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Respiratory interventions for breathlessness in adults with advanced diseases (Protocol)

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TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
BACKGROUND	1
Figure 1.	2
Figure 2.	3
OBJECTIVES	4
METHODS	4
ACKNOWLEDGEMENTS	9
REFERENCES	9
APPENDICES	11
CONTRIBUTIONS OF AUTHORS	12
DECLARATIONS OF INTEREST	12
SOURCES OF SUPPORT	13

[Intervention Protocol]

Respiratory interventions for breathlessness in adults with advanced diseases

Anna Bolzani¹, Stefanie M Rolser¹, Helen Kalies¹, Matthew Maddocks², Eva Rehfuss³, Flavia Swan⁴, Marjolein Gysels⁵, Irene J Higginson⁶, Sara Booth⁷, Claudia Bausewein¹

¹Department of Palliative Medicine, Munich University Hospital, LMU Munich, Munich, Germany. ²Department of Palliative Care, Policy and Rehabilitation, Cicely Saunders Institute, King's College London, London, UK. ³Institute for Medical Informatics, Biometry and Epidemiology, Pettenkofer School of Public Health, LMU Munich, Munich, Germany. ⁴Hull Medical School, University of Hull, Hull, UK. ⁵Amsterdam Institute of Social Science Research, University of Amsterdam, Amsterdam, Netherlands. ⁶Department of Palliative Care, Policy and Rehabilitation, Cicely Saunders Institute, King's College London, London, UK. ⁷Department of Palliative Care, Cambridge University Hospitals, Cambridge, UK

Contact address: Anna Bolzani, Department of Palliative Medicine, Munich University Hospital, LMU Munich, Marchioninstr. 15, Munich, Germany. Anna.Bolzani@med.uni-muenchen.de.

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ABSTRACT

This is a protocol for a Cochrane Review (Intervention). The objectives are as follows:

To assess the effects and safety of interventions targeting respiration as the predominant underlying mechanism of effect to relieve breathlessness in adults suffering from advanced diseases.

BACKGROUND

This protocol is partly based on suggested wording from the Cochrane Pain, Palliative and Supportive Care Review Group (PaPaS CRG). Some wording is used from the original review (Bausewein 2008), which this new review will update and replace.

Description of the condition

Breathlessness or dyspnoea is defined as “subjective experience of breathing discomfort that consists of qualitatively distinct sensations that vary in intensity” (Meek 1999). The term ‘breathlessness’ reflects the patients’ perspective based on the daily experi-

ence whereas the medical term ‘dyspnoea’ focuses more on the clinical sign of an underlying condition (Johnson 2014). “The experience derives from interactions among multiple physiological, psychological, social, and environmental factors, and may include secondary physiological and behavioural responses” (Meek 1999). Since this definition was adopted, new evidence has led to better understanding of the mainly sensory and affective components and that dyspnoea “must generally be distinguished from signs that clinicians typically invoke as evidence of respiratory distress, such as tachypn(o)ea, use of accessory muscles, and intercostal retractions.” (Parshall 2012). Many patients with different conditions including primary and secondary cancer, lung diseases (e.g.

chronic obstructive pulmonary disease (COPD), pulmonary hypertension, cystic fibrosis, interstitial lung disease (ILD)), chronic heart failure (CHF) or motor neuron disease/amyotrophic lateral sclerosis (MND/ALS) suffer from this distressing symptom (Bailey 2010; Booth 2008; Breaden 2011; Lansing 2009; Solano 2006). Breathlessness is a multifactorial and complex symptom and an experience unique to the individual (Booth 2008). It is often expressed as air hunger, work of breathing, laboured breathing, awareness of respiratory distress, and shortness of breath or chest tightness (Barnes 2016; Parshall 2012). Breathing discomfort is described by such phrases as ‘could not breathe fast or deep enough’ or ‘could not get enough air’ or ‘suffocating’ (Guz 1997). Breathlessness is one of the most prevalent and distressing symptoms in advanced stages of malignant and non-malignant diseases. Up to 95% of patients with advanced chronic pulmonary disease, 88% with advanced heart disease, and 70% with end stage cancer experience breathlessness in their last year of life (Graham 2010; Lansing 2009; Moens 2014; Solano 2006; Teunissen 2007). The frequency and severity of breathlessness increase during the course of the disease until death (Bailey 2010; Breaden 2011). It is an extremely distressing symptom for the patient but also for the accompanying family and professional carers (Booth 2008). Overall, breathlessness is still difficult to palliate.

