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## Self-management education for chronic obstructive pulmonary disease (Review)

Monninkhof E, van der Valk PP, van der Palen JJ, van Herwaarden CLA, Partidge MR, Walters EH, Zielhuis GG, van Herwaarden KC, Partridge MM

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[Intervention Review]

# Self-management education for chronic obstructive pulmonary disease

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## ABSTRACT

### Background

In asthma, self-management programmes have been proven to be effective. In COPD, their value is not clear.

### Objectives

To assess the efficacy of COPD self-management/ education programmes on health outcomes and use of health services

### Search methods

We searched the Cochrane Airways Group trial register, MEDLINE (January 1985 to October 2001), reference lists, and abstracts of medical conferences. We also contacted research groups in the field for ongoing trials and unpublished material.

### Selection criteria

Controlled trials (randomised and non-randomised) of self-management education in patients with COPD. Studies focusing mainly on physical pulmonary rehabilitation were excluded.

### Data collection and analysis

Two reviewers independently assessed trial quality and extracted data. Investigators were contacted for additional information.

### Main results

The reviewers included 12 articles describing eight randomised-controlled trials and one controlled clinical trial. Self-management education was compared with usual care in eight studies. The studies in this review assessed a broad-spectrum of outcome measures with different follow-up times. Synthesis of the results using meta-analysis was not always possible. The studies showed no effect of self-management education on hospital admissions, emergency room visits, days lost from work and lung function. Inconclusive results were observed on health-related quality of life (HRQoL): studies using the disease specific St. George's Respiratory Questionnaire (SGRQ) showed a better quality of life in the patients in the intervention group, but only in the activity component where there was

heterogeneity between the results of the two included studies. A potential reason for the absence of convincing effects on HRQoL is the limited use of COPD-specific instruments. Inconclusive results were observed on COPD-symptoms and use of other health care resources such as doctor and nurse visits. Self-management education reduced the need for rescue medication, and led to an increased use of courses of oral steroids and antibiotics for respiratory symptoms.

### Authors' conclusions

The data available for this review are insufficient for forming recommendations. Further research on the effectiveness of self-management programmes should be focussed on behavioural change evaluated in well designed randomised controlled trials with standardised outcomes designed for use in COPD patients, and with long follow-up time so that definite conclusions can be made.

## PLAIN LANGUAGE SUMMARY

**There is insufficient evidence to determine whether self-management education for people with chronic obstructive pulmonary disease (COPD) is effective, and more research is needed**

Chronic Obstructive Pulmonary Disease (COPD) represents a considerable burden to the individual and his or her loved ones, and represents an increasing health burden globally. COPD is characterised by frequent fluctuations, and repeated severe exacerbations are common. The idea of self-management is to prevent exacerbations by means of life style adaptation and to give patients tools to treat their exacerbations in an early stage. It is not clear how much difference education about self-management makes to people with COPD. The medical literature was systematically searched to obtain studies assessing the effects of self-management education in people with COPD. Two reviewers independently assessed each paper for methodological quality and extracted the data. The data from this review are insufficient to formulate recommendations regarding the effectiveness of self-management education programmes for COPD, because of the broad variation of outcome measures used and other limitations to generalisations in the current published literature.

## BACKGROUND

Chronic Obstructive Pulmonary Disease (COPD) is a serious public health problem worldwide. The prevalence, morbidity and mortality are expected to rise, especially in countries with a rapidly aging population and even in populations with reduced smoking rates (Feenstra 2001). A study published by the World Bank / World Health Organisation reported that COPD is likely to rise from being the 12th most burdensome disease in 1990 to the 5th in 2020 (Murray 1996). This will place an enormous burden on the healthcare system and will also cause a loss in health-related quality of life for many COPD patients. Treatment for individuals with COPD is often primarily aimed at improving airflow obstruction by bronchodilator- and anti-inflammatory therapy, despite indications that airflow obstruction is irreversible and apparent lack of effect of pharmacological interventions on the progressive decline in health status. Despite optimal pharmacological treatment many COPD patients experience substantial functional impairment (Wouters 1999). However, airflow obstruction correlates poorly with disease perception by the patient (Curtis 1994; Ketelaars 1996). COPD is a systemic inflammatory disease, and

besides airflow limitation and hyperinflation, due to loss of elastic recoil, as well as intrinsic airway narrowing, systemic deficits such as skeletal and respiratory muscle dysfunction are prominent features. There is a growing need for other forms of therapy for COPD-patients, not only to control and alleviate symptoms and complications of respiratory dysfunction, but also to teach them how to optimally carry out the activities of daily living (ATS 1981) in the face of their physiological impairment. As the benefits from pharmacological interventions (perhaps other than oxygen) are limited, it is essential that the relative merits of self-management education are fully evaluated and any successful components for this intervention clearly identified.

In the treatment of asthma, patient education and self-management programmes have proven to be successful, at least when combined with regular review, in reducing the economic burden of disease, as well as in improving quality of life and lung function (Gibson 2000; Gallefoss 1999a; Lahdensuo 1996; Klein 2001). In COPD, pulmonary rehabilitation has been proven to increase exercise tolerance and quality of life (Goldstein 1994). The drawback is that pulmonary rehabilitation programmes will normally

be more expensive and time consuming for both professionals and patients, than self-management programmes, and may be less widely available.

Worth 1996 was the first to describe the effectiveness of a self-management programme aimed at acquisition by the patient of self-management skills and behavioural change. Unfortunately, this pilot study was uncontrolled and studied a small sample of COPD patients (N = 21). Impressive reductions in the frequency of exacerbations and home visits by the family doctor were observed, but no changes in lung function were found. Several controlled trials have been conducted to evaluate the effectiveness of COPD education and self-management education programmes. This review is being conducted to address the impact of those programmes on health outcomes and health-care utilisation, and to assess the influence of programme characteristics on health outcomes.

## OBJECTIVES

- (1) To evaluate whether self-management education programmes in COPD leads to improved health outcomes.
- (2) To evaluate whether self-management education programmes in COPD leads to a reduction of health care utilisation.

## METHODS

### Criteria for considering studies for this review

#### Types of studies

Studies were included if they were randomised controlled trials or controlled clinical trials, which assessed the efficacy of self-management education in COPD patients. Studies focusing mainly on pulmonary rehabilitation were excluded as well as studies before 1985. In the period before 1985, treatment of COPD was not comparable with current practice.

#### Types of participants

Patients with a clinical diagnosis of COPD and not asthma as primary diagnosis were included.

#### Types of interventions

The interventions were categorised according to whether or not they involved COPD education and/or self-treatment guidelines i.e. an action plan.

Operational definition of COPD education: a programme which transfers information about COPD and treatment of COPD in

any of the following forms: written, verbal, visual or audio. Minimal education included the provision of written material alone or a short structured verbal interaction between a healthcare provider and a patient. However, it had to be embedded in a formal programme, in which the primary goal was to improve the knowledge and understanding of COPD. The educational programme might be directed towards smoking cessation, improving exercise, nutrition, self-treatment of exacerbations, inhalation technique or coping with the activities of daily living, or a combination of these. Operational definition of self-treatment guidelines (action plan): a written plan produced for the purpose of patient self-management of COPD exacerbations. It informs patients about when and how to adjust and/or start medication in case of an exacerbation.

#### Types of outcome measures

Any of the following outcomes: health-related quality of life scores; symptom scores; number and severity of exacerbations; courses of oral steroids or antibiotics; use of rescue medication; hospital admissions; emergency room visits; use of other health care facilities; days lost from work; lung function; and exercise capacity.

### Search methods for identification of studies

We identified studies from the following sources:

Cochrane Airways Group trials register derived from MEDLINE (January 1985 to October 2001), EMBASE (January 1985 to October 2001), CINAHL (January 1985 to October 2001), hand searched respiratory journals and meeting abstracts. We searched the databases using the following terms: 'Self-Care' AND 'Lung-Diseases-Obstructive' AND (COPD OR chronic obstructive pulmonary disease OR emphysema OR chronic bronchitis) AND ('Patient-Education' OR self management OR self-management OR self management). After obtaining articles for inclusion, we handsearched the bibliographic lists for additional papers. We also contacted research groups in this field for unpublished and non-registered ongoing trials.

### Data collection and analysis

#### STUDY ELIGIBILITY

Two reviewers independently coded the studies from the above searches based on the abstract/keywords/ title:

- (1) Include: Randomised Controlled Trial (RCT) or Controlled Clinical Trial (CCT), COPD, education/ self-management
  - (2) Possible RCT or CCT, but undetermined from the abstract
  - (3) Exclude: non-RCT or non-CCT, pulmonary rehabilitation
- Two investigators independently assessed study eligibility, study quality and intervention types. Agreement was examined; disagreement was resolved if possible by consensus, and otherwise by consultation with a third reviewer. We included articles if they were

randomised or controlled clinical trials of COPD education, and reported relevant outcomes.

