals in this comparison. However, 20 of the studies were judged as having a high risk of bias, 13 studies had no baseline data, and sufficient data could not be extracted from three studies. Of the 44 remaining studies, eight studies reported data for continuous outcomes and six studies reported patient outcomes only. Data for each trial in this comparison can be found at [http://www.epoc.cochrane.org/en/newPage2.html](http://www.epoc.cochrane.org/en/newPage2.html).

**Professional practice**

Thirty trials with 36 comparisons reported dichotomous health professional outcomes, had a low or moderate risk of bias, and reported baseline data. The adjusted RDs in compliance with desired practice varied from -2.0% to 36.2%, with a median improvement of 6% (interquartile range 1.8% to 15.9%). When we included the studies judged as having a high risk of bias but that had baseline data (five studies with five comparisons) in a sensitivity analysis it did not change the overall results. The adjusted RDs in compliance with desired practice varied from -2.0% to 36.2% with a median improvement of 6% (interquartile range 2.0% to 14.7%).

The 36 comparisons from the 30 trials were included in univariate meta-regression analyses. There were six factors and two interactions from the univariate analyses that had P < 0.3:

- contribution of educational meetings (P = 0.06);
- attendance at the meetings (P = 0.01);
- interactive versus didactic meetings (P = 0.03);
- complexity of the targeted behaviour (P = 0.02);
- seriousness of the outcome (P = 0.02);
- risk of bias (P = 0.28);
- interaction between interactive versus didactic meetings and multifaceted interventions (P = 0.003);
- interaction between intensity of educational meeting and multifaceted interventions (P = 0.21).

Because of the large number of variables (eight) that were eligible for inclusion in the multivariate analysis relative to the number of included comparisons (36), we chose not to carry out the planned multivariate analysis. This was due to the high risk of spurious findings. For the same reason, the P values from the univariate analyses must be interpreted cautiously. Inspection of the bubble and box plots for the four most statistically significant explanatory factors (P < 0.03) suggested some inconsistent relationships. Higher attendance at the educational meetings was associated with larger adjusted RDs, as hypothesised (P < 0.01). Mixed interactive and didactic education meetings (median adjusted RD 13.6) were more effective than didactic meetings (RD 6.9), as hypothesised; but interactive meetings appeared to be less effective (RD 3.0) (Figure 1).
Figure 1. Box plot of adjusted risk difference (RD) versus the type of education
For complexity of the targeted behaviour, the hypothesis was that the more complex the behaviour the smaller the effect would be. This was the case for high complexity (adjusted RD -0.3) versus moderate (RD 10.5) or low (RD 4.7) complexity behaviours, but not for moderate versus low complexity behaviours. Similarly, we had hypothesised that the more serious the outcome, the greater the effect on the targeted behaviour. This was the case for a high level of seriousness of the targeted behaviour (adjusted RD 7.1) or moderate (RD 9.8) versus low (RD 2.9), but not for high versus a moderate level of seriousness.

The differences in effect estimates between studies with a multifaceted intervention and studies with educational meetings alone were not statistically significant (median adjusted RD 6.0 for both, P = 0.90); and for studies with different baseline compliance rates (P = 0.8).

Eighteen trials reported continuous outcomes, of which eight trials had baseline data and a low or moderate risk of bias. The adjusted relative percentage change varied from 0% to 53%. The median percentage change was 10% (interquartile range 9% to 24%).

**Patient outcomes**

There were 21 trials that reported patient outcomes in this comparison. Out of the 21 trials, 13 trials were of low or moderate risk of bias and had baseline values. For dichotomous outcomes, there were five trials. The adjusted RDs in the achievement of treatment goals varied from -0.9% to 4.6%, with a median improvement of 3.0% (interquartile range 0.1% to 4.0%).

For continuous outcomes there were eight studies (nine comparisons). The adjusted relative percentage change in the patient health indicator varied from -1% to 26%. The median percentage change was 4% (interquartile range 0% to 11%).

See [Summary of findings for the main comparison](www.epoc.uottawa.ca).

**Comparison 2: educational meetings alone compared to no intervention**

There were 56 trials in this comparison, of which 41 trials had dichotomous outcomes and 15 had continuous health professional outcomes. Twenty-four trials (26 comparisons) of educational meetings alone compared to no intervention were judged to have a low or moderate risk of bias and reported baseline data. Data for each trial in this comparison are available at [www.epoc.uottawa.ca](http://www.epoc.uottawa.ca).

**Professional practice**

Of the 24 trials (26 comparisons), 19 trials (21 comparisons) had dichotomous data. The data from these trials were used to calculate the median and interquartile range. The median adjusted RDs varied from -2.0% to 29.3%, with a median of 6% (interquartile range 0% to 12.0%).

Five trials (five comparisons) out of the 24 studies reported continuous outcomes. The data from these trials were used to calculate the median and interquartile range. The median adjusted relative percentage changes ranged from 0% to 50%, with a median of 10% (interquartile range 8% to 32%).

**Patient outcomes**

Seventeen trials reported patient data but only nine studies were judged to be of moderate or low risk of bias and had baseline data. Out of these, there were three trials reporting dichotomous data and six trials comparing continuous data. The adjusted RDs for dichotomous patient outcomes varied from -0.9 to 4.0 with a median improvement of 3.0% (interquartile range -0.9 to 4.0). For continuous patient outcomes the adjusted relative percentage change varied from -1% to 26%, with a median of 8% (interquartile range 0% to 12.0%).

**Comparison 3: educational meetings compared to other interventions**

Two trials compared educational meetings to other interventions. Both trials had a moderate risk of bias and used dichotomous outcomes. The comparison interventions were: a facilitated implementation of an office system to improve services for early detection of cancer (Dietrich 1992), and an educational outreach visit intervention to increase prescribing of recommended non-steroidal anti-inflammatory drugs for patients with osteoarthritis and inflammation (Figueiras 2001). The adjusted RD for the comparison educational meetings versus office meetings was -8.0% decrease in compliance for the educational intervention group. For the comparison of educational meetings versus educational outreach, the adjusted RD was -1.4% decrease in compliance for the educational intervention group.

No patient outcomes were reported in these trials.

**Comparison 4: any intervention in which educational meetings were a component compared to educational meetings alone**

There were seven trials in which a multifaceted intervention that included educational meetings was compared to educational meetings alone. Only one of these trials had a low or moderate risk of bias and reported baseline data (Dietrich 1992). This study compared one-day small group discussions combined with an office system and facilitator with a one-day small group discussions only. The aim was to improve services for early detection of cancer. There was a 12% adjusted relative percentage increase in patients receiving faecal occult blood testing. Another study had a low risk of bias but no baseline data (Browner 1994). A three-hour seminar plus follow-up seminars, patient education materials, and office visits to medical doctors and other staff, plus reminders, were compared to a three-hour seminar. The authors reported that the proportion of patients being screened for hypercholesterolemia was the same for both groups (51%) and the proportion of patients being managed in compliance with guidelines was 34% in the intensive group and 33% in the education only group.

**Comparison 5: interactive educational meetings compared to didactic (lecture based) educational meetings**

Two trials compared interactive to didactic educational meetings. Only one of these had a low or moderate risk of bias and reported baseline data (Santoso 1996). The aim of this study was to improve appropriate drug use in acute diarrhoea, to prevent dehydration.
and death. Locally arranged interactive educational meetings were compared to didactic educational meetings. These were arranged for all prescribers in a health district. Although a somewhat larger improvement was reported for the group receiving interactive education it was not statistically significant (adjusted RD 1.4%).

Comparison 6: any other comparison of different types of educational meetings

One study which was judged as having a high risk of bias; it had no baseline data (Heale 1988).

Publication bias

We visually explored a funnel plot of the 36 studies included in the main comparison (Figure 2). The plot was asymmetrical with more smaller studies spreading far out to the right (a larger adjusted RD) and not to the left (a RD less than zero). While this may be an indication of publication bias, there are other possible explanations for this asymmetry, including poorer methodological quality of the smaller studies, true heterogeneity (for example due to the smaller studies having higher attendance rates), and chance (Cochrane Handbook 2008). It is also plausible that educational meetings may occasionally have (true) large effects and that they rarely have large negative impacts on professional practice. Although, we cannot draw firm conclusions about the existence of publication bias, the asymmetry suggests that studies that include fewer than 100 healthcare professionals may sometimes overestimate the impact of educational meetings on professional practice.
**DISCUSSION**

The main finding of this updated review is consistent with earlier versions of the review. This is that educational meetings can result in small to moderate improvements in professional practice and, as would be expected (Umble 1996), smaller improvements in patient outcomes. However, the results of this update suggest that improvements are most likely to be small (median adjusted RD 6%) even with educational meetings that vary in their impact. The explanation of this, emerging from this update, differs from the previous version of the review. We have included more than twice as many studies as in the previous version and potential explanations for the observed variation in effects are more complex than emerged with the smaller number of studies.

A key finding of the previous review, as well as other reviews (for example Bloom 2005), was that interactive education was more effective than didactic education, which appeared to have little or no effect on professional practice. In this update, we found that mixed interactive and didactic education was most effective, whereas interactive education alone appeared to be least effective; the median adjusted RD for didactic education was 6.9%. A possible explanation for this apparent inconsistency is that the studies often provided minimal descriptions of the interventions, making it difficult to classify them. When in doubt, we categorised interventions as mixed. In future updates, and in practice, it may be more relevant to only use two categories. These are didactic educational meetings and educational meetings that are partially or largely interactive. In the context of this review this would suggest that while didactic education can change professional practice, educational meetings that are partially or largely interactive appear to be more effective.

We did not find a significant difference in the effects of multifaceted interventions and educational meetings alone. The median adjusted RD for both was 6%. This finding of similar effects for multifaceted and single interventions is consistent with the results of two other reviews (Grimshaw 2004; Jamtvedt 2006), although some reviews have concluded that multifaceted interventions are more effective (for example Grimshaw 2001; Mansouri 2007).

Among the other explanatory factors that we explored in this update, those that may help to explain variations in the impact of educational meetings on professional practice are the proportion of professionals in the target audience that attend the meetings, the complexity of the targeted behaviour, and the seriousness or importance of the targeted outcome. The intensity of the educational meetings, the setting, baseline compliance, and the risk of bias did not help to explain the observed variation in effects. These findings should be interpreted cautiously because they are based on indirect (between study) comparisons. There were a large number of potential explanatory factors (10) relative to the number of comparisons included in our primary analysis (36), and several of the explanatory factors were difficult to code. Nonetheless, these findings may provide some useful insights for those planning and evaluating educational meetings.

It is logical that people who do not attend educational meetings would not benefit from them and, therefore, the impact on professional practice would decrease as the proportion of people in the target audience that attend the meetings declines. Only 54% of the included studies had an attendance of 80% or more. Health professionals may select continuing educational activities in areas in which they are interested and, therefore, already performing well. For this reason, the impact on those who did not attend could potentially be greater than on those who did; had they attended (Sibley 1982). Thus, those planning educational meetings may want to consider strategies to increase attendance, particularly amongst those who might not choose to attend based on prior interests, in order to increase the potential impact on targeted practices.

Our findings also support the logic that educational meetings are unlikely to improve practice for highly complex behaviours (median adjusted RD -0.3). Similarly, they suggest that the impact of educational meetings may be smaller for outcomes that health professionals may perceive as not having serious consequences for patients (for example the total number of tests ordered) (median adjusted RD 2.9) compared to outcomes that they may perceive as having moderately or highly serious consequences for patients (median adjusted RD 9.8 and 7.1, respectively).

Although we did not find statistically significant differences in the effects of educational meetings on professional practice, there was a trend suggesting that more intensive interventions might have larger effects, as would be expected. We categorised most of the interventions as non-intensive (27 of the 36 comparisons), none as intensive, and the rest as moderately intensive. Thus we cannot draw conclusions about ‘intensive’ interventions based on these data and it is possible that the approach we used to categorise the intensity of interventions was not adequate to detect relevant differences in intensity amongst the included interventions.

We found only two studies that tested different ways of modifying educational meetings (Dietrich 1992; Santos 1996). Thus there is extremely limited information from direct comparisons of different types of educational meetings to supplement our indirect comparisons and inform decisions about how to modify educational meetings to make them more effective.

We found only two studies that compared the impact on professional practice of educational meetings and other interventions. These studies found that educational meetings alone were less effective than office systems, and slightly less effective than edu-
Other systematic reviews of educational outreach (O'Brien 2007) and audit and feedback (Jamtvedt 2006) have used similar methods to our review. These reviews also omitted studies with high risk of bias from the analysis. The median adjusted RD for professional practice for educational outreach compared to no intervention was 5% (interquartile range 3.0% to 6.2%). The median adjusted RD for audit and feedback compared to no intervention was 4% (interquartile range -0.8% to 9%). These findings are similar to our findings for educational meetings (median adjusted RD 6%, interquartile range 2.9% to 15.3%) suggesting that, with all the limitations of indirect comparisons, the effectiveness of these three interventions may be similar.

**Limitations**

As with any systematic review, our review is limited by the data provided in the included studies. Of the 81 studies that met our inclusion criteria, we judged 20 studies to have a high risk of bias and did not, therefore, include them in our primary analyses. Of the remaining 61 studies, only 30 provided data that could be included in meta-regression analyses exploring the heterogeneity in the effects of educational meetings on professional practice. Thus, despite a large number of relevant studies, these studies provide only a limited basis for informing decisions about when educational meetings are most likely to be effective or how best to design implementation strategies using educational meetings. Also, there may be several relevant studies among the many studies awaiting assessment.

Our findings are further limited by inadequate descriptions of the interventions in many of the included studies, as well as by our ability to characterise studies with respect to the many potential factors that could explain the heterogeneity in the results of the included studies. In particular, we found it difficult to characterise the contribution of educational meetings to multifaceted interventions, the intensity of educational meetings, the format (interactive versus didactic), the complexity of the targeted behaviours, and the seriousness of the targeted outcomes. The need for difficult judgements, and an inadequate basis for making many of these judgements, adds to the need to interpret the results of our analyses cautiously.