Description of the intervention

Management of breathlessness

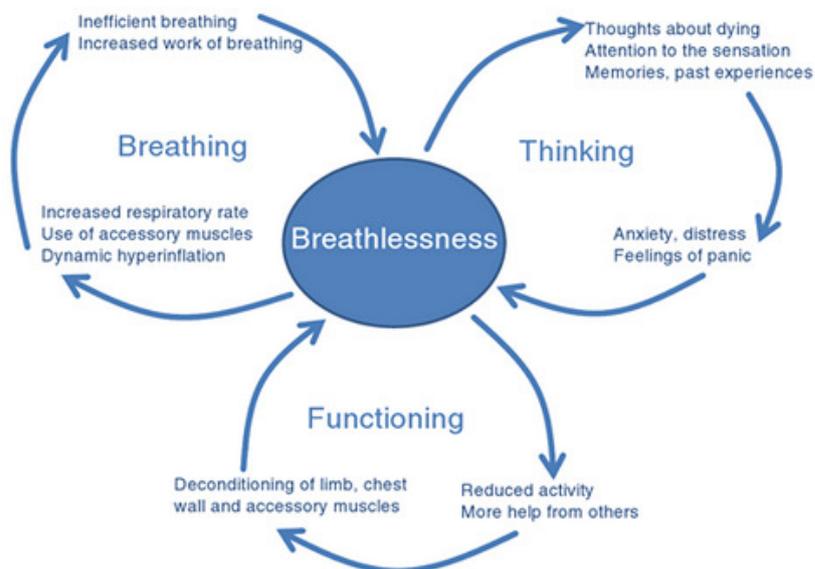
Appropriate management to relieve breathlessness in advanced diseases requires both pharmacological and non-pharmacological interventions. Different systematic reviews and meta-analyses were published in recent years and analysed the effects of pharmacological interventions such as opioids (Barnes 2016; Mahler 2013), benzodiazepines (Simon 2016), and oxygen (Ameer 2014; Cranston 2008; Sharp 2016) for breathlessness in adult patients.

However, the use of drugs to treat breathlessness is sometimes limited as they entail adverse effects and doses need to be titrated carefully. Therefore, non-pharmacological interventions are an important part of the treatment of breathlessness. As mentioned above, many systematic reviews analysed the effects of pharmacological treatments, which is why we are focusing solely on non-pharmacological interventions in this review.

Non-pharmacological interventions

Many non-pharmacological interventions for the relief of breathlessness have been developed and evaluated in recent years. For better clarity, we therefore categorise the interventions based on a theoretical concept developed by Booth 2014, Chin 2016 and Spathis 2017. This concept builds on the effect breathlessness has on patients (Figure 1).

Figure 1. Perpetuation of breathlessness by vicious cycles (Booth 2014)