#### ANALYSIS

We analysed outcomes as continuous and/or categorical variables using standard statistical techniques.

(A) For continuous outcomes, the weighted mean difference (WMD) or standardised mean difference (SMD) with 95% confidence intervals (CI) were calculated as appropriate

(B) Rates (relative risks(RR)/odds ratios(OR)) were pooled.

#### PRIMARY COMPARISON

The primary comparison was self-management versus usual care.

## RESULTS

### Description of studies

See: Table of included studies.

#### DOCUMENTED CHARACTERISTICS OF INCLUDED TRIALS

- (1) Demographics: age, gender, socio-economic status
- (2) Setting of intervention: primary care, hospital based, group education, individual education, community based
- (3) Involvement of the partner in the intervention
- (4) Duration of intervention: number of sessions, hours of teaching, time frame
- (5) Inclusion of physical exercise: type, location, duration
- (6) Sample size/power
- (7) Disease severity
- (8) Intermediate outcomes: COPD specific knowledge, skills, inhalation technique, smoking cessation, compliance

#### MISSING DATA

An attempt was made to contact the author to complete missing data. A reminder was sent if the author did not reply.

### Risk of bias in included studies

Two reviewers independently assessed the methodological quality of the included studies using the criteria list of [Jadad 1996](#).

The quality variables recorded in the criteria list of [Jadad](#) were:

- (1) procedure of allocation
- (2) withdrawals/drop-outs
- (3) blinding of patients and outcome assessment

In general, a maximum of five points could be obtained using this criteria list. As blinding is impossible when assessing behavioral interventions the maximum score is limited to three points. A higher score indicates better methodological quality.

In addition we described the general quality of the data in the studies in terms of sample size, quality of the outcome assessment, length of the follow-up period, etc.

## Effects of interventions

### INCLUSION

The search identified 395 (386 Cochrane database and nine hand searched) titles and abstracts that were screened to identify 33 potentially relevant articles about self-management education in COPD. We obtained full-text versions of these papers, and two reviewers independently assessed them. The reviewers included 12 articles describing eight randomised controlled trials and one controlled clinical trial.

Twenty-one articles were excluded for the following reasons: the design of the study was not a CCT or RCT (10); most of the patients had asthma as primary diagnosis (2); the studies were published before 1985 (2); the studies were primarily focused on pulmonary rehabilitation (6); or the outcome assessed was not appropriate (1).

### MISSING DATA

Replies were received from all authors who are listed in the acknowledgement section. However, not all authors could provide us with the additional requested information.

### QUALITY

The method to generate the sequence of randomisation was not clear in two of the eight randomised-controlled trials. The concealment of allocation was adequate in the other six RCTs. None of the interventions was double blind, because blinding of the patients with respect to study status is almost impossible in behavioural clinical trials. A description of withdrawals and dropouts was given in eight of the nine studies.

Five studies ([Gallefoss 1999a](#), [Gallefoss 1999b](#), [Gallefoss 2000a](#); [Emery 1998](#); [Watson 1997](#); [Littlejohns 1991](#); [Blake 1990](#)) scored the maximum number of three quality points, three studies ([Cockcroft 1987](#); [Gourley 1998](#), [Solomon 1998](#)); [Sassi-Dambron 1995](#)) two points and 1 ([Howland 1986](#)) study scored one point.

### INTERVENTIONS

Eight of nine studies described COPD self-management education compared with usual care. One study compared a rehabilitation programme without exercise component with general health education ([Sassi-Dambron 1995](#)). In four studies the education delivery mode consisted of group education; in four of individual education and one study used written education material only. One intervention also assessed the use of an action plan for self-treatment of exacerbations. The follow-up time was 12 months in four studies ([Gallefoss 1999a](#), [Gallefoss 1999b](#), [Gallefoss 2000a](#)); [Littlejohns 1991](#); [Howland 1986](#); [Blake 1990](#)), 10 months in one study ([Cockcroft 1987](#)), six months in three studies ([Gourley 1998](#), [Solomon 1998](#)); [Watson 1997](#); [Sassi-Dambron 1995](#)) and two months in one study ([Emery 1998](#)).

### COMPARISONS

Eight of the nine studies that compared self-management education with usual care as control have been included in this review.

### SUBJECTS / RECRUITMENT

A total number of 1295 patients were randomised in the eight studies; 1106 patients completed these studies. The drop out rates

ranged from 0% to 22.4%.

Five studies recruited their patients from outpatient clinics; one study recruited from general practice, one from the community and one study recruited patients from a mix of these settings.

#### OUTCOMES

Number of studies reporting outcome:

- (1) Health-related quality of life (HRQoL) (8)
- (2) Symptoms (2)
- (3) Number and severity of exacerbations (1)
- (4) Courses of oral steroids and/or antibiotics (3)
- (5) Use of rescue medication (1)
- (6) Hospital admissions (4)
- (7) Emergency room visits (1)
- (8) Use of other health-care facilities (3)
- (9) Days lost from work (2)
- (10) Lung function (4)
- (11) Exercise capacity (0)

#### HEALTH-RELATED QUALITY OF LIFE

Instruments for measurement of HRQoL differed widely among the studies. COPD-specific HRQoL by means of the St. George's Respiratory Questionnaire (SGRQ) was measured in two studies (Gallefoss 1999a; Watson 1997). SGRQ total scores and domain scores were all lower (indicating a better HRQoL) in the self-management education groups, but these differences did not reach clinical significance. The SGRQ-domain physical activity showed a significant and clinically relevant lower score (WMD -10, 95% CI: -18.5; -2.0); indicating a better HRQoL in the self-management education group, but there was significant heterogeneity between these two studies (Chi Sq 12.54  $p < 0.05$ ).

General HRQoL was measured with the Sickness Impact Profile (SIP) in three studies (Blake 1990; Littlejohns 1991; Emery 1998). Data from these three studies were not suitable for meta-analysis and showed incongruent results. One study (Emery 1998) reported significant improvement in total function measured by the SIP for the control group. However, Blake 1990 showed better physical function and total function in favour of the intervention group, and Littlejohns 1991 showed a significant greater improvement in physical function in the intervention group. Other studies used the Health Status Questionnaire 2.0 (Gourley 1998), the General Health Questionnaire (Cockcroft 1987) and a self-designed questionnaire (Howland 1986) (consisting of elements of the health locus of control scale, respiratory health questionnaire, Zung scales and SIP-scales) to measure general health status. In the latest two studies, general HRQoL was not significantly different between the self-management education and control group, although, Gourley 1998 showed significantly improved scores for the well-being dimension of the Health Status Questionnaire 2.0 in the intervention group.

#### SYMPTOMS

The effects of self-management education on COPD symptoms were examined in two studies (Solomon 1998; Watson 1997). Solomon 1998 assessed symptoms with different instruments: the

BORG-scale measuring breathlessness on a 12-point scale and the Global Assessment Scale measuring symptom severity on a 6-point scale. BORG-scale scores indicated a positive direction of effect of self-management education on breathlessness, although no statistical significant differences were observed. The Global Assessment Scale showed a reduction (not statistically significant) in symptom severity in the self-management education group while in the control group no reduction was observed. In the study by Watson 1997 patients scored their respiratory status in symptom diaries on a four point scale (usual, mild, moderate, severe). They found no significant differences between groups in the proportion of days rated as mild, moderate and severe.

#### NUMBER AND SEVERITY OF EXACERBATIONS

One study (Littlejohns 1991) reported the number of exacerbations. However, no comparison could be made because they only mentioned the number of acute exacerbations in the intervention group. The other studies did not use COPD exacerbation rates as an outcome measure.

Outcome measures such as (increased) symptoms, use of rescue medication, hospitalisations and use of health care facilities can serve as proxy variables for exacerbations of COPD because these variables indicate worsening of COPD.

#### COURSES OF ORAL STEROIDS AND/OR ANTIBIOTICS

Three studies assessed the use of oral corticosteroids for respiratory problems (Gallefoss 1999b; Watson 1997; Littlejohns 1991). Gallefoss 1999b reported that 69% of patients in the intervention group registered steroid courses compared to 44% in the control group, with a median of three and four courses recorded during the study year, respectively. Littlejohns 1991 reported that 49% of the patients in the intervention group used oral steroids and 37% in the control group during the study year. Meta-analysis of these studies showed an increased use of oral steroid courses in the educated patients (Relative Risk 1.39, 95% CI 1.02 to 1.91). Watson 1997 analysed from symptom diaries the days on prednisolone as percentage of the days recorded. The intervention group spent 15% of the days recorded on prednisolone compared to 9% in the control group.