The scope of our review is both a strength and a limitation. Not restricting our scope to a specific clinical problem or area increased the number of studies that could be included and reduced the risks of spurious findings. However, it is not possible to draw firm conclusions based on this review regarding the effects of educational meetings for specific clinical problems, or how best to design educational meetings for specific clinical problems. Nonetheless, we would argue that our review provides a useful context in which to interpret the findings of the individual trials included in this review as well as other studies that address more specific questions about the effects of educational meetings.

**Authors’ Conclusions**

**Implications for practice**

Educational meetings, alone or combined with other interventions, can improve professional practice and patient outcomes. The effect is most likely to be small and similar to other types of continuing medical education, such as audit and feedback and educational outreach visits. Strategies to increase attendance at educational meetings, using mixed interactive and didactic formats and focusing on outcomes that are likely to be perceived as serious, may increase the effectiveness of educational meetings. Educational meetings alone are not likely to be effective for changing complex behaviours.

**Implications for research**

Future reports of trials of continuing medical education should include clear and detailed descriptions of the interventions, including the proportion of the target audience that attended, the size of groups at meetings, the length and number of sessions, the teaching techniques, and whether there was any skills practice. They should adhere to the CONSORT recommendations for reporting RCTs (Altman 2001), including the extensions for cluster randomised trials (Campbell 2004) and for pragmatic trials (Zwarenstein 2008).

Further comparisons of educational meetings alone that are targeted at individuals at one level of an organisation and no intervention are unlikely to further our understanding of when educational meetings are likely to be effective and how to improve their effectiveness. Direct comparisons of different types of education are needed, such as different group sizes and different numbers and lengths of follow up. In particular, evaluations of more intensive interventions, which may be both more effective and more costly, compared to less intensive interventions are needed. Evaluations of conceptual models or theories to tailor continuing medical education in order to maximise its effectiveness are also needed. These evaluations should use cluster randomised designs, whenever possible, together with process evaluations to further our understanding of why interventions do or do not work and the variations in their effects (Northstar).

For the next update of this review we will reconsider the categorisation of potential explanatory factors and re-evaluate the use of a multivariate regression analysis.

**Acknowledgements**

Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)
We wish to thank Jessie McGowan and Doug Salzwedel of the EPOC Group for undertaking the searching of the EPOC databases and advising us about further search strategies.

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Zwarenstein 2008

* Indicates the major publication for the study.
## Characteristics of included studies [ordered by study ID]

### Angunawela 1991

| Methods | Cluster RCT  
| Follow up: | providers: DONE  
| patients: N/A  
| Blinded assessment: DONE  
| Baseline: NOT CLEAR  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: HIGH  
| Participants | 43 prescribers in 15 state health institutions and patients (18,766 episodes of care)  
| Country: Sri Lanka  
| Proportion of eligible providers who participated: 94%  
| Outpatient departments; Academic/Teaching setting: MIXED  
| Type of targeted behaviour: PRESCRIBING (antibiotics)  
| Complexity of targeted behaviour: LOW  
| Interventions | 1. CME: didactic seminar 3 hrs + printed material  
| 2. Printed material  
| 3. No intervention control  
| Outcomes | Professional practice: % patients receiving prescriptions for antibiotics  
| Patient: none  
| Seriousness of outcome: LOW  
| Notes |  

### Risk of bias

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<th>Item</th>
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<td>Blinded assessment: DONE</td>
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<td>Reliable outcomes: NOT DONE</td>
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<td>Protection against contamination: DONE</td>
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<td>Overall quality: MODERATE</td>
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### Participants

- 113 physiotherapists in 68 practices
- Country: Netherlands
- Proportion of eligible providers who participated: 21%
- Community-based care
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (low back pain)
- Complexity of targeted behaviour: LOW

### Interventions

1. CME: 2.5 hrs x 2 didactic and interactive workshops targeted at barriers
2. No intervention control (guidelines by mail)

### Outcomes

- Professional practice: proportion of adherence to guidelines for four recommendations
- Patient: % of patients at sick leave during previous 6 weeks at 52 weeks
- Seriousness of outcome: LOW

### Risk of bias

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<td>A - Adequate</td>
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### Methods

<table>
<thead>
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<tr>
<td>Cluster RCT</td>
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<td>Follow up:</td>
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</tr>
<tr>
<td>providers: DONE</td>
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<tr>
<td>Reliable outcomes: NOT CLEAR</td>
<td></td>
</tr>
<tr>
<td>Protection against contamination: DONE</td>
<td></td>
</tr>
<tr>
<td>Overall quality: HIGH</td>
<td></td>
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</tbody>
</table>

### Participants

- Prescribers (clinical officers and medical officers) in 16 health centres
- Country: Zambia
- Proportion of eligible providers who participated: 84%
- Community-based care
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (quality of patient management and rational drug use)
- Complexity of targeted behaviour: LOW

### Interventions

1. CME: 3-day interactive seminar x 2
2. No intervention control

### Outcomes

- Professional practice: overall proportion of patients adequately managed
- Patient: none
- Seriousness of outcome: LOW

### Risk of bias

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors' judgement</th>
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</tr>
</thead>
<tbody>
<tr>
<td>Allocation concealment</td>
<td>Yes</td>
<td>A - Adequate</td>
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</table>
### Boissel 1995

#### Methods
- Cluster RCT
- Follow up: providers: NOT CLEAR
- patients: N/A
- Blinded assessment: NOT CLEAR
- Baseline: NOT DONE
- Reliable outcomes: NOT CLEAR
- Protection against contamination: DONE
- Overall quality: LOW

#### Participants
- 385 general practitioners in 278 practices providing breast and cervical cancer screening
- Country: France
- Proportion of eligible providers who participated: NOT CLEAR
- Primary care
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: SCREENING (for cancer)
- Complexity of targeted behaviour: LOW

#### Interventions
1. CME: one-day seminar and educational material sent four times over one year
2. No intervention control

#### Outcomes
- Professional practice: average number of prescriptions for mammography and smear tests
- Patient: none
- Seriousness of outcome: LOW

#### Notes
- No baseline data

#### Risk of bias

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Brown 1999

<table>
<thead>
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<th>Methods</th>
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<td>Reliable outcomes: NOT DONE</td>
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<td>Overall quality: MODERATE</td>
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</table>

<table>
<thead>
<tr>
<th>Participants</th>
<th>70 primary care physicians, surgeons, medical subspecialists, physician assistants and nurse practitioners</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Country: USA</td>
</tr>
<tr>
<td></td>
<td>Proportion of eligible providers who participated: 7%</td>
</tr>
<tr>
<td></td>
<td>Community-based care</td>
</tr>
<tr>
<td></td>
<td>Academic/Teaching setting: NOT CLEAR</td>
</tr>
<tr>
<td></td>
<td>Type of targeted behaviour: COMMUNICATION BEHAVIOUR</td>
</tr>
<tr>
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<td>Complexity of targeted behaviour: LOW</td>
</tr>
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</table>

<table>
<thead>
<tr>
<th>Interventions</th>
<th>1. CME: didactic and interactive workshop, 8 hrs</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2. No intervention control</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Professional practice: average score on the Art of Medicine Survey</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Patient: none</td>
</tr>
<tr>
<td></td>
<td>Seriousness of outcome: LOW</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Notes</th>
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Risk of bias

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<tr>
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<td>Protection against contamination: DONE</td>
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<tr>
<td>Overall quality: HIGH</td>
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### Participants

<table>
<thead>
<tr>
<th>197 primary care physicians in 174 practices</th>
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</thead>
<tbody>
<tr>
<td>Country: USA</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: 65%</td>
</tr>
<tr>
<td>GPs/family practitioners</td>
</tr>
<tr>
<td>Academic/Teaching setting: NON-TEACHING setting</td>
</tr>
<tr>
<td>Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (screening for hypercholesterolemia)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour: MEDIUM</td>
</tr>
</tbody>
</table>

### Interventions

1. Intensive CME: didactic and interactive seminar: 3 hrs + 2 hrs seminar + a third seminar a couple of months later + phone calls + 2 visits to MD and staff to explain educational material + laminated cards + chart reminders + post-card reminders to patients
2. Standard CME: didactic and interactive seminar: 3 hrs
3. No intervention control

### Outcomes

<table>
<thead>
<tr>
<th>Professional practice: % of patients whose management complied with guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient: none</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
</tr>
</tbody>
</table>

### Notes

No baseline data

### Risk of bias

<table>
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*Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)*

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**Methods**

Cluster RCT
Follow up: providers: DONE
patients: N/A
Blinded assessment: NOT CLEAR
Baseline: DONE
Reliable outcomes: NOT CLEAR
Protection against contamination: DONE
Overall quality: MODERATE

**Participants**

Two districts were randomly selected from 40 districts in Bangkok to represent each of four types of neighbourhoods (industrial, downtown living, modern living, and suburban), then randomly assigned to the control or intervention group. 78 pharmacies were then randomly selected from the 8 districts and randomly assigned to one of the two groups.

78 pharmacies in 8 districts in Bangkok
Country: Thailand
Proportion of eligible providers who participated: 20%

OTHER: pharmacies
Academic/Teaching setting: NON-TEACHING setting
Type of targeted behaviour: PRESCRIBING (dispensing of antibiotics and corticosteroids)
Complexity of targeted behaviour: MEDIUM

**Interventions**

1. CME: educational intervention performed in 3 groups. Pharmacy owners and assistants in a 2-day seminar (case management and rational use of drugs) + enforcement of regulations performed by 6 inspectors + peer review groups
2. No intervention control

**Outcomes**

Professional practice: % clients receiving steroids at request
Patient: none
Seriousness of outcome: MODERATE

**Risk of bias**

<table>
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</tbody>
</table>
### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: N/A  
Blinded assessment: DONE  
Baseline: NOT CLEAR  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

321 general practitioners in 178 practices with 8135 patients  
Country: Denmark  
Proportion of eligible providers who participated: 100%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: Out of hours contacts  
Complexity of targeted behaviour: LOW

### Interventions

1. CME: 5 CME meetings in small groups + economic incentive for a status consultation of patient + feedback/reminder: patients' name and number of contacts once a month  
2. No intervention control

### Outcomes

Professional practice: none  
Patient: fall in number of out-of-hours service contacts  
Seriousness of outcome: LOW

### Notes

No baseline data

### Risk of bias

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</tbody>
</table>
Clark 1998

| Methods | Cluster RCT  
| Follow up:  
| providers: DONE  
| patients: NOT DONE  
| Blinded assessment: DONE  
| Baseline: NOT CLEAR  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: MODERATE  

| Participants | 74 general practice paediatricians and 637 of their asthma patients  
| Country: USA  
| Proportion of eligible providers who participated: 89%  
| Primary care in paediatrics  
| Academic/Teaching setting: NOT CLEAR  
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (asthma care for children)  
| Complexity of targeted behaviour: MEDIUM  

| Interventions | 1. CME: interactive seminar based on theory of self-regulation, 5 hrs  
| 2. No intervention control  

| Outcomes | Professional practice: % parents reporting on some indicators of physician behaviour  
| Patient: indicators of use of care  
| Seriousness of outcome: MODERATE  

| Notes |  

| Risk of bias |  
| Item | Authors’ judgement | Description |  
| Allocation concealment? | Unclear | B - Unclear |  

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**Methods**

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<td>patients: NOT DONE</td>
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<td>Reliable outcomes: DONE</td>
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<td>Protection against contamination: DONE</td>
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</tr>
<tr>
<td>Overall quality: MODERATE</td>
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</tbody>
</table>

**Participants**

- General practitioners from 68 practices in 53 locations with 1133 of their patients
- Country: Scotland
- Proportion of eligible providers who participated: 91%
- General practice
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (epilepsy care)
- Complexity of targeted behaviour: MEDIUM

**Interventions**

1. CME intensive: postal dissemination of guideline + interactive, accredited workshops + dedicated structured protocol documents (tool to be used in patient treatment) + the services of a nurse specialist in epilepsy: who offered advice and training to practices in establishing epilepsy review programs, promoted the use of the guideline in epilepsy management and provided information on epilepsy for both practitioners and patients
2. CME intermediate: postal dissemination of guideline + interactive, accredited workshops + dedicated structured protocol documents (tool to be used in patient treatment)
3. Postal dissemination of a nationally developed guideline

**Outcomes**

- Professional practice: process of care data
- Patient: SF-36 general health related quality of life measures
- Seriousness of outcome: HIGH

**Risk of bias**

| Allocation concealment | A - Adequate | Description |
**Methods**

<table>
<thead>
<tr>
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<tr>
<td>Follow up:</td>
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<tr>
<td>providers: DONE</td>
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<tr>
<td>patients: N/A</td>
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<td>Blinded assessment: NOT CLEAR</td>
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<tr>
<td>Overall quality: MODERATE</td>
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</table>

**Participants**

- 24 pharmacists in 24 pharmacies
- Country: Australia
- Proportion of eligible providers who participated: NOT CLEAR
- Community-based care
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: PRESCRIBING (inappropriate drug use)
- Complexity of targeted behaviour: LOW

**Interventions**

1. CME: interactive and didactic seminar presenting a pharmacy-based protocol based on the Stages of change model with practicing of skills, 3 hrs
2. No intervention control

**Outcomes**

- Professional practice: pharmacist behaviour observed by pseudo-patrons
- Patient: none

**Risk of bias**

<table>
<thead>
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<tbody>
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<td>Unclear</td>
<td>B - Unclear</td>
</tr>
</tbody>
</table>
**Methods**