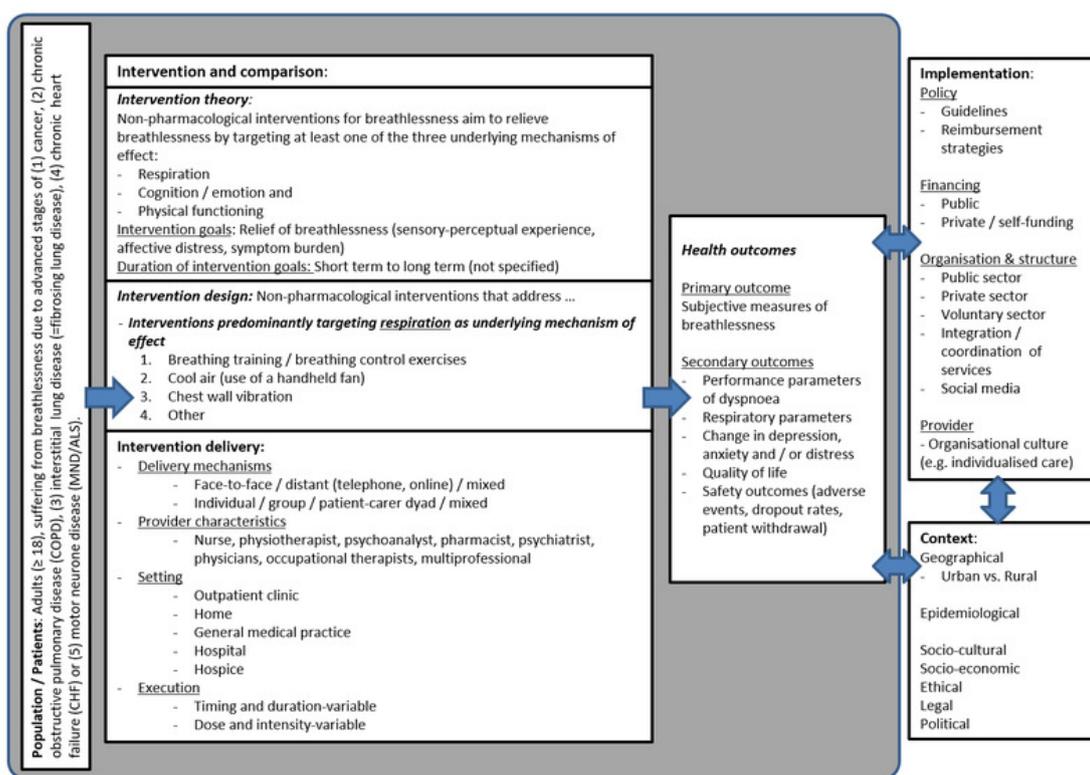


- Respiratory: Inefficient breathing and increased work of breathing can be observed due to dysfunctional breathing patterns with an increased respiratory rate, the need for the use of accessory muscles, and dynamic hyperinflation.
- Cognitive-emotional: Misconceptions and paying too much attention to the sensation of breathlessness such as memories of past or negative experiences lead to anxiety, distress, feelings of panic, and thoughts about dying.
- Physical: Persons suffering from severe breathlessness show reduced physical activity with a tendency to self-isolation and the need for more help from others. This leads to deconditioning of limb, chest wall and accessory muscles.

We expect a huge number of studies and categories of interventions to be included. Therefore, three different reviews, based on the theoretical concept, will be conducted. An additional review is planned, focusing on interventions targeting more than one underlying mechanism as described above.

In this review, we will analyse non-pharmacological interventions targeting primarily respiration to relieve breathlessness in patients suffering from advanced stages of disease, for example breathing training, handheld fan, and chest wall vibration. These interventions may take place in a variety of settings, and can, with guidance of healthcare professionals, mostly be carried out by patients themselves (Figure 2).

Figure 2. System-based logic model on respiration interventions for breathlessness in patients with advanced diseases



Invasive interventions could also be classified as non-pharmacological but they will not be the focus in this review. Therefore, we will exclude surgical procedures such as drainage, tapping, endoscopy, ventilation and catheterisation. We will also exclude the following non-pharmacological interven-

tions as there have been recent Cochrane reviews: pulmonary rehabilitation (McCarthy 2015), and nutrition (Ferreira 2012).

How the intervention might work

As we expect several different interventions to be subsumed in this review, there is no one underlying mechanism but various mechanisms that target respiration by enhancing breathing.

Breathing training interventions address the main physiologic impediment of respiration and focus on gentle and prolonged exhalation and on slower and deeper breathing (Nield 2007; Spahija 1996). Examples are diaphragmatic breathing, pursed lip breathing, body position exercises, and respiratory muscle training.

The mechanism by which a fan or a breeze of cool air reduces breathlessness remains unclear, but is possibly linked to the diving response, which causes ventilatory depression when the trigeminal area of the face is cooled (stimulation of facial and nasopharyngeal receptors) (Galbraith 2010).

In-phase chest wall vibration stimulates chest wall receptors that alter respiratory sensations and reduces the breathing discomfort at rest associated with steady-state hypercapnia. Although there are conflicting findings about which receptors mediate effects of vibration on ventilation, there is evidence that muscle spindles in intercostal muscles may be responsible (Cristiano 1997; Homma 1984; Parshall 2012).

Based on a template by Rohwer 2017 we developed a system-based logic model in which we show how non-pharmacological interventions for breathlessness, with a focus on interventions predominantly targeting respiration, are implemented in the healthcare system (Figure 2).

Why it is important to do this review

Non-pharmacological interventions can complement pharmacological interventions and may offer alternative treatment options in the management of breathlessness occurring in advanced illness. As research into this challenging, poorly managed and burdensome symptom is rapidly evolving, there is a need to synthesise the most recent evidence to inform practice and research. Our review aim is to aid health professionals in the treatment of breathlessness with palliative intent and to inform patients and carers about the evidence of non-pharmacological interventions targeting respiration to relieve breathlessness.