Use of antibiotics for respiratory problems was assessed in two studies (Watson 1997; Littlejohns 1991). Littlejohns et al. reported that 79% of the patients in the intervention group were prescribed antibiotics during the study year compared to 52% in the control group. Watson 1997 analysed from symptom diaries the days on antibiotics as a percentage of the days recorded. The intervention group spent 10% of the days recorded on antibiotics compared with 4% in the control group. Treatment differences in both studies were statistically significant.

#### USE OF RESCUE MEDICATION:

One study (Gallefoss 1999b) reported on the use of short-acting B<sub>2</sub>-agonist as rescue medication. Use of rescue medication was coded as the Defined Daily Dosages (DDD) for comparison of medications within the same chemical therapeutic group. In this study the educated patients received less than half the amount

of rescue medication (median DDD = 125) compared with the control group (median DDD = 290). This reduction is statistically significant.

#### HOSPITAL ADMISSIONS

Four studies reported COPD related hospitalisations (Gallefoss 2000a; Cockcroft 1987; Solomon 1998; Littlejohns 1991). Overall, these studies did not observe significant differences between the self management education group and the control group. Meta-analyses of two studies (Gallefoss 2000a; Littlejohns 1991) reporting data about the number of patients with one or more admissions, showed a non-significant reduction of hospitalisations in favour of the treatment group (Relative Risk 0.80, 95% CI 0.43 to 1.50).

#### EMERGENCY ROOM VISITS

One study (Solomon 1998) reported the effect of self-management education on emergency room (ER) visits. This study showed no significant differences in ER-visits between the self-management education and the control group: four weeks prior to the last follow-up visit the median number of ER-visits per patient was 0.15 and 0.17 respectively for the active and control group.

#### USE OF OTHER HEALTH CARE FACILITIES

Doctor and nurse visits were reported in three studies and used for meta-analysis (Gallefoss 2000a; Solomon 1998; Watson 1997). Two studies (Gallefoss 2000a; Solomon 1998) reported a reduction of these visits, and one study (Watson 1997) reported an increase in the self-management education group compared to the control group. However, this last study (Watson 1997) was only based on three and four visits in the intervention and control group respectively. There was heterogeneity among these studies (Chi Sq 8.48  $p < 0.05$ ). The overall weighted mean difference did not reach statistical significance at -0.36 (95% CI: -0.75 to 0.03) visits per year in favour of the treatment group.

#### DAYS LOST FROM WORK

One study (Gallefoss 2000a) reported no difference between groups in days lost from work. Almost 50% of the COPD-patients in this study were employed, and only three out of 14 and two out of 13 in the intervention and control group respectively, reported an absence from work. Blake 1990 found no between group differences in restricted activity days. Restricted activity days were defined as days in which work was missed or in which activities were significantly reduced because of health problems. So, this study assessed loss from paid and unpaid work which can be seen in the larger numbers reported.

#### LUNG FUNCTION

Lung function was assessed as the forced expiratory volume in 1 second as percent predicted of the normal value (FEV1%-predicted). Of the four studies (Gallefoss 2000a; Emery 1998; Watson 1997; Littlejohns 1991) which reported data on FEV1%-predicted three were entered as post treatment levels. One study (Littlejohns 1991) only reported mean within-group differences so the results were entered as % change in predicted FEV1. No significant difference in deterioration of FEV1%-predicted was seen

between the intervention and control groups (Standardised Mean Difference -0.01, 95% CI -0.24 to 0.22).

#### EXERCISE CAPACITY

No study evaluated the effect of self-management education on exercise capacity.

## DISCUSSION

This review systematically evaluated eight studies of self management education for patients with COPD compared to usual care. The studies showed no effect of self-management education on hospital admissions, ER visits, days lost from work and lung function. Inconclusive results were observed on HRQoL. A potential reason for the absence of convincing effects on HRQoL is the limited use of COPD specific instruments. General HRQoL instruments may not be sensitive enough to detect differences in COPD patients. The studies using the disease specific SGRQ showed a trend towards better quality of life in the educated patients.

Days lost from work might not be an adequate outcome in COPD patients because many are in the older age groups and often retired. Since, in most COPD studies a minority of the patients undertake paid work, we think restricted activity days, indicating days in which the normal activities are reduced by the disease, would be a better outcome measure.

We did not expect to find an effect of self management education on lung function. The accelerated decline in pulmonary function in COPD patients is very difficult to affect. Smoking cessation is the only treatment that has so far been able to reduce this accelerated decline (Fletcher 1977).

Inconclusive results were observed on COPD symptoms and use of other health care facilities, such as doctor and nurse visits.

Self-management education did reduce the need for rescue medication. This may indicate that self-management education leads to a better disease control in COPD-patients. However, use of rescue medication was measured in only one study, so the strength of evidence for this observation is poor.

Self-management education led to an increased use of courses of oral steroids and/or antibiotics for respiratory symptoms. This apparent paradox does not mean that self management education leads to worsening of respiratory symptoms. Seemungal 2000 showed in a well designed study that in COPD patients 50% of the exacerbations were not reported to a doctor. Considering this, it is most likely that the educated patients were more conscious of worsening of their symptoms, and that the threshold for seeking help was decreased. For the future, it will be important to investigate potential beneficial effects (on HRQoL and/or prevention of hospital admissions) but also side effects of increased use of prednisolone over the long term. But to answer these questions there is a need for large studies with a long follow-up period.



The review also identified a number of limitations in the current published literature, which need to be considered:

(1) The studies in this review assessed a broad spectrum of outcome parameters with different follow-up periods. Meta-analytic comparisons could not readily be made. For the main part, we have been limited to describing the effects of self-management education on the different outcome parameters.

(2) The COPD population was defined in varying detail, and included different diagnostic criteria. This could have led to heterogeneity in disease severity. The mode of the self management education programmes varied from group education, individual education, to written education material only. Use of an action plan for self treatment of exacerbations was assessed in one study only. Considering the evidence in a review of asthma self-management (Gibson 2000), we believe that this could be an important part of a self-management education programme. Gibson 2000 showed positive effects on health outcomes of self management interventions in adult asthma patients, using an action plan and regular health practitioner follow-up whereas interventions without action plans were not always of obvious benefit.

(3) The span of 14 years over which the trials were conducted will have introduced changes to the educational contents and the mode of delivery and background therapy. Most studies were not aimed at improving self management skills or behavioural change. However, we believe that self management involves the transfer of knowledge as well as the acquisition by the patient of certain important skills which ultimately leads to a change in their behaviour. This is the only way education can have a long term impact on the daily life of the COPD patient because knowledge about the disease does not directly implicate behavioural change. A theoretical model of behaviour and behavioural change, for example the ASE-model (Bandura 1986), can be very helpful in designing a self-management programme. The programme should be focused on the core elements of behaviour change in the theoretical model (for example: enhancing self-efficacy expectations or social support).

Besides the aforementioned limitations in the literature, absence of positive results in most of the self management education studies could also be caused by non-reversibility of the disease. COPD is a less variable disease than for example asthma, and it is a disease where the scope for therapeutic interventions is much more limited and therefore it is intrinsically harder to show positive results.

## AUTHORS' CONCLUSIONS

### Implications for practice

The data abstracted by this review form an insufficient basis for the formulation of recommendations regarding the effectiveness of self-management programmes for COPD patients, because of the broad spectrum of outcome parameters and other important limitations in the current published literature.

### Implications for research

Future studies with sufficient sample size and longer follow-up time should focus on:

- (1) the effects of self-management programmes that focus on acquisition of self-management skills and behavioural change;
- (2) the core elements of such a self management programme;
- (3) the effects of an action plan; and
- (4) cost/benefit analysis of prednisolone courses in self management educated patients in the long run.

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**Monnikhof** {unpublished data only}

COPE-study. Ongoing study 1–5–1999.

**Tregonning** {unpublished data only}

Randomised controlled trial of a home-based programme of exercise and education in COPD. Ongoing study ?.