- Provider RCT
- Follow up: providers: NOT DONE
- patients: N/A
- Blinded assessment: DONE
- Baseline: DONE
- Reliable outcomes: DONE
- Protection against contamination: NOT CLEAR
- Overall quality: MODERATE

**Participants**

- 72 specialists in medical, surgical oncology, radiotherapy, hematology, gynecology etc., caring for cancer patients
- Country: Belgium
- Proportion of eligible providers who participated: 2%
- Hospital setting, outpatients
- Academic/Teaching setting: NON-TEACHING
- Type of targeted behaviour: COMMUNICATION SKILLS (cancer)
- Complexity of targeted behaviour: MEDIUM

**Interventions**

1. CME: 19-hrs basic training + 3hrs x 6 consolidation small group workshops over 3 months
2. 19-hrs basic training

**Outcomes**

- Professional practice: patients’ satisfaction score with interview
- Patient: none
- Seriousness of outcome: HIGH

**Risk of bias**

<table>
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### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: N/A  
Blinded assessment: NOT CLEAR  
Baseline: DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

98 doctors in 98 practices providing cancer screening for 2595 patients  
Country: USA  
Proportion of eligible providers who participated: NOT CLEAR  
Primary care  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: SCREENING (cancer)  
Complexity of targeted behaviour: MEDIUM

### Interventions

1. CME: 1-day small group discussions of relevant topics  
2. 1-day small group discussions of relevant topics + office system with facilitator  
3. Office system with facilitator  
4. No intervention control

### Outcomes

Professional practice: different cancer screening initiatives measured by patient surveys and chart reviews  
Patient: none  
Seriousness of outcome: MODERATE

### Risk of bias

<table>
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<td>Yes</td>
<td>A - Adequate</td>
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</table>
### Dolan1997

| Methods | Cluster RCT  
|         | Follow up: providers: NOT DONE 
|         | patients: N/A 
|         | Blinded assessment: NOT CLEAR  
|         | Baseline: DONE 
|         | Reliable outcomes: NOT CLEAR  
|         | Protection against contamination: NOT CLEAR  
|         | Overall quality: LOW |

| Participants | 82 internal medicine housestaff and attending physicians  
|              | Country: USA  
|              | Proportion of eligible providers who participated: NOT CLEAR  
|              | Hospital, outpatients  
|              | Academic/Teaching setting: UNIVERSITY BASED 
|              | Type of targeted behaviour: SCREENING (cancer)  
|              | Complexity of targeted behaviour: LOW |

| Interventions | 1. CME: educational workshop 1hr x 2  
|               | 2. No intervention control |

| Outcomes | Professional practice: median mean proportion of moderate to high risk patients per physician reporting skin cancer control practices  
|          | Patient: none  
|          | Seriousness of outcome: MODERATE |

| Notes |

| Risk of bias | Item | Authors’ judgement | Description |
|             | Allocation concealment? | Unclear | B - Unclear |

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### Eaton 1999

| Methods                        | Cluster RCT  
|-------------------------------|-------------------  
| Follow up:                    | NOT CLEAR         
| providers:                    | N/A                
| patients:                     | N/A                
| Blinded assessment:           | DONE               
| Baseline:                     | NOT DONE          
| Reliable outcomes:            | DONE               
| Protection against contamination: | DONE          
| Overall quality:              | MODERATE           

| Participants                   | 1 doctor and 1 nurse from each of 30 primary care practices  
|--------------------------------|---------------------------------------------------------------  
| Country:                       | New Zealand                                                  
| Proportion of eligible providers who participated: | 10%              
| General practice               | NOT CLEAR                                                     
| Academic/Teaching setting:     | NOT CLEAR                                                    
| Type of targeted behaviour:    | SPIROMETRY USE (screening, diagnosis and monitoring of respiratory disease)  
| Complexity of targeted behaviour: | LOW             

| Interventions                 | 1. CME: 2 hr workshop: theoretical + practical aspects of performance + handheld spirometer received  
|--------------------------------|--------------------------------------------------------------------------------------------------------  
|                                | 2. Handheld spirometer received                                                                       

| Outcomes                      | Professional practice: spirometry quality assurance data  
|--------------------------------|----------------------------------------------------------  
| Patient:                      | none                                                     
| Seriousness of outcome:       | MODERATE                                                 

| Notes                          | No baseline data                                         

| Risk of bias                   | Authors' judgement  
|--------------------------------|-------------------  
| Allocation concealment?        | Unclear          
| Item                           | B - Unclear      

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<td>Overall quality: LOW</td>
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### Participants

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<tr>
<th>Item</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>Participants</td>
<td>67 general practitioners with 443 low back pain patients</td>
</tr>
<tr>
<td>Country</td>
<td>Netherlands</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
<td></td>
</tr>
<tr>
<td>General practice</td>
<td></td>
</tr>
<tr>
<td>Academic/Teaching setting: NOT CLEAR</td>
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<tr>
<td>Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (low back pain)</td>
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</tr>
<tr>
<td>Complexity of targeted behaviour: LOW</td>
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### Interventions

<table>
<thead>
<tr>
<th>Item</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. CME</td>
<td>2 hr educational session + tools for collaboration with manual, exercise and physical therapists + two scientific articles + national guideline + patient education card</td>
</tr>
<tr>
<td>2. No intervention control</td>
<td></td>
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</tbody>
</table>

### Outcomes

<table>
<thead>
<tr>
<th>Item</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Professional practice: % referral to a therapist; prescription of pain medication</td>
<td></td>
</tr>
<tr>
<td>Patient: % still having low back pain after 6 weeks</td>
<td></td>
</tr>
<tr>
<td>Seriousness of outcome: LOW</td>
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### Notes

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<tr>
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### Risk of bias

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</table>
Fallowfield 2002

Methods
Cluster RCT
Follow up: providers: DONE
patients: DONE
Blinded assessment: DONE
Baseline: NOT CLEAR
Reliable outcomes: NOT CLEAR
Protection against contamination: NOT CLEAR
Overall quality: MODERATE

Participants
160 oncologists from 34 cancer centres
Country: UK
Proportion of eligible providers who participated: 80%
Outpatient clinic
Academic/Teaching setting: NOT CLEAR
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (communicating with cancer patients)
Complexity of targeted behaviour: LOW

Interventions
1. CME: 3-day residential small group course targeting communication skills + 6 patient simulators to provide feedback
2. Residential course + written feedback
3. Written feedback only
4. No intervention control

Outcomes
Professional practice: counts of communication behaviours
Patient: none
Seriousness of outcome: MODERATE

Notes
Data unextractable

Risk of bias

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors’ judgement</th>
<th>Description</th>
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<tbody>
<tr>
<td>Allocation concealment</td>
<td>Yes</td>
<td>A - Adequate</td>
</tr>
</tbody>
</table>
**Methods**

- Cluster RCT
- Follow up:
  - providers: DONE
  - patients: NOT CLEAR
- Blinded assessment: DONE
- Baseline: NOT DONE
- Reliable outcomes: DONE
- Protection against contamination: NOT CLEAR
- Overall quality: MODERATE

**Participants**

- 205 nurses in a large urban home health care agency and their 371 Medicare congestive heart failure patients
- Country: USA
- Proportion of eligible providers who participated: NOT CLEAR
- Community-based care
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (care for patients with heart failure)
- Complexity of targeted behaviour: HIGH

**Interventions**

1. CME: interactive practitioner training with role-play and audiotaping
2. No intervention control

**Outcomes**

- Professional practice: mean number of skilled nursing visits delivered within 90 days
- Patient: none
- Seriousness of outcome: HIGH

**Notes**

- No baseline data

### Risk of bias

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<td>Allocation concealment?</td>
<td>Unclear</td>
<td>B - Unclear</td>
</tr>
</tbody>
</table>
## Methods

- **Cluster RCT**
- **Follow up:** providers: NOT CLEAR
- **patients:** N/A
- **Blinded assessment:** DONE
- **Baseline:** DONE
- **Reliable outcomes:** DONE
- **Protection against contamination:** DONE
- **Overall quality:** MODERATE

## Participants

- 595 general practitioners in 15 geographical areas
- **Country:** Spain
- **Proportion of eligible providers who participated:** NOT CLEAR
- **General practice**
- **Academic/Teaching setting:** NON-TEACHING
- **Type of targeted behaviour:** PRESCRIBING (improvement of prescription of NSAIDS)
- **Complexity of targeted behaviour:** LOW

## Interventions

1. Educational outreach
2. CME: 45-min educational session
3. No intervention control

## Outcomes

- **Professional practice:** prescription of non-steroidal NSAIDS as first choice
- **Patient:** none
- **Seriousness of outcome:** LOW

## Notes

**Risk of bias**

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<td>A - Adequate</td>
</tr>
</tbody>
</table>
### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: N/A  
Blinded assessment: DONE  
Baseline: DONE  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: HIGH  

| Participants | Approximately 650 general practitioners in 142 practices  
Country: Norway  
Proportion of eligible providers who participated: 49%  
General practice  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (urinary tract infections in women and sore throat)  
Complexity of targeted behaviour: LOW |
| --- | --- |

| Interventions | 1. CME: 1-day interactive course about urinary tract infections + summary of the main recommendations in electronic and poster format + patient educational material in electronic and paper format + computer-based decision support and reminders during consultations + increase in the fee for telephone consultations + printed material to facilitate discussions in the practice + points in the continuing medical education programme of The Norwegian Medical Association  
2. 1-day interactive course about sore throat + summary of the main recommendations in electronic and poster format + patient educational material in electronic and paper format + computer-based decision support and reminders during consultations + increase in the fee for telephone consultations + printed material to facilitate discussions in the practice + points in the continuing medical education programme of The Norwegian Medical Association |
| --- | --- |

| Outcomes | Professional practice: use of laboratory tests  
Patient: None  
Seriousness of outcome: LOW |
| --- | --- |

### Notes

### Risk of bias

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*Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)*

Copyright © 2009 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.
### Methods

<table>
<thead>
<tr>
<th>Provider RCT</th>
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<tbody>
<tr>
<td>Follow up:</td>
</tr>
<tr>
<td>providers: DONE</td>
</tr>
<tr>
<td>patients: N/A</td>
</tr>
<tr>
<td>Blinded assessment: NOT CLEAR</td>
</tr>
<tr>
<td>Baseline: NOT DONE</td>
</tr>
<tr>
<td>Reliable outcomes: NOT CLEAR</td>
</tr>
<tr>
<td>Protection against contamination: DONE</td>
</tr>
<tr>
<td>Overall quality: MODERATE</td>
</tr>
</tbody>
</table>

### Participants

| 148 public health physicians |
| Country: Norway |
| Proportion of eligible providers who participated: 45% |
| General practice |
| Academic/Teaching setting: NON-TEACHING |
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (working evidence-based in a public health practice) |
| Complexity of targeted behaviour: HIGH |

### Interventions

1. CME: 1 - 5 day workshop: (1 day: 10 physicians; 3 days: 21 physicians; 5 days: 18 physicians) + discussion list + help desk and information service + access to 5 databases + 3 newsletters |

### Outcomes

| Professional practice: % of physicians having used research explicitly to some degree (used/not used) |
| Patient: none |
| Seriousness of outcome: MODERATE |

### Risk of bias

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</tbody>
</table>
Methods

Cluster RCT
Follow up: providers: DONE
patients: N/A
Blinded assessment: DONE
Baseline: NOT DONE
Reliable outcomes: NOT CLEAR
Protection against contamination: DONE
Overall quality: MODERATE

Participants

14 districts of low socioeconomic status in Lima (pharmacies and physicians)
Country: Peru
Proportion of eligible providers who participated: 79%
Setting of care: pharmacies
Academic/Teaching setting: NON-TEACHING
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (recognition, management and prevention of STDs)
Complexity of targeted behaviour: HIGH

Interventions

1. CME: 1.5 hrs x 3=4.5-hr luncheon training seminars + physicians in each district invited to attend a 6 hr workshop on management of STD syndromes + referral network + monthly follow-up visits for six months to all certified pharmacies and referral physicians and health centres within their district.
2. Seminar on diarrhoea

Outcomes

Professional practice: recognition of STD symptoms, adequate management of the syndrome, recommendations for use of condoms and recommendations for treatment of partners
Patient: none
Seriousness of outcome: HIGH

Notes

Risk of bias

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<tr>
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<td>Unclear</td>
<td>B - Unclear</td>
</tr>
</tbody>
</table>
### Methods

| Cluster RCT | Follow up: providers: DONE |
| Baseline: NOT DONE |
| Blinded assessment: DONE |
| Reliable outcomes: DONE |
| Protection against contamination: DONE |
| Overall quality: MODERATE |

### Participants

| 38 general practitioners with 189 patients |
| Country: UK |
| Proportion of eligible providers who participated: 7% |
| Setting of care: general practice |
| Academic/Teaching setting: NON-TEACHING |
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (assessment and management of depression) |
| Complexity of targeted behaviour: MEDIUM |

### Interventions

1. CME: 2 hr x 5 approved training course sessions, including role play
2. No intervention control

### Outcomes

| Professional practice: 7 indicators for patient satisfaction with doctor |
| Patient: Hamilton depression score |
| Seriousness of outcome: MODERATE |

