TEMPLEMAN, L. and ROBERTS, F. 2020. Effectiveness of expiratory muscle strength training on expiratory strength, pulmonary function and cough in the adult population: a systematic review. *Physiotherapy* [online], 106, pages 43-51. Available from: <u>https://doi.org/10.1016/j.physio.2019.06.002</u>

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TEMPLEMAN, L., ROBERTS, F.

2020



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Accepted Manuscript

Title: Effectiveness Of Expiratory Muscle Strength Training On Expiratory Strength, Pulmonary Function And Cough In The Adult Population: A Systematic Review

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 PII:
 S0031-9406(19)30076-8

 DOI:
 https://doi.org/10.1016/j.physio.2019.06.002

 Reference:
 PHYST 1111

To appear in: *Physiotherapy*



Please cite this article as: Templeman L, Roberts F, Effectiveness Of Expiratory Muscle Strength Training On Expiratory Strength, Pulmonary Function And Cough In The Adult Population: A Systematic Review, *Physiotherapy* (2019), https://doi.org/10.1016/j.physio.2019.06.002

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Effectiveness Of Expiratory Muscle Strength Training On Expiratory Strength, Pulmonary Function And Cough In The Adult Population: A Systematic Review

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Word Count: 3132 (excluding abstract, references, tables, figures, legends)

Abstract

Background: Respiratory muscle strength declines in certain disease states, leading to impaired cough, reduced airway clearance and an increased risk of aspiration pneumonia. Respiratory muscle training may therefore reduce this risk.

Objectives: To assess current evidence of expiratory muscle strength training (EMST) on maximum expiratory pressure, cough flow and spirometry.

Data sources: Databases including CINAHL, Medline, Science Direct and PEDRo were searched.

Eligibility Criteria: Randomised controlled trials investigating expiratory muscle strength training on maximum expiratory pressure, pulmonary function or cough in any adult population, published before December 2017.

Study appraisal: Data were extracted to a trial description form and study quality evaluated by 2 reviewers. Meta-analysis was performed with calculation of mean differences and 95% confidence intervals.

Results: Nine studies met inclusion criteria and ranged in size from 12 to 42 participants. Trials investigated EMST in healthy adults (2), multiple sclerosis (3), COPD (2), acute stroke (1) and spinal cord injury (1). Overall, EMST improved maximum expiratory pressure (15.95cmH2O; 95% CI: 7.77 to 24.12; p<0.01) with no significant impact on cough flow (4.63L/min; 95%CI -27.48 to 36.74; P=0.78), forced vital capacity (-0.16L; 95%CI -0.35 to 0.02; P=0.09) or forced expiratory volume in 1 second (-0.09L; 95%CI -0.10 to -0.08; P<0.001) versus control or sham training.

Conclusions: Meta-analysis indicated a small significant increase in maximum expiratory pressure following EMST. Improvements in maximum expiratory pressure did not lead to improvements in cough or pulmonary function.

Limitations: Variations in protocol design and population limited the overall effect size.

Funding: None

Systematic Review Registration: PROSPERO (CRD42018104190).

Contribution of Paper

This review synthesises current EMST data on expiratory strength, pulmonary function and cough.

Despite differences in intervention protocols, meta-analysis indicated a small but statistically significant increase in maximum expiratory pressure following training. Few studies included secondary outcome measures of pulmonary function and cough flow with no evidence to support any change in airflow measures following EMST versus control or sham intervention.

The functional relevance of maximum expiratory pressure gains is yet to be determined. Physiotherapists and clinicians involved in pulmonary rehabilitation or with patients prone to aspiration should be advised that use of EMST *alone* for improving airway clearance mechanics is not supported. Further evidence of the relationship between expiratory strength and airway clearance is needed before EMST alone could be justified clinically to prevent aspiration.

Keywords: Expiratory muscle strength training, maximum expiratory pressure, aspiration pneumonia.

INTRODUCTION

Respiratory muscle function is vital for life: creating pressure differences needed for ventilation, eliminating airway secretions and protecting the airways [1].