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\* Indicates the major publication for the study

## CHARACTERISTICS OF STUDIES

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

#### Blake 1990

Methods	RCT FUP = 12 m control: usual care	
Participants	Eligible: 109 Randomised: 94 Completed: 83 Mean age: I: 63 (?) yrs C: 64 (?) yrs Sex (% male): I: 80% C: 82% Diagnosis COPD: respiratory symptoms + abnormal pulmonary function: - daily cough and/or shortness of breath over the preceding six months - FEV1/VC < 75 % or FVC%pred < 75% Recruitment: outpatients Major exclusions: - FEV1%pred: ? FEV1/VC: I: 46% C: 43%	
Interventions	Mode: individual education + patient brochure + audiotape Content: stress management; relaxation exercise; meditation; guided imagery focusing on breathing; social and recreational activities; communication skills Duration: 1 to 4 hrs Action Plan: N	
Outcomes	-Health status: - SIP - Hospital days - Bed-disability days - Restricted-activity days - Physician visits	
Notes	-Follow-up data for 6 and 12 months -19% never smokers	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Unclear risk	D - Not used

**Cockcroft 1987**

Methods	RCT FUP = 10 m control: usual care	
Participants	Eligible: 92 Randomised: 75 Completed: 73 Mean age: I: 69 (?) yrs C: 71 (?) yrs Sex (% male): I: 69 C: 67 Diagnosis of COPD: all pts suffering from chronic respiratory disability that was caused mainly by chronic obstructive airways disease; at least admitted twice during previous 3 yrs. Recruitment: outpatients Major exclusions: disability not caused by a respiratory condition FEV1%pred:? FEV1/VC:?	
Interventions	Mode: individual by home visits of a respiratory health worker Content: COPD knowledge; symptoms; coping behaviour Duration: about 10 hrs Action Plan: N	
Outcomes	-Health Status -GHQ -Hospital admissions -Deaths -Knowledge about medication and condition -VAS-scales concerning physical and psychological aspects	
Notes		
<b><i>Risk of bias</i></b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Unclear risk	B - Unclear

**Emery 1998**

Methods	RCT FUP = 2 m control: usual care	
Participants	Eligible: 92 Randomised: 50 Completed: 49 Mean age: I: 67 (6) yrs C: 67 (6) yrs Sex (% male): I: 42 C: 48 Diagnosis of COPD: stable COPD; > 50 yrs; FEV1/VC < 70; > 6 months clinical symptoms of COPD	

**Emery 1998** (Continued)

	Recruitment: outpatients + GP-patients + advertisements + word of mouth Major exclusions: significant cardiac disease; other diseases affecting exercise tolerance or learning skills last 3 months; asthma without fixed obstruction FEV1%pred: I: 43 (18) C: 39 (16) FEV1/VC:?	
Interventions	Mode: group education Content: COPD knowledge;therapy; coping; interpreting pulmonary function tests; understanding of arterial bloodgases; stressmanagement Duration: 26 hrs Action Plan: N	
Outcomes	-Health status - SIP - HRQoL-MHLC -Health knowledge test - FEV1%pred	
Notes	The third arms was disregarded, because it was focussed on pulmonary rehabilitation	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Low risk	A - Adequate

**Gallefoss 1999a**

Methods	RCT FUP = 12 m control: usual care	
Participants	Eligible: 68 Randomised: 62 Completed: 53 Mean age: I: 57 (9) yrs C: 58 (10) yrs Sex (% male): I:48 C: 52 Diagnosis of COPD: FEV1pred >= 40% and FEV1pred < 80% Recruitment: outpatients Major exclusions: any serious disease FEV1%pred: I:59 (9) C: 56 (11) FEV1/VC: I: 55(9) C:52(10)	
Interventions	Mode:patient brochure + group sessions Content:COPD knowledge; medication; symptoms; exacerbations; inhalation technique;smoking cessation; relaxation; coping Duration: max 6.5 hrs Action Plan: Y	

**Gallefoss 1999a** (Continued)

Outcomes	-Health status -SGRQ -other HRQoL instruments	
Notes	- The same study as Gallefoss 1999b and 2000	
<b><i>Risk of bias</i></b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Low risk	A - Adequate

**Gallefoss 1999b**

Methods	RCT FUP = 12 m control: usual care	
Participants	Eligible: 68 Randomised: 62 Completed: 53 Mean age: I: 57(9) C: 58(10) Sex (% male): I:48 C: 52 Diagnosis of COPD: FEV1%pred >= 40% and FEV1%pred<80% Recruitment: outpatients Major exclusions: any serious disease FEV1%pred: I: 59 (9) C:56 (11) FEV1/VC: I: 55 (9) % C: 52 (10) %	
Interventions	Mode:patient brochure + group sessions Content:COPD knowledge; medication; symptoms; exacerbations; inhalation technique;smoking cessation;relaxation; coping Duration: max 6.5 hrs Action Plan: Y	
Outcomes	-Compliance -Courses of steroids -Use rescue medication	
Notes	- The same study as Gallefoss 1999a and 2000	
<b><i>Risk of bias</i></b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Low risk	A - Adequate



### Gallefoss 2000a

Methods	RCT FUP = 12 m control: usual care	
Participants	Eligible: 68 Randomised: 62 Completed: 53 Mean age: I: 57 (9) yrs C: 58(10) yrs Sex (% male): I: 48% C: 52% Diagnosis of COPD: FEV1pred >= 40 % and FEV1pred < 80% Recruitment: outpatients Major exclusions: any serious disease FEV1%pred: I: 59 (9) C: 56 (11) FEV1/VC: I: 55 (9) % C: 52 (10) %	
Interventions	Mode: patient brochure + group sessions + individual sessions by nurse and physiotherapist Content: COPD knowledge; medication; symptoms; exacerbations; inhalation technique; smoking cessation, relaxation; coping Duration: max 6,5 hrs Action Plan: Y	
Outcomes	-Health status - SGRQ - Hospital admissions - Days lost from work - GP-consultation - FEV1%pred	
Notes	- 6% were never smokers - The same study as Gallefoss 1999a and 1999b	
<b><i>Risk of bias</i></b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Low risk	A - Adequate

### Gourley 1998

Methods	RCT FUP = 6 m control: usual care	
Participants	Eligible: ? Randomised: 98 Completed: 98 Mean age: I: 69 (6) yrs C: 69 (9) yrs Sex (% male): I: 100 C: 100 Diagnosis of COPD: COPD ATS criteria; at least one MDI; > 40 yrs	

**Gourley 1998** (Continued)

	Recruitment: outpatients Major exclusions: life expectancy < 6 months; hospitalisation or ER-visits during past 2 wks; lung infection past 2 wks; decompensated CHF class 3 or 4; other lung disease except for committant asthma FEV1%pred:? FEV1/VC: ?	
Interventions	Mode: individual verbal education Content: COPD knowledge; therapy; coping Duration: 3 hrs Action Plan: N	
Outcomes	- Health status - HSQ2 - Patient satisfaction - Disease knowledge - Disease management knowledge	
Notes	- The same study as no. 33	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Unclear risk	D - Not used

**Howland 1986**

Methods	CCT FUP = 12 m control: usual care	
Participants	Eligible: 923 Randomised: 659 Completed: 538 Mean age: I: 59 (?) yrs C: 60 (?) yrs Sex (% male): I: 54% C: 51% Diagnosis COPD: presence of COAD; FEV1/FVC < 60% or between 60 to 70% with chronic symptoms of cough, weezing or breathlessness Recruitment: community patients Major exclusions: - FEV1%pred: ? FEV1/VC: ?	
Interventions	Mode: group education (one for the mildly and one for the severely impaired patients) Content: Severe group: COPD knowledge; nutrition; exercise; smoking cessation; Mild group: emphasized prevention and impairment oa smoking cessation, building and maintaining physical endurance and reducing stress Duration: Severe group = 12 hrs; Mild group = 6 hrs	

**Howland 1986** (Continued)

	Action Plan: N	
Outcomes	-Health status measured by a self designed questionnaire - FEV1%pred	
Notes	- They assessed a severe and a mild group of patients	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Unclear risk	D - Not used

**Littlejohns 1991**

Methods	RCT FUP = 12 m control: usual care	
Participants	Eligible: 166 Randomised: 152 Completed: 133 Mean age: I: 63 (8) yrs C: 63 (7) yrs Sex (% male): I: 67 C: 63 Diagnosis of COPD: previously documented chronic airflow obstruction; 30 to 75 yrs; FEV1%pred < 60%; stable state Recruitment: outpatients Major exclusions: other major disease; change in medication at least six weeks before recruitment FEV1%pred: I: 45 (22) C: 50 (23) FEV1/VC:?	
Interventions	Mode: individual by respiratory health worker Content: COPD knowledge; inhalation technique; impairment, disability and handicap Duration: ? Action Plan: N	
Outcomes	-Health status - SIP -No. of exacerbations -Courses of oral steroids -Courses of antibiotics -Hospital admissions -FEV1%pred -Exercise capacity -6MW	
Notes		

Littlejohns 1991 (Continued)

<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Low risk	A - Adequate

Sassi-Dambon 1995

Methods	RCT FUP = 6 m control:general health education
Participants	Eligible: 143 Randomised: 98 Completed: 76 Mean age: I: 68 (8) C: 67 (8) Sex (% male): I: 57 C: 53 Diagnosis of COPD: patients report that he or she felt limited in physical function secondary to dyspnea Recruitment: outpatients+ GP-patients+community patients Major exclusions: participation in pulmonary rehabilitation previous 2 years FEV1%pred: I: 50 (21) C:50 (23) FEV1/VC: ?
Interventions	Mode: group education (I: rehab without exercise C: general health education lectures) Content: ? Duration: 6 sessions Action Plan: N
Outcomes	-Health status -QWB -Symptoms -Exercise capacity -6MW
Notes	-Comparison of two interventions. No usual care group.