### Notes

No post-test data

### Risk of bias

<p>| Item | Authors’ judgement | Description |
| Allocation concealment? | Yes | A - Adequate |</p>
<table>
<thead>
<tr>
<th>Method</th>
<th>Description</th>
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<tbody>
<tr>
<td>follow up</td>
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</tr>
<tr>
<td></td>
<td>patients: N/A</td>
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<tr>
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<td>Blinded assessment: DONE</td>
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<td>Baseline: NOT DONE</td>
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<tr>
<td></td>
<td>Reliable outcomes: NOT CLEAR</td>
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<tr>
<td></td>
<td>Overall quality: MODERATE</td>
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<table>
<thead>
<tr>
<th>Participants</th>
<th>49 primary care physicians (two standardized simulated patients)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Country</td>
<td>UK</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated</td>
<td>NOT CLEAR</td>
</tr>
<tr>
<td>Setting of care</td>
<td>General practice</td>
</tr>
<tr>
<td>Academic/Teaching setting</td>
<td>NON-TEACHING</td>
</tr>
<tr>
<td>Type of targeted behaviour</td>
<td>GENERAL MANAGEMENT OF A PROBLEM (depression)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour</td>
<td>MEDIUM</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Interventions</th>
<th>1. CME: 4hr x 2 education sessions given two weeks apart with guidelines + goal-setting + doing a videotape of a patient interview as homework for discussion at last session</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2. No intervention control</td>
</tr>
<tr>
<td>Outcomes</td>
<td>Professional practice: physicians' behaviour as reported by standardised patients</td>
</tr>
<tr>
<td></td>
<td>Patient: None</td>
</tr>
<tr>
<td></td>
<td>Seriousness of outcome: MODERATE</td>
</tr>
</tbody>
</table>

| Notes | No baseline data |

<table>
<thead>
<tr>
<th>Risk of bias</th>
<th>Item</th>
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<th>Description</th>
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<tr>
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<td>Unclear</td>
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### Methods

<table>
<thead>
<tr>
<th>Cluster RCT</th>
<th>Follow up:</th>
</tr>
</thead>
<tbody>
<tr>
<td>providers: NOT CLEAR</td>
<td></td>
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<tr>
<td>patients: N/A</td>
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<tr>
<td>Blinded assessment: NOT CLEAR</td>
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<tr>
<td>Baseline: NOT DONE</td>
<td></td>
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<tr>
<td>Reliable outcomes: DONE</td>
<td></td>
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<tr>
<td>Protection against contamination: DONE</td>
<td></td>
</tr>
<tr>
<td>Overall quality: LOW</td>
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</table>

### Participants

<table>
<thead>
<tr>
<th>Head nurses in 10 community health centres</th>
</tr>
</thead>
<tbody>
<tr>
<td>Country: Malawi</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
</tr>
<tr>
<td>Setting of care: Community-based care</td>
</tr>
<tr>
<td>Academic/Teaching setting: NON-TEACHING</td>
</tr>
<tr>
<td>Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (drug counseling to parents with sick children)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour: LOW</td>
</tr>
</tbody>
</table>

### Interventions

<table>
<thead>
<tr>
<th>1. CME: 11-day training course with nationally adapted guidelines + one supervisory visit from course instructors</th>
</tr>
</thead>
<tbody>
<tr>
<td>2. No intervention control</td>
</tr>
</tbody>
</table>

### Outcomes

<table>
<thead>
<tr>
<th>Professional practice: 10-point composite scale measuring the quality of drug counselling</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient: none</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
</tr>
</tbody>
</table>

### Risk of bias

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<thead>
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</tbody>
</table>
## Goldberg 2001

| Methods | Cluster RCT  
| Follow up:  
| providers: NOT CLEAR  
| patients: DONE  
| Blinded assessment: DONE  
| Baseline: DONE  
| Reliable outcomes: DONE  
| Protection against contamination: DONE  
| Overall quality: MODERATE |
| Participants | Spine surgeons, primary care physicians, patients who were surgical candidates, and hospital administrators in ten communities with annual rates of back surgery above the 1990 national average  
| Country: USA  
| Proportion of eligible providers who participated: 12%  
| Setting of care: hospital setting  
| Academic/Teaching setting: NOT CLEAR  
| Type of targeted behaviour: Surgery (rate of back surgery)  
| Complexity of targeted behaviour: HIGH |
| Interventions | 1. CME: regional study group meetings for neurosurgeons and orthopedists + CME conferences for primary care providers + mailed generalist academic detailing + videodisc patient decision making + small discussion groups of key administrative personnel  
| 2. No intervention control |
| Outcomes | Professional practice: low-back surgical rate  
| Patient: none  
| Seriousness of outcome: HIGH |
| Notes |  
| Risk of bias |  
| Item | Authors' judgement | Description |
| Allocation concealment? | Unclear | B - Unclear |
### Methods

Cluster RCT  
Follow up:  
- providers: DONE  
- patients: NOT DONE  
Blinded assessment: DONE  
Baseline: NOT CLEAR  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

- 12 clusters of 5 community mental health nurses each, based on geographical location  
- Country: UK  
- Proportion of eligible providers who participated: NOT CLEAR  
- Setting of care: community-based care  
- Academic/Teaching setting: NON-TEACHING  
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (compliance therapy)  
- Complexity of targeted behaviour: HIGH

### Interventions

1. CME: 80 hrs of teaching delivered on a day-release basis over 10 weeks  
2. No intervention control

### Outcomes

- Professional practice: none  
- Patient: compliance with schizophrenia medication  
- Seriousness of outcome: HIGH

### Notes

- Cost of training each community mental health nurse: £ 1474

### Risk of bias

<table>
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</tbody>
</table>
Methods

Cluster RCT
Follow up: providers: DONE
patients: N/A
Blinded assessment: DONE
Baseline: DONE
Reliable outcomes: DONE
Protection against contamination: DONE
Overall quality: HIGH

Participants

Doctors, midwives, interns and students in obstetric practices in 22 hospitals in Mexico City and 18 in Thailand
Country: Mexico and Thailand
Proportion of eligible providers who participated: 65%
Hospital, inpatients
Academic/Teaching setting: NON-TTEACHING
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (OBSTETRIC PRACTICES)
Complexity of targeted behaviour: HIGH

Interventions

1. CME: series of 3 workshops at time 0, after 6 weeks and after 6 months + Meeting with hospital directors and department heads + Provision of the database, computers and printers + coordinator from each hospital
2. No intervention control

Outcomes

Professional practice: % change in practice rates in 6 provider behaviours for obstetric care
Patient: none
Seriousness of outcome: HIGH

Risk of bias

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<tbody>
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</tbody>
</table>
Hadiyono 1996

Methods
Cluster RCT
Follow up:
providers: DONE
patients: N/A
Blinded assessment: DONE
Baseline: DONE
Reliable outcomes: NOT CLEAR
Protection against contamination: NOT CLEAR
Overall quality: HIGH

Participants
Prescribers in 24 health centres.
Country: Indonesia
Proportion of eligible providers who participated: 83%
Community-based care
Academic/Teaching setting: NOT CLEAR
Type of targeted behaviour: PRESCRIBING (reduce the use of injections)
Complexity of targeted behaviour: LOW

Interventions
1. CME: Educational workshop 1.5 - 2 hrs with 6 prescribers and 6 patients/community members to discuss reasons for injection used and to arrive at a consensus
2. No intervention control

Outcomes
Professional practice: reduction of injection use
Patient: none
Seriousness of outcome: LOW

Notes

Risk of bias

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</table>
Harmsen 2005

| Methods | Provider RCT  
| Follow up:  
| providers: DONE  
| patients: NOT CLEAR  
| Blinded assessment: DONE  
| Baseline: NOT CLEAR  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: MODERATE |

| Participants | 38 general practitioners with a practice population of at least 25% of non-Western country of origin, resulting in a total of 986 consultations  
| Country: Netherlands  
| Proportion of eligible providers who participated: 22%  
| General practice  
| Academic/Teaching setting: NON-TEACHING  
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (intercultural communication skills)  
| Complexity of targeted behaviour: LOW |

| Interventions | 1. CME, physician intervention: 2.5-day training on intercultural communication based on Pinto’s ‘three-step method’ + patient intervention: 12-min videotaped instruction in the waiting room that the patient should feel free to communicate directly and express any disagreement  
| 2. No intervention control |

| Outcomes | Professional practice: one indicator for mutual understanding and three indicators for patients’ satisfaction  
| Patient: none  
| Seriousness of outcome: HIGH |

| Notes | Risk of bias  
| Item | Authors’ judgement  
| Allocation concealment? | Unclear  
| | B - Unclear |
### Methods

<table>
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<th>Provider RCT</th>
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<tr>
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<td>Baseline: NOT DONE</td>
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<tr>
<td>Reliable outcomes: NOT CLEAR</td>
</tr>
<tr>
<td>Protection against contamination: NOT CLEAR</td>
</tr>
<tr>
<td>Overall quality: LOW</td>
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</tbody>
</table>

### Participants

| 46 family doctors, providing care for patients with one of 6 common problems: transient ischaemic attacks, hypertension, pre-menstrual syndrome, chlamydial infections, dementia, prescribing |
| Country: Canada |
| Proportion of eligible providers who participated: 52% |
| Family practice |
| Academic/Teaching setting: NOT CLEAR |
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (clinical problems within family medicine) |
| Complexity of targeted behaviour: MEDIUM |

### Interventions

| 1. CME: small group, problem-based sessions |
| 2. CME: large group, case problem discussion |
| 3. CME: traditional didactic lecture |

### Outcomes

| Professional practice: performance score as rated by simulated patient visit |
| Patient: none |
| Seriousness of outcome: LOW |

### Risk of bias

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</tr>
</tbody>
</table>
| Methods | Cluster RCT  
| Follow up: providers: DONE  
| patients: N/A  
| Blinded assessment: DONE  
| Baseline: NOT CLEAR  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: MODERATE  
|  
| Participants | Doctors and allied health staff in 37 hospitals with 3240 patients  
| Country: UK  
| Proportion of eligible providers who participated: NOT CLEAR  
| Hospital Academic/Teaching setting: NOT CLEAR  
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (management of unstable angina)  
| Complexity of targeted behaviour: HIGH  
|  
| Interventions | 1. CME: educational session run by a local opinion leader, including feedback on hospital level  
| 2. No intervention control  
|  
| Outcomes | Professional practice: % compliance with guideline for angina  
| Patient: none  
| Seriousness of outcome: HIGH  
|  
| Notes |  

**Risk of bias**

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</table>
### Jennett 1988

| Methods | Cluster RCT  
|---------|----------------------------------|
|         | Follow up: providers: DONE  
|         | patients: N/A  
|         | Blinded assessment: DONE  
|         | Baseline: DONE  
|         | Reliable outcomes: DONE  
|         | Protection against contamination: DONE  
|         | Overall quality: HIGH  

| Participants | 31 family doctors in 25 practices providing care for 2077 episodes of patients with risk of colorectal or prostatic cancer or with hypertension  
|             | Country: Canada  
|             | Proportion of eligible providers who participated: 12%  
|             | Community-based care  
|             | Academic/Teaching setting: NOT CLEAR  
|             | Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (cancer screening and hypertension management)  
|             | Complexity of targeted behaviour: LOW  

| Interventions | 1. CME: 1.5-hour small group meeting + 2 teleconferences over 6-8 weeks + 4 newsletters in cancer screening  
|               | 2. CME: 1.5-hour small group meeting + 2 teleconferences over 6-8 weeks + 4 newsletters in hypertension management  
|               | 2. No intervention control  

| Outcomes | Professional practice: proportion of recommended behaviours in cancer screening and hypertension management  
|          | Patient: none  
|          | Seriousness of outcome: MODERATE  

### Risk of bias

<table>
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<td>Yes</td>
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</tbody>
</table>
### Jones 1998

| **Methods** | Cluster RCT  
|            | Follow up:  
|            | providers: NOT DONE patients: DONE  
|            | Blinded assessment: NOT DONE  
|            | Baseline: DONE  
|            | Reliable outcomes: NOT CLEAR  
|            | Protection against contamination: NOT CLEAR  
|            | Overall quality: LOW  

| **Participants** | 116 nurses in 6 wards in two hospitals  
|                 | Country: UK  
|                 | Proportion of eligible providers who participated: NOT CLEAR  
|                 | Hospital: stroke units and wards  
|                 | Academic/Teaching setting: UNIVERSITY BASED  
|                 | Type of targeted behaviour: REHABILITATION (of stroke patients)  
|                 | Complexity of targeted behaviour: HIGH  

| **Interventions** | 1. CME: mixed teaching format lessons 2hr x 2  
|                  | 2. No intervention control  

| **Outcomes** | Professional practice: % correct positions in a set of observations  
|             | Patient: none  
|             | Seriousness of outcome: HIGH  

| **Risk of bias** |  
|                 | **Item** | **Authors’ judgement** | **Description**  
|                 | Allocation concealment? | Yes | A - Adequate  

### Kasje 2004

| **Methods** | Cluster RCT  
|            | Follow up:  
|            | providers: NOT DONE patients: DONE  
|            | Blinded assessment: DONE  
|            | Baseline: NOT DONE  
|            | Reliable outcomes: DONE  
|            | Protection against contamination: DONE  
|            | Overall quality: MODERATE  

### Kasje 2004 (Continued)

| Participants | 245 general practitioners in 27 peer review groups  
Country: Netherlands  
Proportion of eligible providers who participated: 93%  
General practice  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: PRESCRIBING (chronic heart failure and diabetes mellitus type 2) Complexity of targeted behaviour: LOW |
| Interventions | 1a. CME: 13 peer review groups in arm for condition chronic heart failure: 1-hr interactive peer group session on management of condition with case based discussions  
2a. No intervention control  
1b. CME: 14 peer review groups in arm for condition diabetes mellitus type 2: 1-hr interactive peer group session on management of condition with case based discussions  
2b. No intervention control |
| Outcomes | Professional practice: % of patients receiving prescription of ACE inhibitors  
Patient: none  
Seriousness of outcome: MODERATE |

### Notes

#### Risk of bias

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<thead>
<tr>
<th>Item</th>
<th>Authors' judgement</th>
<th>Description</th>
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<tbody>
<tr>
<td>Allocation concealment?</td>
<td>Yes</td>
<td>A - Adequate</td>
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</table>