Respiratory muscle strength and pressure is known to decline progressively with increasing age [2], possibly leading to inadequate ventilation and undermining normal airway protection [1]. Certainly the increased frequency of aspiration pneumonia in the elderly has been attributed to impaired airway protection such as reduced peak expiratory flow and cough [3]. Impaired respiratory strength is also observed in several disease states including chronic obstructive pulmonary disease (COPD) [4], Parkinson's disease (PD) [5,6], multiple sclerosis (MS) [7,8] and stroke [9], with reduced airway protection increasing the risk of aspiration pneumonia and hospital admission [10-12]. For this reason, respiratory muscle training becomes interesting as a strategy to reduce mortality from pneumonia.

The respiratory musculature comprises inspiratory and expiratory muscles with focus, to date, on inspiratory muscle training (IMT). Studies in healthy populations have demonstrated positive changes in respiratory muscle strength and diaphragm mobility following IMT [13,14] with similar findings in patients with stroke, MS and PD [15,16]. Evidence also suggests respiratory weakness varies between diseases with expiratory muscles shown to weaken to a lesser extent than inspiratory in patients with COPD [17], but to a greater extent in MS [18]. From a functional perspective, expiratory muscles are known to have high activation for force generation during cough [19], therefore, specific expiratory weakness and impaired airway clearance may be responsible for the increased incidence of aspiration in patients prone to respiratory decline. Certainly studies in stroke suggest impairment of cough function is due to weakness of expiratory muscles rather than dysfunction at the level of the glottis [9]. Interventions targeting expiratory muscle strength therefore provide an attractive prophylaxis against aspiration pneumonia.

Studies of Expiratory Muscle Strength Training (EMST) on maximum expiratory pressure (MEP) have been performed in different populations, but there is no consensus on the effects of EMST on MEP, airway clearance or other pulmonary measures with no systematic reviews or meta-analyses and just one narrative review published to date [20].

Consequently the aim of this systematic review was to investigate evidence for EMST on MEP in health and disease states. A secondary objective was to identify whether EMST alters spirometry or cough measures.

Review question

Does expiratory muscle strength training affect maximum expiratory pressure, cough flow or spirometry in adult populations?

Methods

Systematic Review and Meta Analysis

This quantitative systematic review protocol was prospectively registered with PROSPERO (CRD42018104190).

Search Strategy

A three-step search strategy was used. An initial, limited search of Medline and CINAHL was followed by analysis of text words and index terms. Using keywords

identified, a second search was undertaken using CINAHL, Medline, Science Direct, PEDRo and Cochrane Central Register for Controlled Trials. All databases were searched from inception to end December 2017. Search terms were: Expiratory muscle strength training; expiratory training; OR Respiratory muscle strength training; AND expiratory pressure, maximum expiratory pressure, cough, sputum clearance. In a third step, references of retrieved studies were searched to identify further publications. To minimise publication bias, 'grey' literature was searched using Google Scholar. Due to limited resources for translation, only articles published in English were included.

Study Design

Randomised controlled trials (RCTs) only.

Participants

Studies of healthy adults and those with pulmonary, neurological or neuromuscular conditions were included. No upper age limit was defined.

Interventions and Comparators

Reports of threshold or resistance EMST were included. Combinations of interventions were excluded (e.g. inspiratory and expiratory training). Comparators were control intervention, sham EMST or breathing exercises.

Outcome Measures

The primary outcome measure was MEP. Secondary outcome measures included forced vital capacity (FVC), forced expiratory volume in 1 second (FEV₁), and peak expiratory cough flow (PECF).

Study selection

On search completion, references were exported to Refworks, with duplicates identified and removed. Titles and abstracts were independently screened against inclusion criteria by two reviewers (LT and FR) and inclusion validated by discussion and consensus.

Quality Assessment

Methodological quality of each eligible article was assessed by two reviewers using the Critical Appraisal Skills Programme (CASP) RCT checklist [21]. Articles were assessed for risk of bias with any disagreement between reviewers resolved through discussion. In line with the Cochrane method, no quality threshold was defined with which to exclude studies, and only trials with RCT designs included [22].

Data Extraction, Synthesis and Analysis

Study details were extracted to a trial description form. Where sufficient information was available, Forrest plots were constructed using standardised mean differences for MEP, FVC, FEV₁ and cough, based on post intervention means.

Statistical heterogeneity was assessed using the inconsistency statistic I^2 [23]. All analyses were performed using Review Manager Software, version 5.0 (Cochrane Collaboration 2011, 5.0).