<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Allocation concealment (selection bias)	Unclear risk	B - Unclear

### Solomon 1998

Methods	RCT FUP = 6 m control: usual care	
Participants	Eligible: 193 Randomised: 128 Completed: 98 Mean age: I: 69 (6) yrs C: 69 (9) yrs Sex (% male): I: 100 C:100 Diagnosis of COPD: COPD ATS criteria; at least one MDI; > 40 yrs Recruitment: outpatients Major exclusions: life expectancy < months; hospitalisation or ER-visits during past 2 wks; lung infection past 2 wks; decompensated CHF class 3 or 4; other lung disease except for committant asthma FEV1%pred:? FEV1/VC:?	
Interventions	Mode: individual verbal education Content: COPD knowledge; therapy; coping Duration: 3 hrs Action Plan: N	
Outcomes	-Control of disease -Symptoms -Hospital admissions -ER visits -Use of other health care facilities	
Notes		
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Low risk	A - Adequate

### Watson 1997

Methods	RCT FUP = 6 m control: usual care	
Participants	Eligible: 93 Randomised: 69 Completed: 56 Mean age: I: 68 (10) yrs C: 67 (8) yrs Sex (% male): I: 62 C: 67 Diagnosis of COPD: COPD (ATS crriteria) as major limiting disease; smoking history > 10 pack years; FEV1%pred < 65%; FEV1/VC < 70 %; bronchodilator therapy Recruitment: GP-patients	

Watson 1997 (Continued)

	Major exclusions: asthma (onset < 35 yrs) as primary diagnosis; cardiac disease as primary diagnosis, another functionally limiting disease (except cor pulmonale) affecting mortality FEV1%pred: I: 37 (14) C: 36 (16) FEV1/VC: I: 52 (25) C: 48 (15)	
Interventions	Mode: action plan and patient brochure Content: COPD knowledge; exercise; smoking cessation; coping (controlling breathlessness); nutrition Duration: < 1 hr Action Plan: Y	
Outcomes	-Health Status - SGRQ -Symptoms -GP-visits -Courses of prednisolone -Courses of antibiotics -FEV1%pred	
Notes		
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Allocation concealment (selection bias)	Unclear risk	B - Unclear

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

Study	Reason for exclusion
Ashikaga 1980	<1985; asthma + COPD
Brough 1982	< 1985; methodological flaws
Brundage 1993	No CCTor RCT
De Assuncao 1999	No CCTor RCT; asthma
Devine 1996	No CCT or RCT: meta-analysis
Gallefoss 2000b	No adequate outcome
Garrod 1997	Abstract only; primarily focused on pulmonary rehabilitation

(Continued)

Grosbois 1996	No CCT or RCT; pulmonary rehabilitation
Hausen 1998	No CCT or RCT; 76% asthma
Hausen 1999	No CCT or RCT
Janelli 1991	No CCT or RCT
Muller 1996	No CCT or RCT; asthma + COPD group
Ries 1995	Primarily focused on pulmonary rehabilitation
Scherer 1996	No CCT or RCT
Scherer 1998	Primarily focused on pulmonary rehabilitation; no adequate outcome
Toshima 1990	Primarily focused on pulmonary rehabilitation; the same article as refnr. 83
Tougaard 1992	About 90% were asthma patients
Tougaard 1993	The same article as refnr. 100
van den Broek 1995	Primarily focused on pulmonary rehabilitation
Wedzicha 1998	Primarily focused on pulmonary rehabilitation
Worth 1997	No CCT

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

#### Bourbeau

Trial name or title	Living Well with COPD
Methods	
Participants	Eligible: 469 Randomised: 191 Completed: 165 Mean age: I: 69 yrs C: 70 yrs Sex (% male): I: 52 C: 59 Diagnosis COPD: smoking history (> 10 packyears), age 50+, FEV1 post 25-75% of pred, no previous diagnosis of asthma, terminal disease, dementia or L CHF, no participation in pulm. rehabilitation in the prior year, no long term care facility stays Recruitment: outpatients Follow-up: 12 m

**Bourbeau** (Continued)

Interventions	Mode: individual education; skill training; self help workbooks; learning video; supportive telephone calls Content: knowledge of COPD; inhalation technique; energy conservation; breathing and coughing techniques; relaxation and exercise; symptom control; adopting a healthy lifestyle; coping; oxygen therapy Duration: 8 hrs Action Plan: Y Exercercise program: Y
Outcomes	-Health status: -SGRQ -Exacerbations -Courses of oral steroids -Courses of antibiotics -Hospital admissions - Use of other health care facilities - Lung function -Exercercise capacity: -6MW
Starting date	1-2-1998
Contact information	J. Bourbeau Respiratory Epidemiology Unit Lady Meridith House 1110 Pine Avenue West Montreal, Quebec, Canada H3A 1A3 jean.bourbeau@mcgill.ca
Notes	

**Monninkhof**

Trial name or title	COPE-study
Methods	
Participants	Eligible: 509 Randomised: 246 Completed: ? Mean age: I: 64 yrs C: 64 yrs Sex (% male): I: 85.5 C:83.7 Diagnosis COPD: clinical stable COPD, smoking history, age 40-75; FEV1%pred 25-75%; FEV1/VC <= 60; reversibility of FEV1%pred < =12 % Recruitment: outpatients Follow-up: 24 m



**Monninkhof** (Continued)

Interventions	Mode: group education; skill training; self help workbook; Content: knowledge of COPD; inhalation technique; relaxation and exercise; symptom control; coping; nutrition; communication Duration: 7.5 hrs Action Plan: Y Exercise program: Y
Outcomes	- Health status: - SGRQ - Euroqol-5D -Symptoms -Exacerbations - Courses of oral steroids -Courses of antibiotics - Use of rescue medication -Hospital admissions - Use of other health care facilities -Lung function - Exercise capacity: -6MW
Starting date	1-5-1999
Contact information	E. Monninkhof COPE-study Department of lung diseases Medisch Spectrum Twente P.O. Box 50000 7500 KA Enschede The Netherlands emonninkhof@introweb.nl
Notes	

**Tregonning**

Trial name or title	Randomised controlled trial of a home-based programme of exercise and education in COPD
Methods	
Participants	Eligible: ? Randomised: 141 Completed: 117 Mean age: I: 73 yrs C: 73 yrs Sex (% male): I: 66 C: 60 Diagnosis COPD: stable COPD, FEV1 < 80%; FEV1/VC < 70 Recruitment: hospital patients Follow-up: 6 m

**Tregonning** (Continued)

Interventions	Mode: individual education; self help workbook; supportive telephone calls Content: knowledge of COPD; coping; nutrition; smoking cessation; relaxation and exercise; communication; flexible use of medication Duration: 3 visits Action Plan: N Exercise program: Y (home)
Outcomes	-Health status: -CRDQ -Symptoms -Exercise capacity: -6MWT
Starting date	?
Contact information	M. Tregonning North Bristol NHS Trust Bristol UK
Notes	

**Worth**

Trial name or title	AFBE-study
Methods	
Participants	Eligible: 102 Randomised: 89 Completed: 86 Mean age: 62 yrs Sex (% male): 76% Diagnosis COPD: clinical stable COPD; age 45-75 FEV1%pred. 30-80%, FEV1 /VC <= 70% FEV1 -reversibility <= 12% Recruitment: outpatients Follow-up: 12 m
Interventions	Mode: group education ; skill training Content: knowledge of COPD; inhalation technique; monitoring of disease; effects/side effects of medication; self adjustment of medication; behaviour during exacerbations Duration: 6 hrs Action Plan: Y Exercise program: N

**Worth** (Continued)

Outcomes	-Health status: - SGRQ - SF-36 -Symptoms -Exacerbations -Courses of oral steroids -Courses of antibiotics - Use of rescue medication -Lung function
Starting date	1-8-1999
Contact information	Y. Dhein, H. Worth Medical Department Fuerth Hospital Jakob-Henle-Str. 1 90766 Fuerth Germany med1@klinikum-fuerth.de
Notes	