This is an update of a Cochrane review on non-pharmacological interventions for the relief of breathlessness in advanced disease (Bausewein 2008). The former review showed effectiveness of neuromuscular electrical stimulation, chest wall vibration, walking aids, and breathing training. The review included 47 studies that were categorised in different intervention groups (e.g. walking aids, acupuncture, breathing training, psychological therapy). Since its publication, many randomised controlled studies on non-pharmacological interventions have been published, including new intervention groups (e.g. breathlessness services). Therefore, although necessary, a single review as an update of the earlier review seemed infeasible. Based on the interventions used to target breathlessness, we decided to assess the interventions in different reviews.

OBJECTIVES

To assess the effects and safety of interventions targeting respiration as the predominant underlying mechanism of effect to relieve breathlessness in adults suffering from advanced diseases.

METHODS

Criteria for considering studies for this review

Types of studies

We will include randomised controlled trials (RCTs), cluster RCTs, and quasi-RCTs (QRCTs). Quasi-randomisation is defined as some pseudo-random method of allocation such as alternation, date of birth, case record number or date of presentation (Higgins 2011). We will include cross-over studies, if separate data for both time periods are presented. We will only use the data of the first period for analysis to avoid carry-over effects. We will require full journal publication. Where full journal publication is not available, we will try to obtain data by contacting the trial authors, unless sufficient data for analyses are provided in online clinical trial results, summaries of otherwise unpublished clinical trials, or conference abstracts. QRCTs will be included in order to obtain the full breadth of relevant trials, in particular as we expect to find a small number of RCTs for some of the intervention categories; we are aware of the higher risk of bias in these studies and will account for this in the analysis.

Types of participants

Adult patients aged 18 years and above, suffering from advanced diseases with a high prevalence of breathlessness.

We will include studies if the majority ($\geq 50\%$) of participants meet the following criteria.

- Patients suffering from cancer should have advanced local or metastatic disease (e.g. TNM Classification of Malignant Tumours (TNM) state $\geq T3$ or $N \geq 1$ or $M \geq 1$).
- Patients with severe COPD should have a forced expiratory volume in one second (FEV1) predicted of $< 50\%$.
- Patients with pulmonary hypertension will be included if they reach a WHO class level $\geq III$, defined by Barst 2004.
- Patients suffering from CHF should have New York Heart Association (NYHA) stage III or IV.
- Patients with ILD or idiopathic pulmonary fibrosis (IPF) : all studies will be included as breathlessness is the predominant symptom and there are hardly any disease-specific treatment options.
- Patients with neuromuscular diseases (MND, ALS): all studies will be included as advanced disease is marked by the occurrence of breathlessness.

If groups for the inclusion criteria mentioned above were stratified, we will only include the subgroups of interest. We will document difficult decisions in the review. Sensitivity analysis can assess the impact of these decisions on the review's result. Patients included in the studies can be in any setting. We will exclude studies of patients with any condition not regarded as advanced and life-limiting such as acute or chronic asthma, or with pre-existing diagnosis of acute asthma or acute cardiac condition as a primary cause of breathlessness.

Types of interventions

We will include interventions targeting respiration to relieve breathlessness according to the following prespecified categories.

- Breathing training or breathing control exercises (e.g. diaphragmatic breathing, pursed lip breathing, body position exercises, respiratory muscle training).
- Cool air (e.g. use of a handheld fan).
- Chest wall vibration.

If we find interventions of interest that do not fit in the above categories, we will define an additional category 'Other' or add new categories if there is a sufficient number of studies.

The judgement for inclusion will be based on the study authors' description of the intervention; any deviation from this will be explicitly mentioned.

Interventions may take place in any setting, e.g. outpatient clinic, home, hospital, hospice, general medical practice.

The comparator may be no treatment, placebo, attention control, standard care, or a different kind of therapy. We will categorise the control groups into 'active controls' or 'other' based on the description of the comparison group. We will focus on active controls as comparison group in our primary analysis. Concomitant interventions, especially pharmacological treatment, will be accepted, if administered in the same way in both the control and the treatment groups. If these interventions are suspected to have some relevant influence on our outcomes we will consider this in subgroup analysis.

Types of outcome measures

We anticipate that studies will use a variety of outcome measures. To be included, a study must have any measure of breathlessness. Adverse effects of respiration interventions will be measured as absent or present and a narrative description of these effects will be given when reported. We will consider all reliable and validated measures for the following outcomes.