Results

Study Inclusion

Literature searching identified 426 titles with 141 duplicates. Of the 285 abstracts screened, 31 were retrieved for full text review, of which nine [7,17,18,24-29] met all inclusion criteria and underwent quality assessment, data extraction and analysis (Fig. 1).

Methodological Quality

All studies reported random allocation of participants in an effort to reduce selection bias, however assessment of true randomisation was not possible as no studies provided recruitment sampling methods. Blinding of participants and staff to group allocation was performed in six studies [7,24-29]. Compliance of attrition reporting was high with rates from 0 [17,29] to 23 [26] patients, with varied reasons for attrition. Of note, Gosselink *et al.* [18] reported patients who were unable to perform MEP measurements due to severely impaired lung function.

Study Description: RCTs investigated EMST in healthy individuals (2 [17,29]), patients with COPD (2 [35,28]) and subjects with neurological conditions including MS (3 [7,18,27]), spinal cord injury (SCI; 1 [26]) and acute stroke (1 [24]).

Participants

The number of study participants ranged from 12 [17] to 42 [24], with 236 participants across all studies. Only Gosselink *et al.* [18] and Kulnik *et al.* [24] performed power analysis for sample size calculation.

Subjects' mean age ranged from 24 [29] to 66 years [25]. As neither Smeltzer *et al.* [7] nor Silverman *et al.* [27] reported participant age, a mean age across studies could not be calculated (Table 1).

Intervention and comparators

All studies used threshold/resistance loading of the expiratory muscles although training devices varied between studies (Table 2) with five using a modified Threshold Inspiratory Muscle Trainer [7,17,18,25,28] and three using a Threshold Positive Expiratory Pressure (PEP) device [24,27,29]. Roth and colleagues [26] used a closed end, high-pressure force meter with subjects exhaling maximally against a pressure gauge. All studies assessed MEP as a primary outcome measure with three including cough [18,24,29] or spirometry outcomes [25,26,29].

Of the included studies, only those in healthy subjects [17,29] compared EMST to a control with no intervention. All remaining studies included breathing exercises [18] or sham EMST as a comparator [7,24-28].

Outcome 1: Expiratory Muscle Strength

Overall meta-analyses included a total of 213 patients and favoured EMST intervention with a small but significant improvement in MEP of 15.95 cmH₂O (95% CI: 7.77 to 24.12; p<0.001; Fig 2A) compared with control/sham. To investigate sources of heterogeneity (I^2 >50%) across studies, sub-analyses were performed by pathology.

Healthy Populations

In both RCTs investigating EMST with healthy subjects [17,29], populations were of similar age (mean age \leq 30) with baseline MEP values in line with reported norms [30] and no significant differences between groups at baseline. Both studies used a 4-week 30% threshold training programme comprising two 15-

minute sets of training daily for 4 weeks. Each reported significant increases in MEP versus control. Meta-analysis comprised 34 healthy participants, with high homogeneity ($I^2=0\%$), suggesting EMST significantly improved MEP with a mean difference of 33.62cmH₂0 (95%CI 16.38 to 50.85; p<0.0001) versus control (Fig 2B).

COPD

Two studies investigated MEP in patients with COPD [25,28]. Whilst both reported no difference between groups at baseline, actual MEP values varied considerably between studies. Weiner *et al.* [28] reported lower baseline MEP than predicted values for the equivalent age category [31], whilst Mota *et al.* [25] recorded pressures greater than some reported norms for MEP [30] Despite these differences, both studies trained muscle endurance with 30-minute training sessions at 50-60% of MEP, although Weiner [28] used a longer 12-week training period. Forrest plots identified good homogeneity between studies (I^2 =0%) and meta-analysis, with a total of 39 patients with COPD, suggested EMST significantly improved MEP in COPD with a standard mean difference of 19.93cmH₂0 (8.88 to 30.97; p=0.0004) versus control (Fig 2C).

Multiple Sclerosis

Three studies investigated MEP in patients with MS [7,18,27]. Gosselink (2000) and Smeltzer (1996) reported reduced baseline MEP to 29% [18] and 36.9% [7] of predicted normative values respectively. The most recent study [27] reported higher baseline MEP values with no comparison to normal values.