### Kerse 1999

| Methods | Cluster RCT  
Follow up: providers: DONE  
patients: NOT DONE  
Blinded assessment: DONE  
Baseline: NOT CLEAR  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: MODERATE |
| Participants | 42 general practitioners and 267 of their patients aged over 65 years  
Country: Australia  
Proportion of eligible providers who participated: 51% general practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: PREVENTIVE CARE (health promotion for elderly people) |
Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)

Kerse 1999  (Continued)

<table>
<thead>
<tr>
<th>Complexity of targeted behaviour: MEDIUM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Interventions</td>
</tr>
<tr>
<td>1. CME: 3-hr didactic seminar on health issues + audit &amp; feedback + 15 mins outreach + card-based prompt system + resource directory</td>
</tr>
<tr>
<td>2. None</td>
</tr>
<tr>
<td>Outcomes</td>
</tr>
<tr>
<td>Professional practice: patients’ recall of discussions with the general practitioner</td>
</tr>
<tr>
<td>Patient: several health outcomes</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
</tr>
<tr>
<td>Notes</td>
</tr>
<tr>
<td>No baseline data</td>
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Risk of bias

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Kiessling 2002

Methods

<table>
<thead>
<tr>
<th>Cluster RCT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Follow up: providers: DONE</td>
</tr>
<tr>
<td>patients: DONE</td>
</tr>
<tr>
<td>Blinded assessment: DONE</td>
</tr>
<tr>
<td>Baseline: DONE</td>
</tr>
<tr>
<td>Reliable outcomes: DONE</td>
</tr>
<tr>
<td>Protection against contamination: DONE</td>
</tr>
<tr>
<td>Overall quality: HIGH</td>
</tr>
</tbody>
</table>

Participants

| 54 general practitioners and 88 of their patients with coronary artery disease |
| Country: Sweden |
| Proportion of eligible providers who participated: NOT CLEAR |
| General practice |
| Academic/Teaching setting: NON-TEACHING |
| Type of targeted behaviour: PREVENTIVE CARE (coronary artery disease) |
| Complexity of targeted behaviour: LOW |

Interventions

| 1. CME: guidelines distributed and presented in a lecture + 1 hr x 3 (4) of case-based education |
| 2. Guidelines distributed and presented in a lecture |
Kiessling 2002  (Continued)

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Professional practice: none (only self-reported)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient</td>
<td>low density lipoprotein cholesterol (mmol/l)</td>
</tr>
<tr>
<td>Seriousness of outcome</td>
<td>MODERATE</td>
</tr>
</tbody>
</table>

Notes

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<th>Risk of bias</th>
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<td>A - Adequate</td>
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</table>

Kimberlin 1993

<table>
<thead>
<tr>
<th>Methods</th>
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<tbody>
<tr>
<td>Cluster RCT</td>
</tr>
<tr>
<td>Follow up:</td>
</tr>
<tr>
<td>providers: DONE</td>
</tr>
<tr>
<td>patients: NOT CLEAR</td>
</tr>
<tr>
<td>Blinded assessment: NOT CLEAR</td>
</tr>
<tr>
<td>Baseline: NOT DONE</td>
</tr>
<tr>
<td>Reliable outcomes: DONE</td>
</tr>
<tr>
<td>Protection against contamination: DONE</td>
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<tr>
<td>Overall quality: MODERATE</td>
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</table>

<table>
<thead>
<tr>
<th>Participants</th>
</tr>
</thead>
<tbody>
<tr>
<td>102 community-based pharmacists providing prescriptions for 762 elderly patients</td>
</tr>
<tr>
<td>Country: USA</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: 24%</td>
</tr>
<tr>
<td>Community-based care</td>
</tr>
<tr>
<td>Academic/Teaching setting: NON-TEACHING</td>
</tr>
<tr>
<td>Type of targeted behaviour: PRESCRIBING (detection of drug problems)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour: MEDIUM</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Interventions</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. CME: homework with post-test + 1-day seminar (mixed format) + follow-up service (help desk)</td>
</tr>
<tr>
<td>2. No intervention control</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Professional practice: % of patients reporting positively about pharmacists' counselling about prescriptions (7 behaviours)</td>
</tr>
<tr>
<td>Patient: none</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
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</table>

Notes

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No baseline data
Kimberlin 1993  
(Continued)

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<td>B - Unclear</td>
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</table>

King 2002

Methods
- Cluster RCT
- Follow up:
  - providers: NOT DONE
  - patients: NOT CLEAR
- Blinded assessment: NOT CLEAR
- Baseline: DONE
- Reliable outcomes: DONE
- Protection against contamination: DONE
- Overall quality: LOW

Participants
- 84 general practitioner principals and 272 patients attending their practices who scored above the threshold for psychological distress
- Country: UK
- Proportion of eligible providers who participated: 10%
- General practice
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (brief cognitive therapy for depression)
- Complexity of targeted behaviour: MEDIUM

Interventions
- 1. CME: 4 half-day training sessions with introduction of guideline for cognitive behaviour therapy
- 2. No intervention control

Outcomes
- Professional practice: (physicians’ self reported attitudes and knowledge)
- Patient: patients’ score on anxiety scales, Beck inventory and SF-36 dimensions
- Seriousness of outcome: MODERATE

Notes

Risk of bias

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<tr>
<td>Allocation concealment?</td>
<td>Unclear</td>
<td>B - Unclear</td>
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</tbody>
</table>
## Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: NOT CLEAR  
Blinded assessment: NOT CLEAR  
Baseline: NOT DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: LOW

## Participants

66 doctors general/family practices providing smoking cessation interventions for 1653 patients  
Country: USA  
Proportion of eligible providers who participated: 6%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: PREVENTIVE CARE (smoking counselling)  
Complexity of targeted behaviour: MEDIUM

## Interventions

1. CME: 6-hr workshop of mixed format + 100 copies of 'Quit and win' smoking cessation manual for patients  
2. 100 copies of 'Quit and win' smoking cessation manual for patients  
3. No intervention control

## Outcomes

Professional practice: patients’ reports of smoking cessation counselling behaviours  
Patient: number of patients smoking after 1 year  
Seriousness of outcome: MODERATE

## Risk of bias

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<th>Item</th>
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<th>Description</th>
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<tr>
<td>Allocation concealment</td>
<td>Unclear</td>
<td>D - Not used</td>
</tr>
</tbody>
</table>
### Levinson 1993

| Methods | Cluster RCT  
|---------|------------------
|         | Follow up:  
|         | providers: DONE  
|         | patients: N/A  
|         | Blinded assessment: DONE  
|         | Baseline: NOT CLEAR  
|         | Reliable outcomes: NOT DONE  
|         | Protection against contamination: NOT CLEAR  
|         | Overall quality: MODERATE  

| Participants | 31 general internists, family doctors in practices encouraged to improve communications skills for 473 patients  
|--------------|-----------------------------------------------------------------
|              | Country: USA  
|              | Proportion of eligible providers who participated: NOT CLEAR  
|              | General practice  
|              | Academic/Teaching setting: NOT CLEAR  
|              | Type of targeted behaviour: DOCTOR - PATIENT COMMUNICATION  
|              | Complexity of targeted behaviour: MEDIUM  

| Interventions | 1. CME: 4.5-hour didactic presentation + case-based discussion  
|---------------|----------------------------------------------------------------------
|               | 2. No intervention control  

| Outcomes | Professional practice: indicators for communication skills with patients in primary care (mean number of statements)  
|----------|-----------------------------------------------------------------  
|          | Patient: none  
|          | Seriousness of outcome: MODERATE  

| Notes | No baseline data  
|-------|------------------

### Risk of bias

| Item                        | Authors' judgement | Description |
|-----------------------------|--------------------|-------------
| Allocation concealment?     | Unclear            | B - Unclear |
Leviton 1999

### Methods
- Cluster RCT
- Follow up: providers: DONE
- patients: N/A
- Blinded assessment: NOT CLEAR
- Baseline: DONE
- Reliable outcomes: DONE
- Protection against contamination: DONE
- Overall quality: HIGH

### Participants
- Estimated 405 obstetricians in 27 hospitals and their preterm delivery cases
- Country: UK
- Proportion of eligible providers who participated: 90%
- Community-based care
- Academic/Teaching setting: MIXED
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (preterm delivery)
- Complexity of targeted behaviour: MEDIUM

### Interventions
1. CME: opinion leaders (one physician and one nurse) appointed from each hospital by the director to serve as local experts + grand rounds lecture on antenatal corticosteroid therapy given by a nationally respected expert + chart reminder system for prompting for therapy + 1-hour group discussion with scenarios, led by opinion leaders + monitoring care to provide feedback
2. No intervention control

### Outcomes
- Professional practice: % of patients receiving antenatal corticosteroids
- Patient: none
- Seriousness of outcome: HIGH

### Risk of bias

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<tr>
<td>Allocation concealment</td>
<td>Yes</td>
<td>A - Adequate</td>
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</tbody>
</table>
Methods
Cluster RCT
Follow up:
providers: DONE
patients: N/A
Blinded assessment: DONE
Baseline: NOT CLEAR
Reliable outcomes: DONE
Protection against contamination: DONE
Overall quality: HIGH

Participants
Clinic staff in 24 nurse-managed municipal primary health clinics with treatment success rates below 70%
Country: South Africa
Proportion of eligible providers who participated: 62%
Community-based care
Academic/Teaching setting: NON-TEACHING
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (tuberculosis) Complexity of targeted behaviour: MEDIUM

Interventions
1. CME: 7 training modules each constituting one training session of 3 hrs + homework + meetings and telephone discussions, drawing on a number of theoretical models
2. No intervention control

Outcomes
Professional practice: none
Patient: % of patients with successful TB treatment completion
Seriousness of outcome: HIGH

Notes

Risk of bias

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<tr>
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<td>Yes</td>
<td>A - Adequate</td>
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</tbody>
</table>
Maiman 1988

### Methods
- Cluster RCT
- Follow up:
  - providers: DONE
  - patients: DONE
- Blinded assessment: NOT DONE
- Baseline: NOT CLEAR
- Reliable outcomes: NOT CLEAR
- Protection against contamination: NOT DONE
- Overall quality: LOW

### Participants
- 83 paediatricians in practices, encouraged to provide medication compliance strategies to patients with otitis media
- Country: USA
- Proportion of eligible providers who participated: 94%
- General practice
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: PRESCRIBING (compliance enhancing practices)
- Complexity of targeted behaviour: LOW

### Interventions
1. CME: tutorial and accompanying printed materials: 2.5 x 2-hr didactic and group discussion + education material
2. Mailed printed materials
3. No intervention control

### Outcomes
- Professional practice: compliance-enhancing strategies and patients with no missed doses
- Patient: none
- Seriousness of outcome: LOW

### Notes
- No baseline data

### Risk of bias

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<tr>
<td>Allocation concealment</td>
<td>Unclear</td>
<td>B - Unclear</td>
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</tbody>
</table>
**Methods**

Cluster RCT
- Follow up: providers: DONE
- Baseline: DONE
- Reliable outcomes: NOT CLEAR
- Protection against contamination: DONE
- Overall quality: MODERATE

**Participants**

- 29 public health nurses in seven older-adult clinics providing care for patients with arthritis
- Country: USA
- Proportion of eligible providers who participated: NOT CLEAR
- Community-based care
- Academic/Teaching setting: NON-TEACHING
- Type of targeted behaviour: Screening and management (of arthritis in elderly patients)
- Complexity of targeted behaviour: LOW

**Interventions**

1. CME: inservice education program on arthritis screening and management in older adults: distribution of required readings; 3hrs mixed format + help desk + laminated screening and management guide + telephone consultation service + 1 hr individual skills training
2. No intervention control

**Outcomes**

- Professional practice: % of clients screened for joint swelling
- Patient: none
- Seriousness of outcome: MODERATE

**Notes**

- Risk of bias

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**Methods**

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<tr>
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<tr>
<td>patients: N/A</td>
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<td>Blinded assessment: NOT CLEAR</td>
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<tr>
<td>Baseline: NOT CLEAR</td>
</tr>
<tr>
<td>Reliable outcomes: NOT CLEAR</td>
</tr>
<tr>
<td>Overall quality: MODERATE</td>
</tr>
</tbody>
</table>

**Participants**

| Primary health care nurses in 22 primary health care clinics |
| Country: South Africa |
| Proportion of eligible providers who participated: 51% |
| Community-based care |
| Academic/Teaching setting: NOT CLEAR |
| Type of targeted behaviour: PRESCRIBING (prescribing practices) |
| Complexity of targeted behaviour: MEDIUM |

**Interventions**

1. CME: 4 days problem-based educational sessions using material from the WHO’s “Guide to Good Prescribing”
2. No intervention control

**Outcomes**

| Professional practice: indicators for rational prescribing in respiratory tract infection and diarrhoea/vomiting |
| Patient: none |
| Seriousness of outcome: HIGH |

**Risk of bias**

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<td>Description</td>
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</table>

| Yes |
| A - Adequate |
Moore 2003 a

| Methods | Cluster RCT  
|---------|------------------------------------------------|
|         | Follow up: NOT CLEAR  
|         | patients: DONE  
|         | Blinded assessment: DONE  
|         | Baseline: NOT CLEAR  
|         | Reliable outcomes: NOT CLEAR  
|         | Protection against contamination: DONE  
|         | Overall quality: MODERATE  
|         |  
| Participants | 84 providers: general practitioners, health visitors, district nurses, midwives and nurse practitioners in 12 general practices  
|             | Country: UK  
|             | Proportion of eligible providers who participated: 23%  
|             | General practice  
|             | Academic/Teaching setting: NOT CLEAR  
|             | Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (nutrition counselling)  
|             | Complexity of targeted behaviour: LOW  
|         |  
| Interventions | Phase 1: 90-min x 3 education for each practice: small groups, multidisciplinary general practice teams, conducted by local clinical opinion leader + Phase 2: 90-min x 2 held on practices’ premises focusing on practicing skills (6 months period) + diet sheets and patient teaching aids  
|         |  
| Outcomes | Professional practice: % of patients reporting having discussed diet with doctor or 7 indicators for phys’ behaviour as reported by patients  
|          | Patient: none  
|          | Seriousness of outcome: LOW  

| Notes | No baseline data  

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<td>Allocation concealment?</td>
<td>Yes</td>
<td>A - Adequate</td>
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</tbody>
</table>
### Methods