Primary outcomes

Breathlessness, measured by self-reported instruments with a focus on breathlessness or mastery of breathlessness (e.g. Baseline Dyspnoea Index (BDI), Borg Dyspnoea Scale (BDS), Medical Research

Council (MRC) Breathlessness Scale, or Chronic Respiratory Disease Questionnaire (CRQ)). Other terms for breathlessness such as dyspnoea, shortness of breath and difficulty breathing will also be accepted.

Secondary outcomes

- Performance parameters (e.g. walking tests, International Physical Activity Questionnaire (IPAQ)).
- Respiratory parameters (e.g. change in FEV1 (%)).
- Change in depression, anxiety and/or distress (e.g. Hospital Anxiety and Depression Scale (HADS)).
- Quality of life (e.g. 36-Item Short Form Health Survey (SF-36)).
- Safety outcomes:
 - Adverse events (measured as absent or present);
 - Dropout rates; and
 - Patient withdrawal from the trial, due to any reason (if mentioned).

Search methods for identification of studies

Electronic searches

We will search the following databases from their inception to the present, without date or language restrictions.

- Cochrane Database of Systematic Reviews (CDSR), the Cochrane Library.
- Cochrane Central Register of Controlled Trials (CENTRAL), the Cochrane Library.
- MEDLINE (Ovid).
- Embase (Ovid).
- PsycINFO (Ovid).
- LILACS (Bireme).
- CINAHL (Ebsco).

We will search MEDLINE and Embase using both controlled vocabulary (namely, MeSH in MEDLINE and Emtree in Embase) and a wide range of free-text terms. To detect all RCTs we will perform the search on MEDLINE using the Cochrane Highly Sensitive Search Strategy, sensitivity-maximising version ([Higgins 2011](#)).

The search strategy for MEDLINE is in [Appendix 1](#).

Searching other resources

We will search the meta-register of controlled trials (mRCT) (www.controlled-trials.com/mrct), clinicaltrials.gov (www.clinicaltrials.gov) and the WHO International Clinical Trials Registry Platform (ICTRP) (apps.who.int/trialsearch/) for ongoing trials. In addition, we will check reference lists of reviews

and retrieved articles for additional studies, and we will perform citation searches on key articles. We will contact experts in the field for unpublished and ongoing trials. We will contact study authors where necessary for additional information.

We will perform the search in collaboration with the Information Specialist of the Cochrane Pain, Palliative and Supportive Care Group.

Data collection and analysis

Selection of studies

Two review authors (AB, SR) will independently screen all titles and abstracts retrieved by the search to identify all trials that may be eligible and for which the full paper should be obtained. Independent review authors will eliminate studies that clearly do not satisfy inclusion criteria, and obtain full copies of the remaining studies. Two review authors (AB, SR) will read these studies independently to select relevant studies, and in the event of disagreement or unclear decision to include, we will resolve disagreement with a third author (MM or CB, depending on the topic). We will not anonymise the studies in any way before assessment.

We will include a PRISMA flow chart in the full review which will show the status of identified studies (Moher 2009) as recommended in Part 2, section 11.2.1 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We will include studies in the review irrespective of whether measured outcome data are reported in a 'usable' way.

Data extraction and management

Two review authors (SR, AB, FS or MM) will independently extract data using a data collection form based on a standard form released by the Cochrane Effective Practice and Organisation of Care Group (EPOC) and check for agreement before entry into Review Manager (RevMan 2014). Where there is disagreement, a third author (CB or SB) will be consulted to resolve differences. We will include information about the following.

Participant characteristics

- Demographic characteristics (age, gender, nationality).
- Underlying disease characteristics (type and stage of condition).

Intervention

- Intervention theory.
- Type of intervention (description of intervention, frequency, duration (total and per session)).

- Types of control condition (control intervention, control group).
- Type of delivery (delivery mechanisms such as face-to-face, distant; group, individual; provider characteristics such as nurses, physicians, multiprofessional; setting such as outpatient clinic, home, hospital).

Methods

- Study design.
- Size of intervention and control group at baseline and follow-up.
- Study duration and follow-up.
- Sources of bias (sequence generation, allocation sequence concealment, blinding, incomplete outcome data, selective reporting, other concerns about bias).

Outcomes

- Key outcomes with measurement instruments.
- Timing, duration and frequency of follow-up.
- Adverse events.
- Number of withdrawals and dropouts.

Context

- Country of origin.