All three studies used a similar protocol using daily sets (15-25 repetitions) of MEP at 50%, 75% or maximal resistance. The earlier studies [7,18] both

employed a 12-week protocol and reported significant improvements in MEP following EMST. Meta-analysis of the three studies, with a total of 69 patients showed considerable heterogeneity (I^2 =61%) and no significant change in MEP following EMST (11.29cmH₂0 (-0.32 to 22.91; p<0.06) versus sham (Fig 2D).

Stroke

One study [24] investigated EMST in patients within 2 weeks of stroke onset (n=42). Baseline MEP values were not significantly different between groups but reduced compared to age-matched normal values. Only 52% of patients completed more than 70% of the 4-week training programme with significant improvement in MEP in intervention *and* control groups (Mean (SD) change in MEP EMST 12(15)cmH₂0 versus sham 12(18) cmH₂0, p=0.35).

Spinal Cord Injury

One RCT [26] investigated EMST in 29 patients with motor-complete spinal cord injury at or above T1. There was no difference between groups at baseline with MEP lower than age-matched norms [31]. Following a strength training protocol of 10 maximal expirations twice daily for 6-weeks, significant improvements in MEP were seen in the training group versus sham (Mean (SD) change in MEP EMST 35(38.4)cmH₂0 versus sham 8(19.1) cmH₂0, p=0.002).

Outcome 2: Cough

Three studies included cough as a secondary outcome measure [18,24,29]. Sasaki [29] and Kulnik *et al.* [24] assessed PECF outcomes following 4-week EMST in healthy subjects and patients following acute stroke, respectively. Baseline PECF was normal in patients after acute stroke [24]. Both intervention and sham groups demonstrated increases in PECF over time, with no significant difference between groups. In healthy subjects, Sasaki [29] found no change in

PECF following EMST. Despite differences in age and baseline MEP, PECF data between studies showed good homogeneity (I^2 =8%) and indicated no significant impact of EMST on PECF (4.63L/min; 95%CI -27.48 to 36.74; P=0.78; Figure 3).

Gosselink *et al.* [18] assessed cough efficacy by means of the validated pulmonary index (PI) for patients with MS [32], reporting significant improvements versus control following EMST. The PI includes patient- and examiner-rated ability to clear the airway, however the authors provided no detail on components of the index that improved following training.

Outcome 3: Pulmonary Function

Three studies [25,26,29] included pulmonary function as outcome measures, reporting FEV1 or FVC. Roth *et al.* [26] demonstrated small increases in FVC and FEV₁ in intervention and sham groups, whilst Mota [25] and Sasaki [29] found no change in any measure of pulmonary function following EMST. Forrest plots indicated good homogeneity (I^2 =0%) with no evidence to support EMST in improving FVC (-0.16L; 95%CI -0.35 to 0.02; P=0.09; Fig 4A) or FEV₁ (-0.09L; 95%CI -0.10 to -0.08; P<0.001; Fig 4B).

Discussion

This review aimed to assess the evidence for EMST on maximum expiratory pressure in different adult populations. Meta-analysis indicated a small but significant increase in expiratory pressure following EMST, representing an improvement of approximately 15% in MEP. There was no evidence of any effect of EMST on peak cough flow or spirometry.

As expiratory training is a potential intervention to improve airway clearance, determining its effectiveness by clinical population is paramount. Further meta-

analysis by patient population also indicated an increase in MEP following expiratory training that reached significance in young healthy adults and those with COPD, but not in patients with MS. This is in line with a recent Cochrane review of respiratory training in MS that failed to find sufficient evidence to support expiratory training or breathing exercises on MEP [33].

Only single RCTs were available in patients with SCI or stroke, precluding metaanalysis. Similar to MS findings, data in patients following acute stroke showed no significant increase in MEP following training, however in patients with spinal cord injury, maximal resistance training significantly increased MEP. Differences in disease pathology across these groups may impact patients' ability to comply with training and account for some of this variation. Facial weakness and reduced lip closure strength have been documented in patients with MS [34] and stroke [35] respectively, affecting ability to achieve and maintain a mouth seal during expiratory training. In the current MS studies, Gosselink et al, [18] excluded patients due to inability to generate mouth pressures, while Silverman et al, [27] mentioned insufficient facial strength as possible exclusion criteria. As each of these studies used near-maximal or maximal strength training, it is conceivable that difficulties maintaining the buccal pressures generated during EMST may have limited the effects of muscle training. Conversely in patients following SCI, where oromotor control is not routinely affected, maximal resistance training was tolerated resulting in significant improvements in MEP versus control.