## DATA AND ANALYSES

### Comparison 1. Self-management versus control

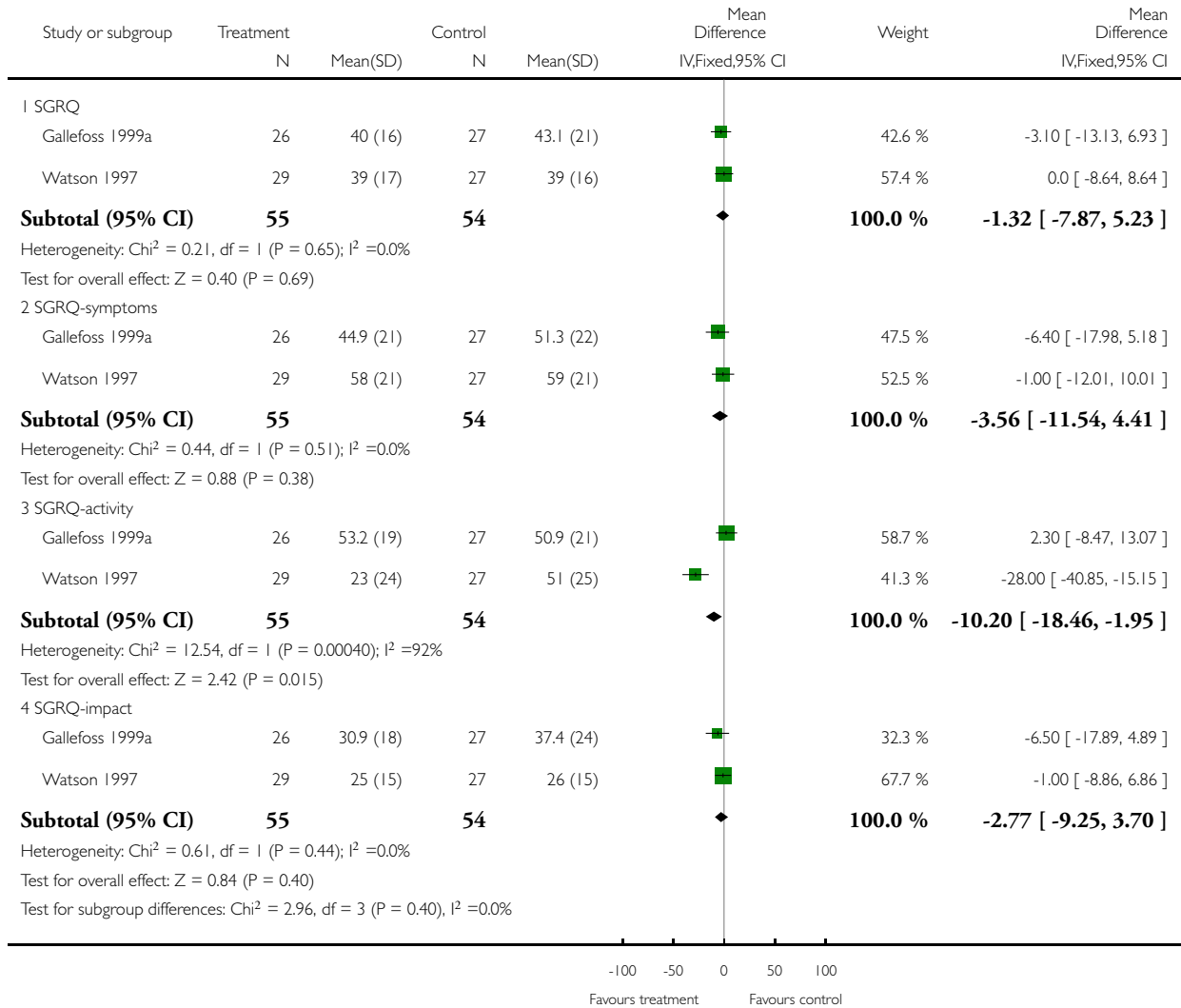
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 HRQOL: SGRQ Total + domains	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
1.1 SGRQ	2	109	Mean Difference (IV, Fixed, 95% CI)	-1.32 [-7.87, 5.23]
1.2 SGRQ-symptoms	2	109	Mean Difference (IV, Fixed, 95% CI)	-3.56 [-11.54, 4.41]
1.3 SGRQ-activity	2	109	Mean Difference (IV, Fixed, 95% CI)	-10.20 [-18.46, -1.95]
1.4 SGRQ-impact	2	109	Mean Difference (IV, Fixed, 95% CI)	-2.77 [-9.25, 3.70]
3 Doctor and nurse visits: mean number per person per year	3	197	Mean Difference (IV, Fixed, 95% CI)	-0.36 [-0.75, 0.03]
3.1 doctor and nurse visits: mean number per person per year	3	197	Mean Difference (IV, Fixed, 95% CI)	-0.36 [-0.75, 0.03]
4 Exercise capacity: 6MW	1	133	Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Lung function: FEV1%pred	4	290	Std. Mean Difference (IV, Fixed, 95% CI)	-0.01 [-0.24, 0.22]
6 Days lost from work: mean number per person per year	2	121	Mean Difference (IV, Fixed, 95% CI)	-17.5 [-50.05, 15.05]
7 General HRQOL: SIP Total score	3	249	Mean Difference (IV, Fixed, 95% CI)	0.30 [-4.00, 4.60]
8 General HRQoL: SIP physical	2	201	Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9 General HRQoL : SIP psychosocial	2	201	Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10 Patients using at least one course of oral steroids	2	186	Risk Ratio (M-H, Fixed, 95% CI)	1.39 [1.02, 1.91]
11 Hospital admissions: one or more	2	195	Risk Ratio (M-H, Fixed, 95% CI)	0.80 [0.43, 1.50]

## Analysis 1.1. Comparison 1 Self-management versus control, Outcome 1 HRQOL: SGRQ Total + domains.

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 1 HRQOL: SGRQ Total + domains

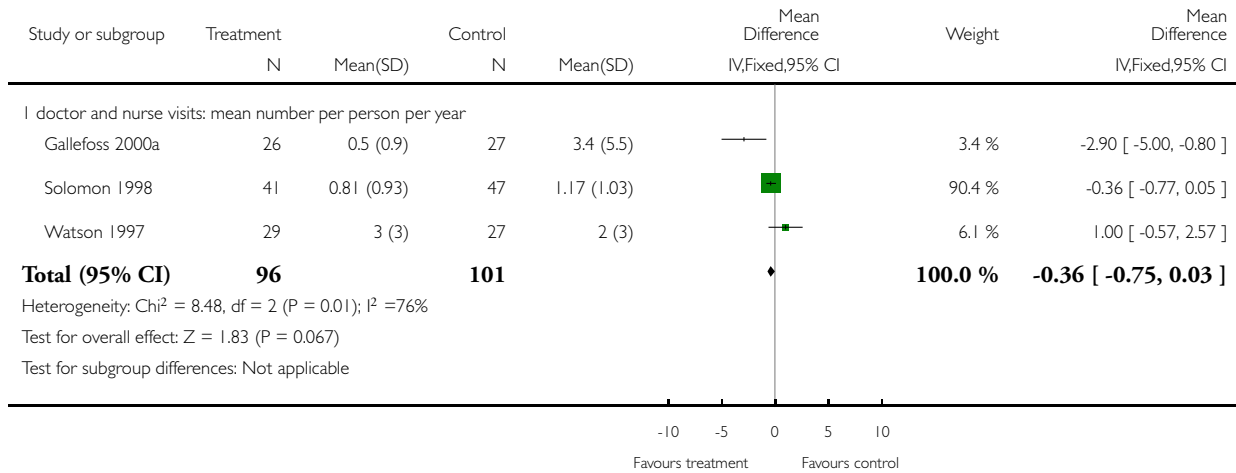


**Analysis 1.3. Comparison 1 Self-management versus control, Outcome 3 Doctor and nurse visits: mean number per person per year.**

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 3 Doctor and nurse visits: mean number per person per year