In case multiple reports of the same study are found, we will extract data of all these reports independently of each other and compare; if data differ between reports, all authors will make a decision how to treat this study and this will be documented in the review. We will collate multiple reports of the same study, so that each study rather than each report is the unit of interest in the review. We will collect characteristics of the included studies in sufficient detail to populate a table of 'Characteristics of included studies' in the full review. Review authors will not be involved in the data extraction of studies they authored or co-authored.

Assessment of risk of bias in included studies

Two authors (AB, MM) will independently assess risk of bias for each study, using the criteria outlined in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011) and adapted from those used by the Cochrane Pregnancy and Childbirth Group.

We will assess the following for each study.

- Random sequence generation (checking for possible selection bias). We will assess the method used to generate the allocation sequence as: low risk of bias (any truly random process, e.g. random number table; computer random number generator); unclear risk of bias (method used to generate sequence not clearly stated). Studies using a non-random process

(e.g. odd or even date of birth; hospital or clinic record number) will be assessed as high risk of bias.

- Allocation concealment (checking for possible selection bias). The method used to conceal allocation to interventions prior to assignment determines whether intervention allocation could have been foreseen in advance of, or during, recruitment, or changed after assignment. We will assess the methods as: low risk of bias (e.g. telephone or central randomisation; consecutively numbered sealed opaque envelopes); high risk of bias (studies that do not conceal allocation (e.g. open list); unclear risk of bias (method not clearly stated).

- Blinding of outcome assessment (checking for possible detection bias). We will assess the methods used to blind study participants and outcome assessors from knowledge of which intervention a participant received. We will assess the methods as: low risk of bias (study states that it was blinded and describes the method used to achieve blinding); high risk of bias (no or incomplete blinding); unclear risk of bias (study states that it was blinded but does not provide an adequate description of how it was achieved). We will also report if study participants are asked about their expectations of benefit of intervention/control if blinding is not feasible.

- Incomplete outcome data (checking for possible attrition bias due to the amount, nature and handling of incomplete outcome data). We will assess the methods used to deal with incomplete data as: low risk (< 10% of participants did not complete the study and/or data have been imputed using appropriate methods); high risk of bias (used 'completer' analysis); unclear risk of bias (insufficient information for low/high risk of bias category).

- Selective reporting (checking for reporting bias). We will assess the methods as: low risk of bias (where it is clear that all of the study's prespecified outcomes and all expected outcomes of interest to the review have been reported); high risk of bias (where not all the study's prespecified outcomes have been reported; one or more reported primary outcomes were not prespecified; outcomes of interest are reported incompletely and so cannot be used; study fails to include results of a key outcome that would have been expected to have been reported); unclear risk of bias (insufficient information for low/high risk of bias category).

- Other bias (e.g. checking for possible biases confounded by small size. We will assess studies as being at low risk of bias (\geq 200 participants per treatment arm); unclear risk of bias (50 to 199 participants per treatment arm); high risk of bias (< 50 participants per treatment arm)).

We will use the Review Manager tool to complete a 'Risk of bias' table (RevMan 2014). Any discrepancy between the two authors will be resolved by discussion involving a third author (CB).

Measures of treatment effect

We will analyse dichotomous outcomes using risk ratios (RRs) with 95% confidence intervals (CIs). We will recategorise any categorical outcomes with more than two categories into two groups. We will analyse continuous data using standardised mean differences (SMDs) with 95% CIs. We will calculate standard deviations, if not reported, using the methods described in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We plan to report the proportion of participants experiencing any adverse effects of respiration interventions, and combine studies using RRs with 95% CIs.

Unit of analysis issues

We will reanalyse data, if possible, for cluster trials which have not taken clustering into account in their analysis. We will calculate effective sample sizes and adjusted standard errors using the design effect method. We will try to obtain estimates for intracluster correlation coefficients from study authors or will use external estimates obtained from comparable studies, as recommended by Cochrane guidelines (Higgins 2011). We will document if reanalysis is not feasible.

In studies with more than two arms, we will consistently choose the active control arms in the main analysis, and, if possible, do a sensitivity analysis, in which we will choose the other control arm. We will combine individually randomised controlled trials and cluster RCTs in the same meta-analyses or harvest plots, but these will be clearly identified (Higgins 2011).

Dealing with missing data

We will contact study authors if missing data on study characteristics or outcome measures precludes study inclusion or limits use of a study at further stages of the review. If studies do not report outcomes based on intention-to-treat analyses this will be considered as a source of bias during 'Risk of bias' assessment. We will try to calculate effect measures or CIs wherever possible from available data, if we get no response.