In COPD, where evidence suggests that up to 20% of patients with severe disease have insufficient inspiratory strength to generate peak flow requirements for inhalation delivery devices [36], patients tolerated EMST well with a significant increase seen in MEP. This may be due to use of a lower intensity "resistive breathing" protocol (up to 60% expiratory pressure training), rather

than strength-based near maximal "threshold loading". Although resistive breathing and threshold loading have been shown to generate similar workloads in *inspiratory* training [37], the higher pressures generated during maximal resistance loading may not be tolerable in patient populations prone to respiratory muscle weakness.

The small effect size of EMST may have been due to the lack of a standardized intervention protocol and duration across studies. Firstly, the different training devices used were neither designed nor validated for expiratory muscle training. The Threshold IMT is validated as a reliable method of loading inspiratory muscles [38] but there is no equivalent study on modification for expiratory loading. Similarly, Threshold PEP was designed as an adjunct to mobilise airway secretions and prevent atelectasis, with studies validating its reliability for muscle training/loading lacking.

Secondly, a lack of training specificity may also have weakened the size of the training effect seen. In terms of muscle physiology, short 4-week periods of training are sufficient to initiate neural changes such as increased motor unit recruitment and firing rates [39], but not to elicit changes in muscle fibre strength [39]. Current studies in healthy individuals both identified increases in MEP after just 4 weeks in line with broader literature [40]. However, evidence suggests training programmes of at least 6 weeks duration, are needed to generate sustainable, measurable changes in muscle fibre hypertrophy [41]. From a clinical perspective, as post-intervention MEP values in COPD, stroke and MS studies all remained below normal for their respective age category [30], extending the duration of EMST may have been more effective in achieving outcomes closer to normal/predicted MEP levels, although patient adherence to longer interventions may prove difficult.

Gains of 15% expiratory strength may represent statistically significant changes in MEP, however the clinical significance of these changes is unclear. Critical MEP levels have been proposed (40cmH_20), below which secretions are thought to accumulate in the larynx [42], however the minimal MEP improvement needed to impact airway clearance has yet to be determined. Correlations between MEP and cough have been documented in SCI where a 10cmH₂0 increase in MEP has been shown to generate 0.15L/sec improvements in PECF [43], however the functional benefits of this have not been determined. The current meta-analysis did not support EMST effects on cough flow, with several factors likely affecting this. In addition to the paucity of data, the lack of cough flow impairment in the populations studied may have limited the potential for improvement, there may have been insufficient improvement in MEP in order to impact cough efficacy, or these findings may be the result of inadequate training specificity. It has been suggested that expiratory muscles need to be trained close to residual volume to improve expiratory capacity and flow [32], however current studies performed training closer to total lung capacity. This highlights the importance of accurate intervention design and the selection of outcome measures that are proven to be sensitive to the intervention in question. It may be that other measures of cough, such as the pulmonary index [18], are more sensitive to changes in MEP.

In line with previous quasi-experimental studies [8] current meta-analysis suggested no significant effect of EMST on either FEV₁ or FVC. Proposed explanations for this suggest elastic recoil and properties of the lung tissue, rather than expiratory muscle strength, may determine maximum expiratory flow [29]. Certainly changes to lung tissue in chronic diseases such as MS are known to impair lung compliance to a greater extent than skeletal muscle weakness [8]. One recent systematic review has suggested combined expiratory and inspiratory training may have a greater impact on pulmonary function by

enhancing inspiratory reserve volume and elastic recoil, and generating significant improvements in FEV₁ [44].

Limitations

The potential to overestimate the current treatment effect, due to the small number of eligible studies, small sample sizes, variation in age, population and intervention design, must be considered when interpreting the present results.