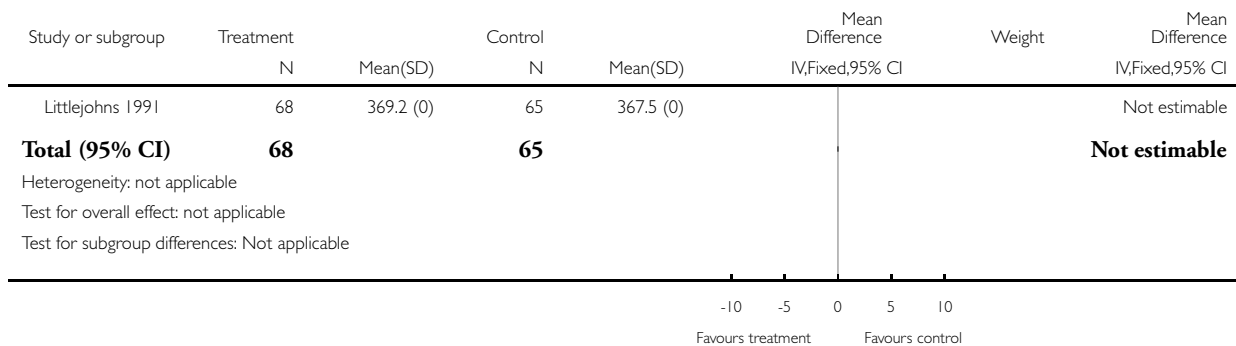


**Analysis 1.4. Comparison 1 Self-management versus control, Outcome 4 Exercise capacity: 6MW.**

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 4 Exercise capacity: 6MW

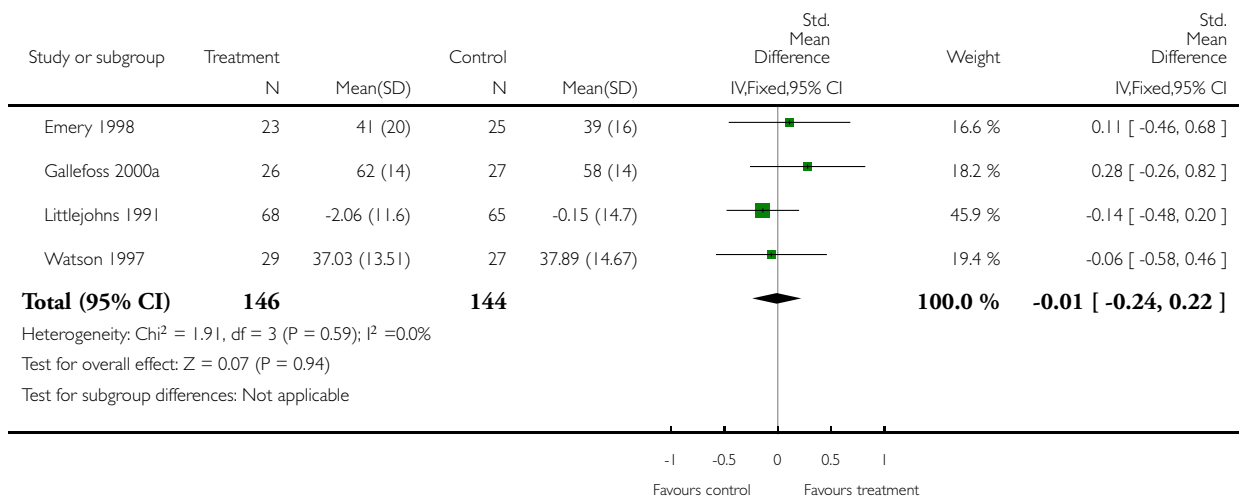


### Analysis 1.5. Comparison 1 Self-management versus control, Outcome 5 Lung function: FEV1%pred.

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 5 Lung function: FEV1%pred

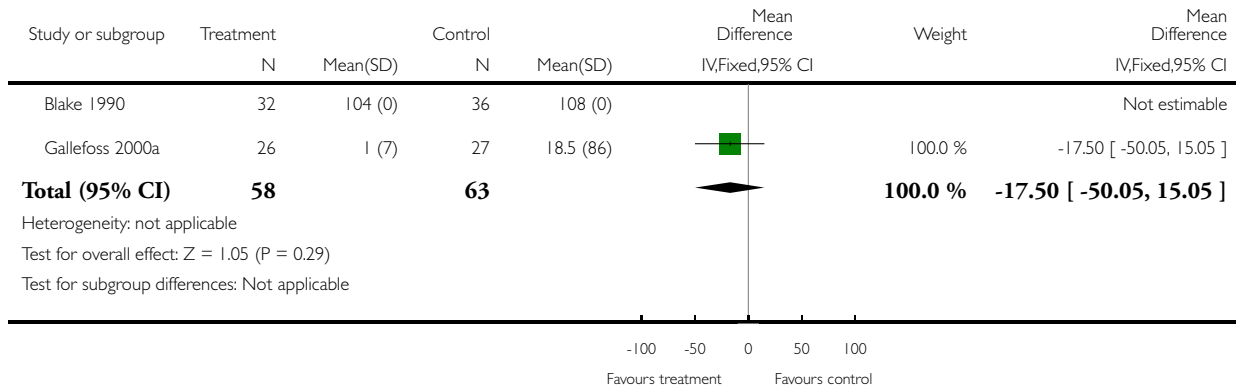


**Analysis 1.6. Comparison 1 Self-management versus control, Outcome 6 Days lost from work: mean number per person per year.**

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 6 Days lost from work: mean number per person per year

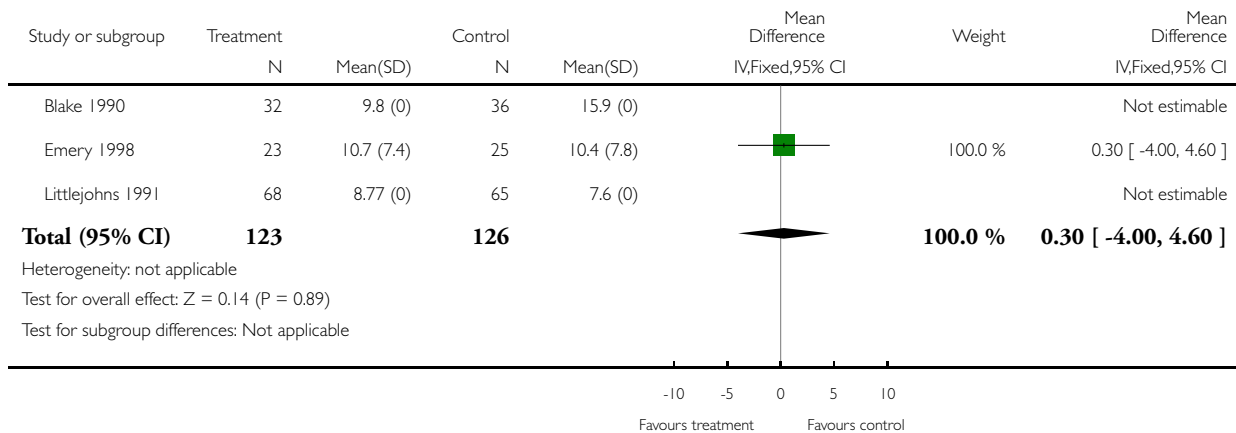


**Analysis 1.7. Comparison 1 Self-management versus control, Outcome 7 General HRQOL: SIP Total score.**

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 7 General HRQOL: SIP Total score



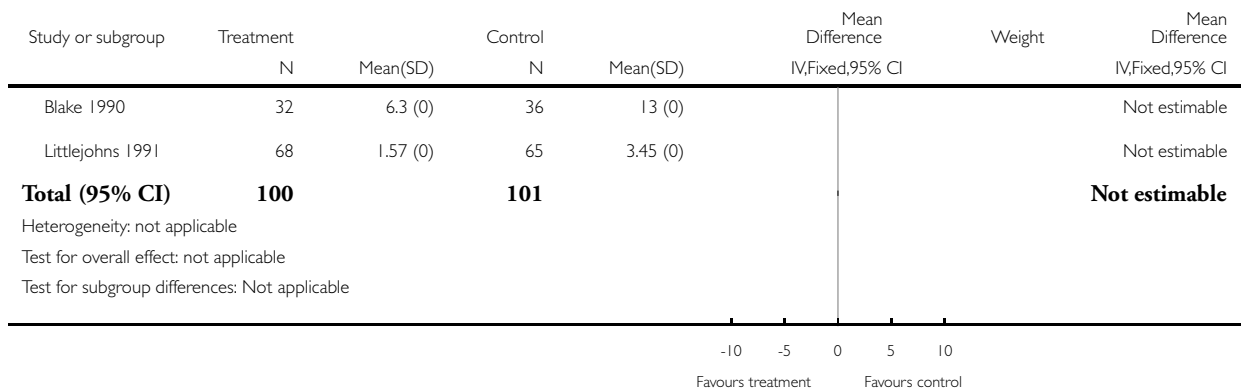


**Analysis 1.8. Comparison 1 Self-management versus control, Outcome 8 General HRQoL: SIP physical.**

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 8 General HRQoL: SIP physical