Assessment of heterogeneity

We will assess methodological and clinical heterogeneity with tables documenting the following characteristics of the included studies.

- Intervention components (e.g. breathing training; cool air; chest wall vibration).
- Intervention delivery mechanism (e.g. face-to-face, distant).
- Provider characteristics (e.g. nurses, physiotherapists, physicians).
- Setting (e.g. outpatient clinic, hospice, home).
- Patients (e.g. COPD, cancer, fibrosing lung disease).
- Methods (outcome measures, outcome assessment).

For those studies assessing the impacts of a given intervention category on comparable outcomes, thus making pooling through

meta-analysis feasible, we will assess statistical heterogeneity graphically with a forest plot by examining the extent to which CIs overlap, and statistically with the I^2 statistic. We will consider an I^2 value greater than 50% to indicate substantial statistical heterogeneity, and will consider it statistically significant if the P value for the Chi^2 test is < 0.1 . We will document statistical heterogeneity but this will not have any direct consequences for meta-analysis (see below). We will create forest plots and I^2 calculations using Review Manager 5.3 (RevMan 2014).

Assessment of reporting biases

We will try to minimise publication bias by searching trials registers for projected and registered studies that have never been published. We will contact the authors to get unpublished information if there are such studies registered or some relevant information is missing and can therefore narrow the risk of reporting bias. We will assess the possibility that publication bias affects the review using funnel plots when at least 10 studies are available for meta-analysis.

Data synthesis

We will attempt to pool all studies within a given intervention category assessing the same outcome by conducting a meta-analysis using Review Manager 5.3 (RevMan 2014). We will use the random-effects model due to the expected large heterogeneity in delivery mechanisms, provider characteristics, setting and study population.

We will report results as RRs for dichotomous outcomes and SMDs for continuous outcomes. We will undertake meta-analysis only if studies are judged to be similar enough to give a clinically meaningful answer. We will provide an outcome table and summarise the results narratively if meta-analysis is not possible.

In the case of skewed data, we will log transform these data for our analysis or, if that approach is not feasible, summarise them narratively.

'Summary of findings' table

We will include a 'Summary of findings' table using the GRADE profiler software (GRADEpro GDT 2015) as set out in the PaPaS author guide (AUREF 2012) and recommended in the *Cochrane Handbook for Systematic Reviews of Interventions*, Chapter 4.6.6 (Higgins 2011) to evaluate the quality of evidence in our review. The 'Summary of findings' table will include outcomes: of a) change of breathlessness, b) objective parameters of breathlessness c) quality of life indicators, d) change of depression or anxiety, e) adverse events, f) characteristics of the patient population that benefits most.

Quality of the evidence

This section is taken from the Cochrane Drugs and Alcohol Group recommended text. The overall quality of the evidence for each outcome in our review will be assessed using the GRADE system (GRADEpro GDT 2015) and presented in the 'Summary of findings' tables, to present the main findings of a review in a transparent and simple tabular format. In particular, we will include key information concerning the quality of evidence, the magnitude of effect of the interventions examined, and the sum of available data on the main outcomes.

The GRADE system uses the following criteria for assigning grade of evidence.

- High: we are very confident that the true effect lies close to that of the estimate of the effect.
- Moderate: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of effect, but there is a possibility that it is substantially different.
- Low: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect.
- Very low: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect.

We will decrease grade rating by one (-1) or two (-2) if we identify:

- serious (-1) or very serious (-2) limitation to study quality;
- important inconsistency (-1);
- some (-1) or major (-2) uncertainty about directness;
- imprecise or sparse data (-1); or
- high probability of reporting bias (-1).

Subgroup analysis and investigation of heterogeneity

We will undertake subgroup analysis for the primary outcomes to examine factors that may explain variation in the effectiveness, if numbers are sufficiently large. We will perform stratification as follows.

- Type of intervention.
- Intervention delivery (delivery mechanisms such as face-to-face, distant; group, individual; provider characteristics such as nurses, physicians, multiprofessional; setting such as outpatient clinic, home, hospital).
- Patient characteristics (underlying disease, disease stage, age, gender).
- Underlying therapy.

Sensitivity analysis

We will conduct sensitivity analysis where possible, to test the effect of different methodological decisions made throughout the review process on the primary outcome. We will test the robustness of the results by removing from the pooled effect estimate:

- studies with a high risk of bias for two or more key domains;

- quasi-randomised clinical trials;
- outcome measures;
- intervention of varying duration.