Cluster RCT  
Follow up:  
providers: NOT DONE  
patients: NOT DONE  
Blinded assessment: DONE  
Baseline: DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

245 general practitioners and practice nurses in 44 practices  
Country: UK  
Proportion of eligible providers who participated: 28%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (obesity)  
Complexity of targeted behaviour: MEDIUM

### Interventions

90-min x 3 education for each practice: small groups, multidisciplinary general practice teams, conducted by dietitians + tool for estimation of a patient's daily energy requirement + diet sheets and supporting written resources to facilitate the dietary prescription to patients

### Outcomes

Professional practice: none  
Patient: difference in mean weight of patients 12 months after the intervention  
Seriousness of outcome: LOW

### Notes

#### Risk of bias

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**Morrison 2001**

<table>
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<td>Follow up:</td>
<td>NOT CLEAR</td>
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<tr>
<td>providers:</td>
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<td>patients: N/A</td>
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<td>Blinded assessment:</td>
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<td>Baseline:</td>
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<td>Reliable outcomes:</td>
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<td>Protection against contamination:</td>
<td>DONE</td>
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<tr>
<td>Overall quality:</td>
<td>LOW</td>
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<table>
<thead>
<tr>
<th>Participants</th>
<th>335 general practitioners from 221 practices with 689 referrals</th>
</tr>
</thead>
<tbody>
<tr>
<td>Country:</td>
<td>UK</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated:</td>
<td>50%</td>
</tr>
<tr>
<td>General practice</td>
<td></td>
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<tr>
<td>Academic/Teaching setting:</td>
<td>NOT CLEAR</td>
</tr>
<tr>
<td>Type of targeted behaviour:</td>
<td>GENERAL MANAGEMENT OF A PROBLEM (infertility)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour:</td>
<td>MEDIUM</td>
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</table>

<table>
<thead>
<tr>
<th>Interventions</th>
<th>1. CME: guideline + invitation to a discussion meeting + invitation to have an individual visit + individual meetings with key personnel to inform about the project</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>2. Information that a guideline would be received in 12 months</td>
</tr>
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</table>

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Professional practice: mean number of tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient:</td>
<td>none</td>
</tr>
<tr>
<td>Seriousness of outcome:</td>
<td>MODERATE</td>
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### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: N/A  
Blinded assessment: NOT CLEAR  
Baseline: NOT DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: LOW

### Participants

| 45 internists in a managed care setting, providing nutrition counselling in hyperlipidemia | Country: USA | Proportion of eligible providers who participated: 98% | General practice | Academic/Teaching setting: NON-TEACHING |

| Type of targeted behaviour: PREVENTIVE CARE (nutrition counselling for patients with hyperlipidemia) | Complexity of targeted behaviour: LOW |

### Interventions

1. CME: 2.5-hr mixed session (role play, didactic) + patient dietary form, followed by 0.5-hour individualised tutorial  
2. 2.5-hr mixed session (role play, didactic) + patient dietary form, followed by 0.5-hour individualised tutorial + structured office management system  
3. Usual care

### Outcomes

Professional practice: indicators for nutrition counselling and referral (patients’ responses)  
Patient: none  
Seriousness of outcome: LOW

### Notes

No baseline data

### Risk of bias

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors’ judgement</th>
<th>Description</th>
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<tbody>
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<td>Yes</td>
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### Methods

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Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)  
Copyright © 2009 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.
### Methods
- Cluster RCT
- Follow up: providers: NOT DONE
- patients: NOT DONE
- Blinded assessment: NOT CLEAR
- Baseline: NOT DONE
- Reliable outcomes: NOT CLEAR
- Protection against contamination: DONE
- Overall quality: LOW

### Participants
- 35 nursing staffs from four long-term care facilities with a minimum of 20 residents who had been diagnosed with diabetes
- Country: USA
- Proportion of eligible providers who participated: 10%
- Long-term care facilities
- Academic/Teaching setting: NON-TEACHING
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (improving diabetes care)
- Complexity of targeted behaviour: LOW

### Interventions
1. CME: 20-min didactic sessions x 7, slides followed by questions and answers
2. No intervention control

### Outcomes
- Professional practice: 5 indicators of care for patients with diabetes
- Patient: none
- Seriousness of outcome: MODERATE

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### Risk of bias

<table>
<thead>
<tr>
<th>Item</th>
<th>Authors' judgement</th>
<th>Description</th>
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<tbody>
<tr>
<td>Allocation concealment?</td>
<td>Unclear</td>
<td>B - Unclear</td>
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</tbody>
</table>
### Methods

<table>
<thead>
<tr>
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<tbody>
<tr>
<td>Allocation concealment</td>
<td>Yes</td>
<td>A - Adequate</td>
</tr>
</tbody>
</table>

#### Participants

- 22 psychotherapists in 3 outpatient clinics in 3 midwestern cities
- Country: USA
- Proportion of eligible providers who participated: NOT CLEAR
- Outpatient clinics
- Academic/Teaching setting: NON-TEACHING
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (duration of psychotherapy)
- Complexity of targeted behaviour: MEDIUM

#### Interventions

1. CME mixed: 1-day workshop (didactic presentation, skills training, case review, ‘homework’ given for the follow-up) + follow-up: 1.5 hours held once with supervision and discussion of participants’ report of a session with a patient that had been selected for brief therapy
2. None

#### Outcomes

- Professional practice: none extracted
- Patient: scores on the Brief Symptom Inventory
- Seriousness of outcome: MODERATE
<table>
<thead>
<tr>
<th>Methods</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cluster RCT</td>
</tr>
<tr>
<td>Follow up: providers: DONE</td>
</tr>
<tr>
<td>patients: NOT CLEAR</td>
</tr>
<tr>
<td>Blinded assessment: NOT CLEAR</td>
</tr>
<tr>
<td>Baseline: DONE</td>
</tr>
<tr>
<td>Reliable outcomes: DONE</td>
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<tr>
<td>Protection against contamination: DONE</td>
</tr>
<tr>
<td>Overall quality: MODERATE</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Participants</th>
</tr>
</thead>
<tbody>
<tr>
<td>26 primary care physicians at nine clinics</td>
</tr>
<tr>
<td>Country: USA</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
</tr>
<tr>
<td>General practice</td>
</tr>
<tr>
<td>Academic/Teaching setting: NOT CLEAR</td>
</tr>
<tr>
<td>Type of targeted behaviour: PREVENTIVE CARE (teaching sigmoidoscopy in prevention of cancer)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour: MEDIUM</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Interventions</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. CME: two afternoons with didactic presentation, educational material, discussions and skills training (7 sigmoidoscopies)</td>
</tr>
<tr>
<td>2. No intervention control at first, then same as above</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Professional practice: rate of sigmoidoscopy per 1000 patients before and after sigmoidoscopy skills preceptornship for both groups</td>
</tr>
<tr>
<td>Patient: none</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Notes</th>
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<tbody>
<tr>
<td>Not analysed as a randomised controlled trial - data not extractable</td>
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### Risk of bias

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<tbody>
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<td>Unclear</td>
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</tbody>
</table>
| Methods | Cluster RCT  
Follow up:  
providers: NOT DONE  
patients: DONE  
Blinded assessment: DONE  
Baseline: NOT DONE  
Reliable outcomes: DONE  
Protection against contamination: NOT CLEAR  
Overall quality: LOW |
|---|---|
| Participants | 88 primary care physicians and 648 of their patients  
Country: USA  
Proportion of eligible providers who participated: 16%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (physicians’ communication skills for detection of emotional distress)  
Complexity of targeted behaviour: MEDIUM |
| Interventions | 1. CME: 4-hr session x 2 focusing on emotion-handling skills, one week apart with homework in between: tape recording of one patient to be discussed at the second session  
2. 4-hr session x 2 focusing on problem defining skills, one week apart with homework in between: tape recording of one patient to be discussed at the second session  
3. No intervention control |
| Outcomes | Professional practice: number of emotional and problem-defining responses of physicians in patients visits  
Patient: patients’ distress scores at 6 months  
Seriousness of outcome: MODERATE |

### Risk of bias

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### Methods

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<tr>
<th>Cluster RCT</th>
<th>Follow up: providers: DONE</th>
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<tbody>
<tr>
<td></td>
<td>patients: N/A</td>
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<tr>
<td></td>
<td>Blinded assessment: DONE</td>
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<tr>
<td></td>
<td>Baseline: NOT CLEAR</td>
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<tr>
<td></td>
<td>Reliable outcomes: DONE</td>
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<td></td>
<td>Protection against contamination: DONE</td>
</tr>
<tr>
<td></td>
<td>Overall quality: MODERATE</td>
</tr>
</tbody>
</table>

### Participants

- 26 practices with four or more partners
- Country: UK
- Proportion of eligible providers who participated: 9%
- General practice
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: other (referrals to specialist services)
- Complexity of targeted behaviour: LOW

### Interventions

1. CME: several within-practice educational meetings, total of 5 hrs on average for each practice
2. No intervention control

### Outcomes

- Professional practice: referral rates
- Patient: none
- Seriousness of outcome: LOW

### Risk of bias

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<tbody>
<tr>
<td>Allocation concealment?</td>
<td>Yes</td>
<td>A - Adequate</td>
</tr>
</tbody>
</table>
### Methods

Clustering RCT  
Follow up:  
Providers: NOT CLEAR  
Patients: N/A  
Blinded assessment: NOT CLEAR  
Baseline: DONE  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

6 districts with 15 health centers randomly selected from each district.  
Country: Indonesia  
Proportion of eligible providers who participated: 100%  
Community-based care  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: Prescribing (appropriate use of drugs)  
Complexity of targeted behaviour: LOW

### Interventions

1. CME: 2-hr interactive small group face-to-face intervention at the health centre, 8-12 participants  
2. CME: 2-hr didactic seminar with 60-80 participants  
3. No intervention control

### Outcomes

Professional practice: % increase in use of oral rehydration therapy  
Patient: none  
Seriousness of outcome: HIGH

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<tr>
<td><strong>Methods</strong></td>
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</tr>
<tr>
<td>Cluster RCT</td>
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<tr>
<td>Follow up:</td>
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<tr>
<td>providers: DONE</td>
<td></td>
<td></td>
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<tr>
<td>patients: N/A</td>
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<tr>
<td>Blinded assessment: DONE</td>
<td></td>
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<tr>
<td>Baseline: DONE</td>
<td></td>
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<tr>
<td>Reliable outcomes: DONE</td>
<td></td>
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<tr>
<td>Protection against contamination: DONE</td>
<td></td>
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<tr>
<td>Overall quality: HIGH</td>
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<table>
<thead>
<tr>
<th><strong>Participants</strong></th>
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</thead>
<tbody>
<tr>
<td>14 groups with 120 primary care physician and associate practitioners from 2 group model HMO practices</td>
</tr>
<tr>
<td>Country: USA</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
</tr>
<tr>
<td>General practice</td>
</tr>
<tr>
<td>Academic/Teaching setting: NOT CLEAR</td>
</tr>
<tr>
<td>Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (guideline consistent behaviour for provision of services for low back pain)</td>
</tr>
<tr>
<td>Complexity of targeted behaviour: MEDIUM</td>
</tr>
</tbody>
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<table>
<thead>
<tr>
<th><strong>Interventions</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>1. CME: 1.5 hrs physician education + audit &amp; feedback + follow-up visit</td>
</tr>
<tr>
<td>2. Patient education materials (videotape and pamphlet) + one visit from the study investigators + 2 written reminders</td>
</tr>
<tr>
<td>3. Both interventions</td>
</tr>
<tr>
<td>4. No intervention control</td>
</tr>
</tbody>
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<table>
<thead>
<tr>
<th><strong>Outcomes</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>Professional practice: % utilization of at least 1 of 4 indicators of guideline consistent behaviour (on basis of patient care episodes)</td>
</tr>
<tr>
<td>Patient: none</td>
</tr>
<tr>
<td>Seriousness of outcome: MODERATE</td>
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<table>
<thead>
<tr>
<th><strong>Risk of bias</strong></th>
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<tbody>
<tr>
<td><strong>Item</strong></td>
</tr>
<tr>
<td>Authors’ judgement</td>
</tr>
<tr>
<td><strong>Description</strong></td>
</tr>
<tr>
<td>Allocation concealment?</td>
</tr>
</tbody>
</table>
### Methods

- **Cluster RCT**
- Follow up: providers: DONE
- patients: DONE
- Blinded assessment: DONE
- Baseline: NOT DONE?
- Reliable outcomes: DONE
- Protection against contamination: DONE
- Overall quality: HIGH

### Participants

- General practitioners, back pain patients, their employers and local National Insurance Administration staff in 65 municipalities in 3 counties
- Country: Norway
- Proportion of eligible providers who participated: 100%
- Primary care
- Academic/Teaching setting: NON-TEACHING
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (use of active sick leave for back pain)
- Complexity of targeted behaviour: MEDIUM

### Interventions

1. CME: continuing education workshop for GPs on low back pain and active sick leave + targeted information to patients, the local National Insurance Administration staff and employers + a new check box in the form for reporting sick leave (=reminder) + a standard agreement plan between employer and employee for rehabilitation + desktop summary for GPs of clinical practice guidelines + resource person for each region to support GPs and follow-up patients on sick leave for >16 days
2. No intervention control + a third group having a passive strategy: targeted information, check box in report of sick leave, standard agreement and desktop summary