Conclusion

This review aimed to assess and synthesise current EMST data and determine the effect of EMST on expiratory strength, pulmonary function and cough. Nine studies examined MEP as a primary outcome measure and, despite differences in intervention protocols, meta-analysis indicated a small but statistically significant increase in MEP following EMST. Limited studies included outcome measures of pulmonary function or cough with no evidence to support any change in airflow measures following EMST.

Implications for Research

Methodological variation across studies offers direction for future research to determine the most effective training protocol for MEP gains. As cough and spirometry measures were not sensitive to MEP changes, investigation of other functional outcomes such as dyspnoea may help identify a role for improvements in MEP. Future EMST research may best be conducted as one arm of larger interventions using combined inspiratory and expiratory muscle training.

Implications for Clinical Practice

Populations prone to expiratory muscle weakness have been shown to respond to EMST, however the functional relevance of this is yet to be determined. As a link

has not yet been demonstrated between increased MEP and improved airway clearance, physiotherapists and clinicians involved in treatment of patients prone to aspiration should undertake EMST with caution.

Ethical Approval: None

Funding: This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Conflict of Interest: None

Declarations of interest: None

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Figure 1: PRISMA Flowchart of Search and Selection Criteria



Table 1: Characteristics of Included Studies

Study (year)	Patients	Population	Outcome n	neasures	9	Key Findings
Country [ref]	(N, mean age, sex)		МЕР	PFT	Cough	
Sasaki (2007) Japan [29]	N=33, 24y, 70% male (n=22 completed)	Healthy	-	FVC FEV1	PECF	* MEP improved in EMST group compared with control $(p<0.05)$, with no change in spirometry or PECF
Suzuki et al. (1995) Japan [17]	N=12, 30y, 100% male	Healthy	-	-	-	* MEP improved in EMST group compared with control $(p < 0.05)$.
Gosselink et al. (2000) Belgium [18]	N=21, 58y, 46% male (n=18 completed)	MS	-	-	PI	No significant increase in MEP post training. PI increased (p <0.05) vs baseline and control.
Silverman (2017) USA [27]	N=42, not reported, 26% male (n=36 completed)	MS	~	-	-	No significant increase in MEP post training vs sham
Smeltzer et al. (1996) USA [7]	N=15, not reported, 46% males	MS	~	-	-	* MEP increased (p<0.005) post training vs sham.
Mota et al. (2007) Spain [25]	N=16, 66y, 100% male	COPD GOLD III & IV	~	FVC FEV ₁	-	* MEP improved in EMST group vs sham (p<0.05), with no significant changes in spirometry.
Weiner et al., (2003) Israel [28]	N=26, 62y, 85% male (n=23 completed)	COPD, GOLD III & IV	~	-	-	* MEP improved in EMST group compared with control $(p < 0.05)$
Kulnik et al. (2015) UK [24]	N=42, 64y, 65% male	Stroke	~	-	PECF	MEP and PEFR improved in treatment and sham groups with no significant differences due to training.
Roth et al. (2010) USA [26]	N=29, 30y, 76% male	SCI	~	FVC, FEV ₁	-	MEP increased in training and sham groups but only reached significance in training group ($p<0.05$) with no significant changes in spirometry.

Abbreviations: N: number of subjects; y: years; MS: multiple sclerosis; COPD: chronic obstructive pulmonary disease; GOLD: Global initiative for Chronic Obstructive Lung Disease classification; SCI: spinal cord injury; MEP: maximum expiratory pressure; PFT: pulmonary function tests; FVC: forced vital capacity; FEV₁: forced expiratory volume in 1 second; PI: pulmonary index; PECF: peak expiratory cough flow; vs: versus; EMST: expiratory muscle strength training, * significant change (p<0.05) in MEP vs control/sham.