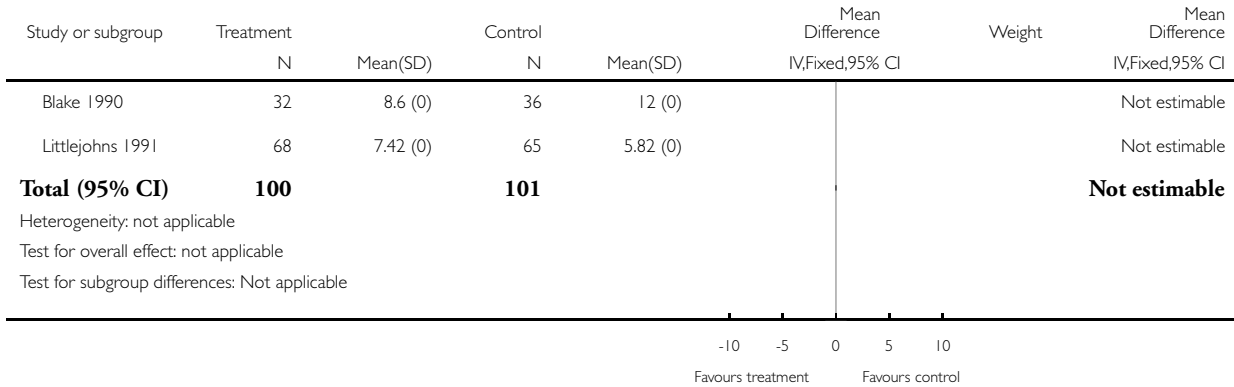


### Analysis 1.9. Comparison 1 Self-management versus control, Outcome 9 General HRQoL : SIP psychosocial.

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 9 General HRQoL : SIP psychosocial

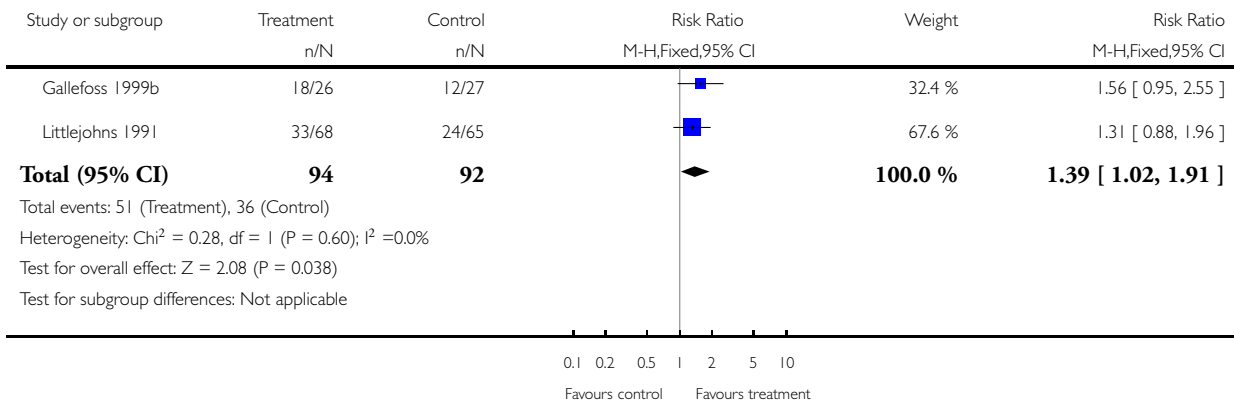


### Analysis 1.10. Comparison 1 Self-management versus control, Outcome 10 Patients using at least one course of oral steroids.

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 10 Patients using at least one course of oral steroids

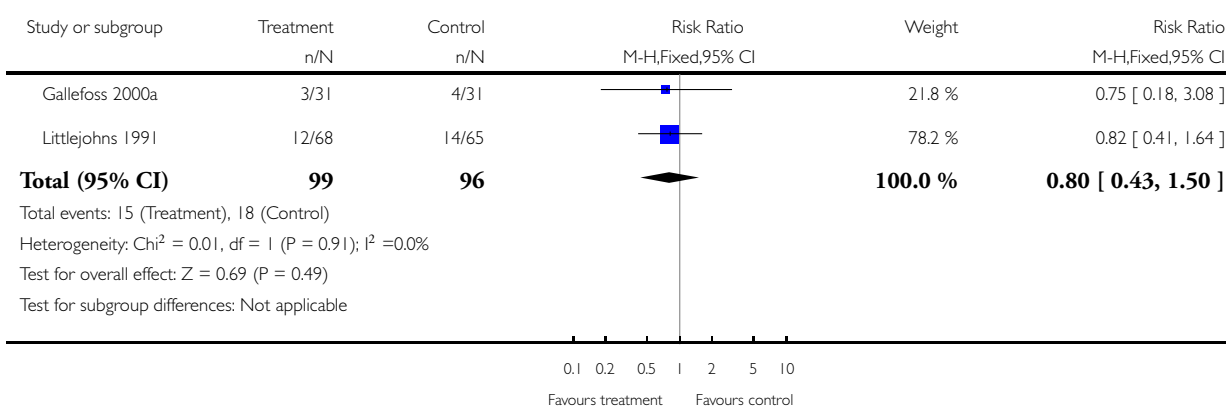


## Analysis 1.11. Comparison 1 Self-management versus control, Outcome 11 Hospital admissions: one or more.

Review: Self-management education for chronic obstructive pulmonary disease

Comparison: 1 Self-management versus control

Outcome: 11 Hospital admissions: one or more



## FEEDBACK

### Error quoting dosage figure

#### Summary

In the results section “Use of Rescue Medication”, the Gallefoss 1999 study is quoted in this review as follows:

“In this study the educated patients received less than half the amount of rescue medication (median DDD = 125) compared with the control group (median DDD=209)”.

In figure 3 of the original Gallefoss paper it is reported that in the control group the median DDD= 290 not 209.

#### Reply

I agree with the comment of Susan Varney. The median DDD rescue medication in the control group should be 290 not 209. This should be changed.

## Contributors

Varney S

## WHAT'S NEW

Last assessed as up-to-date: 20 August 2007.

Date	Event	Description
10 May 2017	Amended	Converted to new review format.

## HISTORY

Protocol first published: Issue 2, 2001

Review first published: Issue 1, 2003

Date	Event	Description
16 July 2002	New citation required and conclusions have changed	Substantive amendment

## CONTRIBUTIONS OF AUTHORS

E. Monninkhof:

Designed and co-ordinated the review

All steps belonging to data collection for the review

Data management for the review

Analysis of data

Interpretation of data

Writing the review

Updating the review

P. van der Valk:

Screened search results against inclusion criteria

Screened retrieved papers against inclusion criteria

Independent arbitrator to reach "consensus agreement"

Provided a clinical perspective

J. van der Palen:

Conceived the review

Screened search results against inclusion criteria  
Screened retrieved papers against inclusion criteria  
Abstracted data  
Obtained and screened data from unpublished studies  
Appraised quality of papers  
Supported datamanagement and data-analysis  
Provided a methodological perspective  
G. Zielhuis:  
Designed the review  
Provided advice on data-analysis  
Provided a methodological perspective  
Provided general advice on the review  
Critically revised the review  
C. van Herwaarden:  
Provided a clinical perspective  
Critically revised the review  
Provided a policy perspective  
M. Partridge:  
Provided a policy perspective  
Provided a clinical perspective  
Provided general advice on the review  
EH Walters:  
Editorial support throughout.

## **DECLARATIONS OF INTEREST**

Netherlands Asthma Foundation, Glaxo Wellcome BV, Boehringer Ingelheim provided funding for this review but these parties were in no way be able to influence the results of the review.

## SOURCES OF SUPPORT

### Internal sources

- Boehringer Ingelheim, Netherlands.

### External sources

- Netherlands Asthma Foundation, Netherlands.
- Glaxo Wellcome BV, Netherlands.
- Boehringer Ingelheim, Netherlands.
- Amicon Health Care Insurance Company, Netherlands.
- Euregio, Belgium.
- Garfield Weston Foundation, UK.

## INDEX TERMS

### Medical Subject Headings (MeSH)

\*Patient Education as Topic; \*Self Care; Health Status; Hospitalization [statistics & numerical data]; Outcome Assessment (Health Care); Patient Compliance; Program Evaluation; Pulmonary Disease, Chronic Obstructive [\*therapy]; Quality of Life; Randomized Controlled Trials as Topic

### MeSH check words

Humans