### Outcomes

- Professional practice: % of patients on active sick leave
- Patient: none
- Seriousness of outcome: LOW

### Notes

- No baseline data

### Risk of bias

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<td>Cluster RCT</td>
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<td>Follow up:</td>
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<td>providers: NOT CLEAR</td>
<td></td>
</tr>
<tr>
<td>patients: NOT DONE</td>
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<td>Blinded assessment: NOT CLEAR</td>
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<td>Baseline: NOT DONE</td>
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<td>Reliable outcomes: NOT CLEAR</td>
<td></td>
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<tr>
<td>Protection against contamination: DONE</td>
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<tr>
<td>Overall quality: MODERATE</td>
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### Participants

<table>
<thead>
<tr>
<th>Item</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td>Health visitors in 18 clinics</td>
<td></td>
</tr>
<tr>
<td>Country: UK</td>
<td></td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
<td></td>
</tr>
<tr>
<td>Community-based care</td>
<td></td>
</tr>
<tr>
<td>Academic/Teaching setting: NON-TEACHING</td>
<td></td>
</tr>
<tr>
<td>Type of targeted behaviour: PREVENTIVE CARE (help couples with relationship problems after birth)</td>
<td></td>
</tr>
<tr>
<td>Complexity of targeted behaviour: MEDIUM</td>
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</tr>
</tbody>
</table>

### Interventions

- 1. CME: 4-day training
- 2. No intervention control

### Outcomes

- Professional practice: % of mothers remembering having discussed relationship with health visitor
- Patient: none
- Seriousness of outcome: LOW

### Risk of bias

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<td>A - Adequate</td>
</tr>
</tbody>
</table>
### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: NOT DONE  
Blinded assessment: NOT CLEAR  
Baseline: NOT DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: MODERATE

### Participants

34 general practitioners with 433 asthma/COPD patients  
Country: Netherlands  
Proportion of eligible providers who participated: 63%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (improve care for asthma patients)  
Complexity of targeted behaviour: MEDIUM

### Interventions

1. CME: 2hr x 4 interactive meetings with peer review included + one educational session for practice assistants  
2. No intervention control

### Outcomes

Professional practice: only self-reported  
Patient: patients' self-reported scores of quality of life  
Seriousness of outcome: HIGH

### Risk of bias

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<tbody>
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</tbody>
</table>
**Smith 1995**

| Methods       | Cluster RCT  
|               | Follow up:  
|               | providers: NOT DONE  
|               | patients: N/A  
|               | Blinded assessment: DONE  
|               | Baseline: NOT DONE  
|               | Reliable outcomes: NOT CLEAR  
|               | Protection against contamination: NOT CLEAR  
|               | Overall quality: LOW  

| Participants | 87 obstetricians and midwives in five hospitals agreed to participate  
|             | Country: UK  
|             | Proportion of eligible providers who participated: 69%.  35 providers completed the study  
|             | Hospital, inpatients  
|             | Academic/Teaching setting: MIXED  
|             | Type of targeted behaviour: SCREENING (communication of test results)  
|             | Complexity of targeted behaviour: LOW  

| Interventions | 1. CME: 1-hour video-based training session  
|               | 2. 1-hour video-based training session + feedback on audio taped consultations  
|               | 3. No intervention control  

| Outcomes | Professional practice: scores for information-giving and communication skills  
| Patient: none  
| Seriousness of outcome: LOW  

| Notes |  

| Risk of bias |  
| Item | Authors' judgement | Description  
| Allocation concealment? | Unclear | B - Unclear  

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Methods

Provider RCT
Follow up:
providers: DONE
patients: DONE
Blinded assessment: NOT CLEAR
Baseline: DONE
Reliable outcomes: NOT CLEAR
Protection against contamination: NOT CLEAR
Overall quality: LOW

Participants

21 rheumatologists at 1 large academic arthritis practice
Country: USA
Proportion of eligible providers who participated: 100%
Hospital, outpatients
Academic/Teaching setting: university-based
Type of targeted behaviour: PRESCRIBING (glucocorticoid-induced osteoporosis)
Complexity of targeted behaviour: LOW

Interventions

1. CME: 1.5 hr educational dinner meeting, mixed format (feedback material discussed) + list of patients with rheumatoid arthritis sent to doctors (feedback) 3 weeks later + reminder guideline
2. No intervention control

Outcomes

Professional practice: % of patients having medication for osteoporosis
Patient: none
Seriousness of outcome: LOW

Notes

Risk of bias

<table>
<thead>
<tr>
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### Strecher 1991

<table>
<thead>
<tr>
<th>Methods</th>
<th>Cluster RCT</th>
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<tr>
<td></td>
<td>Follow up:</td>
</tr>
<tr>
<td></td>
<td>providers: DONE</td>
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<tr>
<td></td>
<td>patients: NOT DONE</td>
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<td></td>
<td>Protection against contamination: NOT DONE</td>
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<tr>
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<td>Overall quality: LOW</td>
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<table>
<thead>
<tr>
<th>Participants</th>
<th>261 residents in 11 primary care training programmes providing smoking counselling</th>
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<tbody>
<tr>
<td></td>
<td>Country: USA</td>
</tr>
<tr>
<td></td>
<td>Proportion of eligible providers who participated: 96%</td>
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<td></td>
<td>Hospital, outpatients</td>
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<tr>
<td></td>
<td>Academic/Teaching setting: UNIVERSITY-BASED</td>
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<tr>
<td></td>
<td>Type of targeted behaviour: PREVENTIVE CARE (smoking counselling)</td>
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<tr>
<td></td>
<td>Complexity of targeted behaviour: LOW</td>
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<table>
<thead>
<tr>
<th>Interventions</th>
<th>1. CME: 1-hour tutorial including 10-min slide presentation, 10 min counselling approach, 20-min group discussion + 1-hour small group or individual follow up</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>2. Same tutorial + prompting program (chart-based reminders)</td>
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<tr>
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<td>3. Prompting program</td>
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<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Professional practice: counselling frequency, mean number of five techniques used per patient, 5 counselling techniques</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Patient: none</td>
</tr>
<tr>
<td></td>
<td>Seriousness of outcome: MODERATE</td>
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</tbody>
</table>
## Methods

Cluster RCT  
Follow up:  
providers: NOT CLEAR  
patients: NOT CLEAR  
Blinded assessment: NOT CLEAR  
Baseline: DONE  
Reliable outcomes: NOT DONE  
Protection against contamination: DONE  
Overall quality: LOW

## Participants

88 medical residents and 250 of their patients  
Country: USA  
Proportion of eligible providers who participated: 100%  
General practice  
Academic/Teaching setting: UNIVERSITY-BASED  
Type of targeted behaviour: the making of advance directives  
Complexity of targeted behaviour: LOW

## Interventions

1. CME: 0.5-hr pre clinic lecture on the topic of advance directives + videotape included + a videotaped session of their own patient interview with feedback afterwards=skills training  
2. Control: NONE

## Outcomes

Professional practice: % of charts documenting a discussion regarding advance directives  
Patient: none  
Seriousness of outcome: LOW

## Risk of bias

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</tbody>
</table>
## Thom 1999

### Methods
- Cluster RCT
- Follow up: providers: DONE
- Patients: NOT CLEAR
- Blinded assessment: DONE
- Baseline: DONE
- Reliable outcomes: NOT CLEAR
- Protection against contamination: DONE
- Overall quality: MODERATE

### Participants
- 20 family practice physicians
- Country: USA
- Proportion of eligible providers who participated: 31%
- Family practitioners
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: OTHER (building patients’ trust in their physician)
- Complexity of targeted behaviour: HIGH

### Interventions
1. CME: problem-based, small group discussions with brief didactic presentations, viewing of videotaped patient encounters and role-playing, 7 hrs
2. Control: NONE

### Outcomes
- Professional practice: no primary specified - no baseline so cannot choose the highest?
- Patient: mean trust score
- Seriousness of outcome: HIGH

### Risk of bias

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<td>Allocation concealment?</td>
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<td>A - Adequate</td>
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</table>
| Methods | Cluster RCT  
| Follow up: providers: DONE  
| patients: DONE  
| Blinded assessment: DONE  
| Baseline: DONE (for patient data)  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: HIGH |

| Participants | 169 physicians in 59 primary care practices  
| Country: UK  
| Proportion of eligible providers who participated: 26%  
| General practice  
| Academic/Teaching setting: NON-TEACHING  
| Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (detection and outcome of depression)  
| Complexity of targeted behaviour: LOW |

| Interventions | 1. CME: 4-hr mixed format seminar: videos, role play, small-group discussion cases + guideline + educators remained available to the practices for about 9 months  
| 2. Control: NONE |

| Outcomes | Professional practice: % detection of depressive patients  
| Patient: % of patients with HAD depression score >= 8  
| Seriousness of outcome: MODERATE |

| Notes |  

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<td>Overall quality: MODERATE</td>
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| Participants     | 208 providers (mixed) in 5 primary care clinics of a large health maintenance organization |
|                  | Country: USA |
|                  | Proportion of eligible providers who participated: NOT CLEAR |
|                  | General practice |
|                  | Academic/Teaching setting: NOT CLEAR |
|                  | Type of targeted behaviour: PREVENTIVE CARE (identification and management of domestic violence) |
|                  | Complexity of targeted behaviour: LOW |

| Interventions    | 1. CME: 2 half-day training sessions + extra training for opinion leaders + bimonthly newsletter + 4 clinic educational rounds + system support: posters, cue cards, questionnaires + feedback of results |
|                  | 2. Control: NONE |

| Outcomes         | Professional practice: % of patients being asked about domestic violence |
|                  | Patient: none |
|                  | Seriousness of outcome: HIGH |

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</table>
Van der Weijden 1999

### Methods

Cluster RCT  
Follow up:  
providers: DONE  
patients: NOT CLEAR  
Blinded assessment: DONE  
Baseline: DONE  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: HIGH

### Participants

32 general practitioners in 20 practices, 3950 patient records  
Country: Netherlands  
Proportion of eligible providers who participated: NOT CLEAR  
General practitioners  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: TEST ORDERING (performance in daily practice regarding targeted cholesterol testing)  
Complexity of targeted behaviour: LOW

### Interventions

1. CME: interactive teaching by local opinion leaders in moderate group size: 3 hrs + consultation registration form (reminders) + desktop flow chart of guideline + patient education leaflet + 2 outreach visits/face-to-face instruction at workplace  
2. Control: postal distribution of guideline with scientific background materials

### Outcomes

Professional practice: median proportion of patients for whom the GP performed repeat testing to diagnose hypercholesterolaemia  
Patient: none  
Seriousness of outcome: LOW

### Risk of bias

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### Varroud-Vial 2004

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<table>
<thead>
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<th>Participants</th>
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<tr>
<td>67 general practitioners in four separate districts of one region</td>
</tr>
<tr>
<td>Country: France</td>
</tr>
<tr>
<td>Proportion of eligible providers who participated: NOT CLEAR</td>
</tr>
<tr>
<td>Family practitioners</td>
</tr>
<tr>
<td>Academic/Teaching setting: NON-TEACHING</td>
</tr>
<tr>
<td>Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (glycaemic control of patients with Type 2 diabetes in a primary care setting)</td>
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<td>Complexity of targeted behaviour: LOW</td>
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<table>
<thead>
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<tr>
<td>1. CME, staged diabetes management program: 3.5 hrs x 3 educational sessions</td>
</tr>
<tr>
<td>2. Control: no intervention control</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Professional practice: indicators for treatment of patients</td>
</tr>
<tr>
<td>Patient: HbA1c</td>
</tr>
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<td>Seriousness of outcome: MODERATE</td>
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### Risk of bias

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<td>B - Unclear</td>
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</table>
**Methods**

Cluster RCT  
Follow up:  
providers: NOT CLEAR  
patients: N/A  
Blinded assessment: DONE  
Baseline: DONE  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: MODERATE

**Participants**

665 physicians from general practice  
Country: Netherlands, Sweden, Slovakia and Norway  
Proportion of eligible providers who participated: NL 24%; S 35%; SK 20%; N 31%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: PRESCRIBING (asthma care)  
Complexity of targeted behaviour: LOW

**Interventions**

1. CME: 2 educational meetings (self-learning method in small peer groups) on asthma care + individual feedback presented in group for discussion  
2. 2 educational meetings (self-learning method in small peer groups) on the care of urinary tract infection + individual feedback presented in group for discussion

**Outcomes**

Professional practice: % correct prescribing for asthma  
Patient: none  
Seriousness of outcome: LOW

**Risk of bias**

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</table>

Continuing education meetings and workshops: effects on professional practice and health care outcomes (Review)
**Methods**

Cluster RCT  
Follow up: providers: DONE  
patients: N/A  
Blinded assessment: DONE  
Baseline: DONE  
Reliable outcomes: DONE  
Protection against contamination: DONE  
Overall quality: HIGH

**Participants**

174 primary care physicians in 26 local primary care practice groups in 5 health care regions  
Country: Netherlands  
Proportion of eligible providers who participated: 70%  
Primary care  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: TEST ORDERING (reducing inappropriate test ordering)  
Complexity of targeted behaviour: LOW

**Interventions**

1. CME: multifaceted with interactive educational small group meeting + audit and feedback (moderate) for 3 selected clinical problems  
2. Control: multifaceted with interactive educational small group meeting + audit and feedback (moderate) for 3 other selected clinical problems

**Outcomes**  
Professional practice: mean number of inappropriate tests, per physician per 6 months  
Patient: none  
Seriousness of outcome: LOW

**Risk of bias**

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The table below contains information from the document "Verstappen 2004".