Table 7	Evoiratory	Mucclo	Ctronath	Training	Intonyontion	Daramatara
I able Z	EXDITATOL	Muscle	Suenaur	rraininu -	· Intervention	Parameters

Study (year)	Subjects	Threshold Intensity	Time/ reps	Frequency (times per week)	Duration (weeks)	Comparator	Device
Sasaki (2007) [29]	Healthy	30%	2x15 min	7	4	Control*	PEP
Suzuki et al. (1995)[17]	Healthy	30%	2x15 min	7	4	Control*	Threshold IMST
Mota et al. (2007) [25]	COPD	50%	30 min	3	5	Sham*	Threshold IMST
Weiner et al. (2003) [28]	COPD	15-60%	30 min	6	12	Control*	Threshold IMST
Gosselink et al. (2000)[18]	MS	60%	6x15 reps	7	12	Control	Threshold IMST
Silverman et al. (2017) [27]	MS	75%	5x5 reps	5	5	Sham**	PEP
Smeltzer et al. (1996) [7]	MS	Patient- selected maximum	6x15 reps	7	12	Sham*	Threshold IMST
Kulnik et al. (2015) [24]	Stroke	50%	5x10 reps	7	4	Sham**	PEP
Roth et al. (2010) [26]	SCI	Maximum resistance	2x10 reps	5	6	Sham*	Closed end pressure meter

* Significant change (p<0.05) in MEP vs control/sham; ** significant increase in MEP vs baseline but not vs sham.

Abbreviations: MS: multiple sclerosis; COPD: chronic obstructive pulmonary disease; SCI: spinal cord injury; PT: physiotherapy. *Significant (p<0.05) increase in MEP following EMST vs control/sham.

Figure 2: Forrest plots illustrating change in maximum expiratory pressure (MEP) following expiratory muscle strength training (EMST) versus control: (A) all studies; (B) healthy subjects; (C) chronic obstructive pulmonary disease (COPD); (D) multiple sclerosis (MS).

A) All studies

		EMST		c	Control			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	IV, Random, 95% CI
GOSSELINK, R. et al.,	8	14	9	-4	6.149	9	15.5%	12.00 [2.01, 21.99]	
KULNIK, S.T. et al.,	12	15	21	12	18	21	15.5%	0.00 [-10.02, 10.02]	
MOTA, S., et al., 2007	24	37	10	1	1.26	6	7.8%	23.00 [0.05, 45.95]	
ROTH, E.J., et al 2010	35	38.4	16	8	19.1	13	8.4%	27.00 [5.51, 48.49]	
SASAKI, M., 2007.	27.8	36.3	11	-3.8	6.46	11	8.3%	31.60 [9.81, 53.39]	
SILVERMAN E.P., et al., 2017	20.4	25.2	20	23.82	27.6	16	10.5%	-3.42 [-20.88, 14.04]	
SMELTZER, S.C., LEVIETES, M.H. and COOK, S.D.,	19.4	9.9	10	-1.2	11.1	5	14.4%	20.60 [9.10, 32.10]	
SUZUKI, S., SATO, M. and OKUBO, T.,	37	35.2	6	0.01	0.01	6	5.9%	36.99 [8.82, 65.16]	
WEINER, P. et al., 2003	18	22.2	12	-1	1.7	11	13.6%	19.00 [6.40, 31.60]	- • -
Total (95% CI)			115			98	100.0%	15.95 [7.77, 24.12]	•
Heterogeneity: Tau ² = 86.18; Chi ² = 20.36, df =	8 (P = 0).009);	$l^2 = 61$.%				_	-50 -25 0 25 50
Test for overall effect: $Z = 3.82$ (P = 0.0001)									Favours [Control] Favours [EMST]

Healthy

		EMST		Control				Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
SASAKI, M., 2007.	27.8	36.3	11	-3.8	6.46	11	62.6%	31.60 [9.81, 53.39]	
SUZUKI, S., SATO, M. and OKUBO, T.,	37	35.2	6	0.01	0.01	6	37.4%	36.99 [8.82, 65.16]	-
Total (95% CI)			17			17	100.0%	33.62 [16.38, 50.85]	
Heterogeneity: $Chi^2 = 0.09$, $df = 1$ (P = Test for overall effect: Z = 3.82 (P = 0	= 0.77); .0001)	l ² = 0	%					-	-50 -25 0 25 50 Favours [Control] Favours [EMST]

C)

		EMST		С	ontrol			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
MOTA, S., et al., 2007	24	37	10	1	1.26	6	23.2%	23.00 [0.05, 45.95]	
WEINER, P. et al., 2003	18	22.2	12	-1	1.7	11	76.8%	19.00 [6.40, 31.60]	
Total (95% CI)			22			17	100.0%	19.93 [8.88, 30.97]	-
Heterogeneity: Chi ² = 0.0 Test for overall effect: Z =)9, df = = 3.54 (1 (P = 0.	0.76); 0004)	l ² = 0%	5				-50 -25 0 25 50 Favours [Control] Favours [EMST]