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* Indicates the major publication for the study

APPENDICES

Appendix I. Search strategy for MEDLINE (Ovid)

1. exp Dyspnea/
2. dyspn?ea.tw.
3. (short* adj2 breath).tw.
4. (urge* adj2 breath*).tw.
5. breathless*.tw.
6. ((labo?red or difficult* or small) adj3 breath*).tw.
7. ((respirat* or breath*) adj3 (distress* or comfort* or discomfort*)).tw.
8. (air adj3 (hunger or starve* or need* or gasp* or pant*)).tw.
9. suffocat*.tw.
10. unsatisf* inspiration.tw.
11. or/1-10
12. Neoplasms/ or Lung Neoplasms/
13. ((lung* or bronchi* or pulmo*) adj3 (neoplasm* or cancer* or tumo?* or metasta* or malignan*)).mp.
14. Lung diseases/
15. exp Pulmonary Disease, Chronic Obstructive/
16. (COPD or COAD).tw.
17. Lung Diseases, Obstructive/
18. (obstruct* adj3 (pulmonary or lung* or airway* or airflow* or bronch* or respirat*)).tw.
19. hypertension, pulmonary/
20. or/12-19
21. exp Heart Failure/
22. ((heart or cardia* or myocard*) adj2 (fail* or insufficienc*)).tw.
23. (decompensat* adj2 (heart* or cardia*)).tw.
24. decompensatio cordis.tw.
25. insufficientia cordis.tw.
26. ((cardiac or heart) adj2 incompetenc*).tw.
27. cardiac stand still.tw.
28. or/21-27
29. exp Lung Diseases, Interstitial/
30. (interstitial adj3 (disease* or pneumoni* or fibrosis)).tw.
31. pulmonary fibrosis.tw.
32. fibrosing alveolitis.tw.
33. Cystic Fibrosis/
34. (cystic fibrosis or mucoviscidosis).tw.

35. or/29-34
36. exp Motor Neuron Disease/
37. (MND or ALS).tw.
38. motor neuron disease*.tw.
39. sclerosis.tw.
40. Amyotrophic Lateral Sclerosis/
41. charcot disease*.tw.
42. lou gehrig disease*.tw.
43. encephalomyelitis disseminate.mp.
44. or/36-43
45. 20 or 28 or 35 or 44
46. ((end stage or advanc* or final or terminal* or limit*) adj3 (disease* or illness*)).tw.
47. Terminally Ill/
48. Terminal Care/
49. Palliative Care/
50. Prognosis/
51. ((advanc* or terminal or limit*) adj3 (prognos* or prospect* or prediction*)).tw.
52. disease progression/
53. ((incurable or worsen* or chronic) adj3 (illness* or disease*)).tw.
54. or/46-53
55. 45 or 54
56. randomized controlled trial.pt.
57. controlled clinical trial.pt.
58. randomized.ab.
59. placebo.ab.
60. drug therapy.fs.
61. randomly.ab.
62. trial.ab.
63. groups.ab.
64. 56 or 57 or 58 or 59 or 60 or 61 or 62 or 63
65. exp animals/ not humans.sh.
66. 64 not 65
67. 11 and 55
68. 66 and 67

CONTRIBUTIONS OF AUTHORS

Developed concept of review: HK, CB, MM, ER, SB, IJH, MG.

Drafted the protocol: HK, AB, SR.

Checked and approved the draft: CB, SB, MM, MG, ER.

Developed search strategy: SR.

DECLARATIONS OF INTEREST

AB: none known.

SR: none known.

HK: none known.

MM: none known. MM is investigator on studies that might be included in this review; MM is a specialist physiotherapist and manages patients with breathlessness and advanced disease.

ER: none known.

FS: none known.

MG: none known.

IJH: none known. IJH is investigator on studies that might be included in this review; IJH is a specialist physician in palliative medicine and manages patients with breathlessness and advanced disease.

SB: has received payment for talks in Feb 2016 from Novartis. SB is investigator on studies that might be included in this review; SB is a specialist physician in palliative medicine and manages patients with breathlessness and advanced disease.

CB: has received payment for one talk in Oct 2015 from Bayer Health Care. CB is investigator on studies that might be included in this review; CB is a specialist physician in palliative medicine and manages patients with breathlessness and advanced disease.

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