<table>
<thead>
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<td>Reliable outcomes: DONE</td>
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<td>Overall quality: HIGH</td>
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</tbody>
</table>

| Participants           | 194 primary care physicians in 27 local primary care practice groups in 5 health care regions |
|                       | Country: Netherlands         |
|                       | Proportion of eligible providers who participated: 71% |
|                       | Primary care                |
|                       | Academic/Teaching setting: NON-TEACHING |
|                       | Type of targeted behaviour: TEST ORDERING (reducing inappropriate test ordering) |
|                       | Complexity of targeted behaviour: LOW |

| Interventions         | 1. CME: multifaceted with interactive educational small group meeting + audit and feedback (moderate) |
|                       | 2. Control: audit and feedback |

| Outcomes              | Professional practice: mean number of tests per physician |
|                       | Patient: none |
|                       | Seriousness of outcome: LOW |

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</tbody>
</table>
Methods

Cluster RCT
Follow up:
providers: DONE
patients: DONE
Blinded assessment: DONE
Baseline: DONE
Reliable outcomes: DONE
Protection against contamination: DONE
Overall quality: MODERATE

Participants

34 general practice trainees providing preventive care for 1500 patients
Country: Australia
Proportion of eligible providers who participated: 50%
General practice
Academic/Teaching setting: UNIVERSITY-BASED
Type of targeted behaviour: PREVENTIVE CARE (stop-smoking counselling)
Complexity of targeted behaviour: MODERATE

Interventions

1. CME, mixed: 3-day workshop (didactic presentation, small group skill practice, role-play)
2. Control: 3-day workshop in rational prescribing

Outcomes

Professional practice: number of patients asked about smoking status
Patient: none
Seriousness of outcome: MODERATE

Notes

Risk of bias

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</table>
Methods

Cluster RCT
Follow up: providers: DONE
patients: N/A
Blinded assessment: DONE
Baseline: DONE
Reliable outcomes: DONE
Protection against contamination: DONE
Overall quality: HIGH

Participants

12 peer review groups including 100 general practitioners with their collaborating pharmacists
Country: Netherlands
Proportion of eligible providers who participated: 29%
General practice
Academic/Teaching setting: NOT CLEAR
Type of targeted behaviour: PRESCRIBING (of antibiotics for respiratory tract symptoms)
Complexity of targeted behaviour: LOW

Interventions

1. CME: peer-group education meeting with communication skills training + feedback presented at practice level + 2-hr group education for assistants + education material for patients at practice site
2. Control: No intervention control

Outcomes

Professional practice: rate of antibiotic prescribing in %
Patient: patient satisfaction
Seriousness of outcome: LOW

Notes

Risk of bias

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</tbody>
</table>
### Westphal 1995

| Methods | Cluster RCT  
| Follow up: providers: DONE  
| patients: N/A  
| Blinded assessment: NOT CLEAR  
| Baseline: DONE  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: MODERATE |

| Participants | Health professionals in 8 maternity hospitals providing advice about breast feeding  
| Country: Brazil  
| Proportion of eligible providers who participated: NOT CLEAR  
| Hospital setting: Academic/Teaching setting: University based/teaching setting  
| Type of targeted behaviour: PREVENTIVE CARE (breast-feeding practice)  
| Complexity of targeted behaviour: HIGH |

| Interventions | 1. CME: mixed format full-time for 14 days over 3 weeks  
| 2. Control: no intervention control |

| Outcomes | Professional practice: score (1-10) for institutional change according to WHO’s 10 steps for successful breast-feeding  
| Patient: none  
| Seriousness of outcome: LOW |

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<td>MODERATE</td>
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### Participants

- 103 family doctors or general internists in 12 communities caring for in-patients post-myocardial infarction
- Country: USA
- Proportion of eligible providers who participated: 71%
- Family practitioners
- Academic/Teaching setting: NOT CLEAR
- Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (care for acute myocardial infarction)
- Complexity of targeted behaviour: MEDIUM

### Interventions

1. CME: 3.5-hr educational session: 2 hrs with traditional methods and 1.5 h of discussions and case examples
2. Control: no intervention control

### Outcomes

- Professional practice: overall measures of desired patient care
- Patient: none
- Seriousness of outcome: HIGH

### Risk of bias

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</table>
### Wilson 1992

| Methods | Cluster RCT  
| Follow up:  
| providers: DONE  
| patients: DONE  
| Blinded assessment: DONE  
| Baseline: DONE  
| Reliable outcomes: NOT CLEAR  
| Protection against contamination: DONE  
| Overall quality: MODERATE |

| Participants | 22 family doctors providing exercise counselling for 420 patients  
| Country: CANADA  
| Proportion of eligible providers who participated: 12%  
| General practice  
| Academic/Teaching setting: NON-TEACHING  
| Type of targeted behaviour: PREVENTIVE CARE (exercise counselling)  
| Complexity of targeted behaviour: LOW |

| Interventions | 1. CME: 2-hour workshop with three components: discussion forum, practical teaching and overview of resources + mailed material  
| 2. Control: no intervention control |

| Outcomes | Professional practice: % of patients with whom physicians discussed exercise, as reported by patients  
| Patient: none  
| Seriousness of outcome: MODERATE |

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<td>patients: NOT CLEAR</td>
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<td>Reliable outcomes: NOT DONE</td>
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<td>Protection against contamination: DONE</td>
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<tr>
<td>Overall quality:</td>
<td>LOW</td>
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</tbody>
</table>

### Participants

- Estimated 1097 doctors (family practitioners + paediatricians) and nurses in 40 community hospitals caring for newborns in neonatal intensive care units
- Country: USA
- Proportion of eligible providers who participated: NOT CLEAR

### Hospital

- Academic/Teaching setting: NON-TEACHING

### Type of targeted behaviour

- GENERAL MANAGEMENT OF A PROBLEM (respiratory distress in neonatals)
- Complexity of targeted behaviour: HIGH

### Interventions

1. CME: 1.5 hrs x 2 given 6-8 months apart: mediated lecture/case study presentation + monthly newsletter
2. 1.5 hrs x 2 given 6-8 months apart: mediated lecture/case study presentation + protocol for treatment of respiratory distress + newsletter
3. Control: newsletter

### Outcomes

- Professional practice: combined process score
- Patient: neonatal mortality
- Seriousness of outcome: HIGH

### Risk of bias

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### Methods

Cluster RCT  
Follow up:  
providers: NOT CLEAR  
patients: NOT DONE  
Blinded assessment: DONE  
Baseline: NOT DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: LOW

### Participants

107 general practitioners and nurses in 41 practices  
Country: UK  
Proportion of eligible providers who participated: NOT CLEAR  
General practice  
Academic/Teaching setting: NON-TEACHING  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (patient centred diabetes care)  
Complexity of targeted behaviour: LOW

### Interventions

1. CME: general practitioners: 0.5-day group training; nurses: 1.5 days  
2. Control: no intervention control

### Outcomes

Professional practice: patients' recognition of a patient booklet, recognition of an insert to the booklet, patients' reports of nurse consulting behaviour  
Patient: none  
Seriousness of outcome: MODERATE

### Notes

No baseline data

### Risk of bias

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</table>
Worrall 1999

| Methods | Cluster RCT  
Follow up:  
providers: DONE  
patients: NOT CLEAR  
Blinded assessment: NOT CLEAR  
Baseline: DONE  
Reliable outcomes: NOT CLEAR  
Protection against contamination: DONE  
Overall quality: MODERATE |
|----------|--------------------------------------------------|
| Participants | 42 family physicians  
Country: Canada  
Proportion of eligible providers who participated: 41%  
General practice  
Academic/Teaching setting: NOT CLEAR  
Type of targeted behaviour: GENERAL MANAGEMENT OF A PROBLEM (management of depression)  
Complexity of targeted behaviour: MEDIUM |
| Interventions | 1. CME: 3-hr small group educational session: teaching and case-based discussion + psychiatrist help service available once a week + clinical guidelines  
2. Control: mailed clinical guidelines on the management of depression |
| Outcomes | Professional practice: % of correct diagnoses of depression  
Patient: mean patient score on the Centre for Clinical Epidemiological Studies Depression scale  
Seriousness of outcome: MODERATE |

Notes

**Risk of bias**

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

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<tr>
<td>Campbell 1991</td>
<td>Outcomes were not measured in a clinical situation</td>
</tr>
<tr>
<td>Carlsson 1998</td>
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</tr>
<tr>
<td>Casebeer 1999</td>
<td>Audioconferences</td>
</tr>
<tr>
<td>Davidoff 1989</td>
<td>Control group received education</td>
</tr>
<tr>
<td>Doyne 2004</td>
<td>Educational outreach</td>
</tr>
<tr>
<td>Dunn 1992</td>
<td>Outcomes were not measured in a clinical situation</td>
</tr>
<tr>
<td>Eckstrom 1999</td>
<td>Quasirandomised</td>
</tr>
<tr>
<td>Francke 1997</td>
<td>Outcomes were based on self-report</td>
</tr>
<tr>
<td>Gifford 1999</td>
<td>Outcomes measured in a test situation</td>
</tr>
<tr>
<td>Huang 2002</td>
<td>Unclear whether the outcomes were reported by participants or whether they were observed by others</td>
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<td>Langewitz 1998</td>
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<td>Lundgren 1999</td>
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<td>Martin 2001 1</td>
<td>Testing of an algorithm for improving nutritional support, not of an educational intervention</td>
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<tr>
<td>Morrison 2001 1</td>
<td>Testing the effectiveness of guidelines, not an educational meeting</td>
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<tr>
<td>O’Neill 1999</td>
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<td>Ockene 1995</td>
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<tr>
<td>Pinkerton 1980</td>
<td>The intervention was video-watching</td>
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<td>Premaratne 1999</td>
<td>Testing of the effectiveness of an asthma centre, not of an educational meeting</td>
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<td>Educational outreach</td>
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<tr>
<td>Quirk 1991</td>
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<tr>
<td>Ratanajamit 2002</td>
<td>The participants were drugstore personnel</td>
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</table>
We defined the intervention as educational outreach

Outcomes were not measured in a clinical situation

Outcomes were measured by patients that the physicians knew were fake (test situation)

Outcomes were based on self-report

The intervention was printed material, not educational meeting

Outcomes were not measured in a clinical situation

The participants were physicians under education

Outcomes were not measured in a clinical situation

The outcome measures (adherence scores) were a mixture of subjective and objective measures.

Not described as educational meeting

Outcomes were based on self-report
DATA AND ANALYSES
This review has no analyses.

APPENDICES
Appendix 1. Search strategies

Search strategy for previous review
The previous review (O'Brien 2001) searched the Cochrane Effective Practice and Organisation of Care (EPOC) Group Specialised Register, MEDLINE (1966 to January 1999) without language restrictions, and the Research and Development Resource Base in Continuing Medical Education (RDRB/CME) (Davis 1991). The reference lists of related systematic reviews and all articles obtained were reviewed. The terms for the MEDLINE search follow: education/; exp education, continuing/; exp education, graduate/; internship and residency/; exp inservice training/; preceptorship/; exp teaching/. The educational terms were combined with methodological terms.


Database: EMBASE <1980 to 2007 Week 49>

Search Strategy:
--------------------------------------------------------------------------------
1  *medical education/ (27217)
2  *continuing education/ (4249)
3  *postgraduate education/ (2708)
4  ((education$ or train$) adj (program$ or intervention? or meeting? or session? or strateg$ or workshop?)).tw. (30225)
5  ((education$ or train$) adj (lecture? or symposi$ or course?)).tw. (1947)
6  or1-5 (61610)
7  (random$ or placebo$).tw. (402856)
8  ((single$ or double$ or triple$ or treble$) and (blind$ or mask$)).tw. (94419)
9  controlled clinical trial:.tw. (9545)
10  or1-7 (423112)
11  6 and 10 (5020)
12  Animals/ (18216)
13  Humans/ (5951720)
14  12 not (12 and 13) (14454)
15  11 not 14 (5020)
16  limit 15 to yr="2003 - 2007" (2210)

WHAT’S NEW
Last assessed as up-to-date: 29 June 2008.
New citation required and conclusions have changed

This is an update of previously published review. Forty-nine new studies have been added to the 32 studies from the previous review, making a total of 81 included studies. The search was re-run in December 2007: Seventy-seven references are listed under 'Studies awaiting classification'.

New search has been performed

All searches updated.

HISTORY

Review first published: Issue 2, 2001

19 June 2008 Amended Converted to new review format.

CONTRIBUTIONS OF AUTHORS

<table>
<thead>
<tr>
<th>Task</th>
<th>Contributor</th>
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<tr>
<td>Drafted the protocol</td>
<td>LF, ADO, GJ, AB</td>
</tr>
<tr>
<td>Searched for trials</td>
<td>EPOC</td>
</tr>
<tr>
<td>Scanned titles and abstracts for eligibility</td>
<td>LF, AB</td>
</tr>
<tr>
<td>Obtained copies of potentially eligible trials</td>
<td>LF</td>
</tr>
<tr>
<td>Appraised and select which trials to include</td>
<td>LF/AB; LF/AR; LF/GJ; LF/MAO; LF/FW; LF/DAD</td>
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<td>Extracted data from trials</td>
<td>LF/AB; LF/AR; LF/GJ; LF/MAO; LF/FW; LF/DAD</td>
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<tr>
<td>Entered data</td>
<td>LF</td>
</tr>
<tr>
<td>Carried out the analysis</td>
<td>JOJ</td>
</tr>
<tr>
<td>Interpreted the analysis</td>
<td>All authors</td>
</tr>
<tr>
<td>Drafted the final review</td>
<td>LF, ADO</td>
</tr>
</tbody>
</table>
Prepared tables and figures

JOJ, ADO, LF

All authors reviewed the draft of the protocol and the systematic review

DECLARATIONS OF INTEREST

None

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

None

INDEX TERMS

Medical Subject Headings (MeSH)

*Congresses as Topic; Education, Continuing [methods; *standards]; *Process Assessment (Health Care); Professional Practice [*standards]; Randomized Controlled Trials as Topic

MeSH check words

Humans