COPD

D) MS

Study or Subgroup	Mean	EMST SD	Total	C Mean	Control SD	Total	Weight	Mean Difference IV, Random, 95% CI	Mean Difference IV, Random, 95% CI
GOSSELINK, R. et al.,	8	14	9	-4	6.149	9	39.4%	12.00 [2.01, 21.99]	
SILVERMAN E.P., et al., 2017	20.4	25.2	20	23.82	27.6	16	24.6%	-3.42 [-20.88, 14.04]	
SMELTZER, S.C., LEVIETES, M.H. and COOK, S.D.,	19.4	9.9	10	-1.2	11.1	5	36.0%	20.60 [9.10, 32.10]	│
Total (95% CI)			39			30	100.0%	11.29 [-0.32, 22.91]	
Heterogeneity: $Tau^2 = 63.20$; $Chi^2 = 5.11$, $df = 2$ Test for overall effect: $Z = 1.91$ (P = 0.06)	(P = 0.	08); I²	= 61%						-50 -25 0 25 50 Favours [Control] Favours [EMST]

Figure 3: Forrest plot of change in peak expiratory cough flow (PECF) following expiratory muscle strength training (EMST) versus control

	1	EMST		Co	ontro	1		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
KULNIK, S.T. et al.,	49	121	21	84	146	21	15.7%	-35.00 [-116.10, 46.10]	
SASAKI, M., 2007.	30	51	11	18	30	11	84.3%	12.00 [-22.97, 46.97]	
Total (95% CI) Heterogeneity: Chi ² =	1.09, d	f = 1	32 (P = 0,	30); I ² =	= 8%	32	100.0%	4.63 [-27.48, 36.74]	
Test for overall effect:	Z = 0.2	28 (P	= 0.78)				-100 -50 0 50 100 Favours [Control] Favours [EMST]		

Figure 4: Forrest plot illustrating change in A) Forced vital capacity (FVC) or B) Forced expiratory volume in 1 second (FEV₁) following expiratory muscle strength training (EMST) versus control.

A)									FVC
Study or Subgroup	E Mean	MST SD	Total	C Mean	ontrol. SD	Total	Weight	Mean Difference IV, Fixed, 95% CI	Mean Difference IV, Fixed, 95% CI
MOTA, S., et al., 2007 ROTH, E.J., et al 2010 SASAKI, M., 2007.	0.16 0.28 -0.3	0.3 0.4 0.5	10 13 11	0.24 0.34 0.001	0.3 0.59 0.001	6 13 11	37.5% 23.0% 39.5%	-0.08 [-0.38, 0.22] -0.06 [-0.45, 0.33] -0.30 [-0.60, -0.01]	
Total (95% CI) Heterogeneity: Chi ² = 1. Test for overall effect: Z	40, df = = 1.72	= 2 (F (P =	34 9 = 0.50 0.09)	0); l ² = (0%	30	100.0%	-0.16 [-0.35, 0.02]	-1 -0.5 0 0.5 1 Favours [Control] Favours [EMST]
B)									FEV ₁

Study or Subgroup	Mean	EMST SD	Total	C Mean	ontrol SD	Total	Weight	Mean Difference IV, Fixed, 95% CI		Mean Difference IV, Fixed, 95% CI	
MOTA, S., et al., 2007	0.01	0.05	10	0.07	0.05	6	5.1%	-0.06 [-0.11, -0.01]			
ROTH, E.J., et al 2010	0.26	0.02	13	0.35	0.01	13	87.5%	-0.09 [-0.10, -0.08]			
SASAKI, M., 2007.	-0.2	0.05	11	-0.1	0.05	11	7.4%	-0.10 [-0.14, -0.06]		=	
Total (95% CI)			34			30	100.0%	-0.09 [-0.10, -0.08]		•	
Heterogeneity: Chi ² = 1. Test for overall effect: Z	55, df = = 15.38	= 2 (P = 8 (P <	= 0.46) 0.0000); I ² = 0)1)	%				-1	-0.5 0 0.5 Favours (Control) Favours (EMST)	1