

ORIGINAL RESEARCH

Effects of inspiratory muscle training on dynamic hyperinflation in patients with COPD

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Abstract: Dynamic hyperinflation has important clinical consequences in patients with chronic obstructive pulmonary disease (COPD). Given that most of these patients have respiratory and peripheral muscle weakness, dyspnea and functional exercise capacity may improve as a result of inspiratory muscle training (IMT). The aim of the study was to analyze the effects of IMT on exercise capacity, dyspnea, and inspiratory fraction (IF) during exercise in patients with COPD. Daily inspiratory muscle strength and endurance training was performed for 8 weeks in 10 patients with COPD GOLD II and III. Ten patients with COPD II and III served as a control group. Maximal inspiratory pressure (Pimax) and endurance time during resistive breathing maneuvers (tlim) served as parameter for inspiratory muscle capacity. Before and after training, the patients performed an incremental symptom limited exercise test to maximum and a constant load test on a cycle ergometer at 75% of the peak work rate obtained in the pretraining incremental test. ET was defined as the duration of loaded pedaling. Following IMT, there was a statistically significant increase in inspiratory muscle performance of the Pimax from 7.75 ± 0.47 to 9.15 \pm 0.73 kPa (P < 0.01) and of tlim from 348 \pm 54 to 467 \pm 58 seconds (P < 0.01). A significant increase in IF, indicating decreased dynamic hyperinflation, was observed during both exercise tests. Further, the ratio of breathing frequency to minute ventilation (bf/V'_p) decreased significantly, indicating an improved breathing pattern. A significant decrease in perception of dyspnea was also measured. Peak work rate during the incremental cycle ergometer test remained constant, while ET during the constant load test increased significantly from 597.1 ± 80.8 seconds at baseline to 733.6 ± 74.3 seconds (P < 0.01). No significant changes during either exercise tests were measured in the control group. The present study found that in patients with COPD, IMT results in improvement in performance, exercise capacity, sensation of dyspnea, and improvement in the IF prognostic factor.

Keywords: chronic obstructive pulmonary disease, cycle ergometer test, dynamic hyperinflation, dyspnea, inspiratory muscle training

Introduction

Chronic obstructive pulmonary disease (COPD) is a common disease that can substantially impair one's quality of life and increase the risk of premature death.

It is a progressive disorder characterized by a partially reversible expiratory airflow limitation. The pathological hallmarks of COPD are inflammation of the peripheral airways and destruction of lung parenchyma or emphysema. It encompasses both emphysema and chronic bronchitis. The airflow limitation is usually progressive and is associated with an abnormal inflammatory response of the lungs to inhaled noxious particles or gases.

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http://dx.doi.org/10.2147/COPD.\$23784

Expiratory airflow limitation and loss of the lung's elastic recoil promotes air trapping with an increase in end-expiratory lung volume (EELV), a decrease in inspiratory capacity (IC), and lung hyperinflation. Consistent reduction in IC during physical activities reflects dynamic hyperinflation. Dynamic hyperinflation appears to be the main cause of exertional breathlessness and exercise performance in patients with COPD.¹

Inspiratory muscle training (IMT) is known to be "an adjunctive therapy in pulmonary rehabilitation, primarily in patients with suspected or proven respiratory muscle weakness." IMT positively influences exercise capacity in COPD, but its effects on dynamic hyperinflation are still controversial.

We examined the benefits of a short outpatient program of IMT on inspiratory muscle performance, exercise capacity, perception of dyspnea, and inspiratory fraction ([IF] = IC/total lung capacity [TLC]). The objective of this study was to analyze the effects of IMT on IF during exercise, exercise capacity, and sensation of dyspnea in patients with COPD.

Materials and methods Subjects

Twenty subjects with moderate to severe COPD GOLD II and III were recruited from our outpatient clinic for the study. According to the criteria of the American Thoracic Society (ATS),⁴ COPD is defined as postbronchodilator forced expiratory volume in 1 second (FEV₁) of 30% to 79% of the predicted value and a ratio of FEV₁ to forced vital capacity (FVC) <70%.

Ten subjects (6 male and 4 female) underwent IMT, while the other ten patients (5 male and 5 female) served as a control group. The baseline characteristics of the patients are summarized in Table 1. There were no significant differences in anthropometric data, lung function parameters, and exercise capacity between groups.

All patients were initially observed over a 4-week observation period while their regular treatment was maintained. During this period, no patients experienced symptoms of respiratory infection. Inclusion criteria required subjects to be less than 70 years old and to have moderate to severe nonreversible airflow obstruction. Exclusion criteria were the exacerbation of COPD within the previous 6 weeks, neoplastic disease or the presence of a disease that could contribute to dyspnea, or exercise limitation (cardiovascular, neuromuscular, or other respiratory diseases). No patients had received treatment with systemic cortisone over the last 6 months.

Table I Baseline characteristics of the patients

Parameter	Control group	Training group
Subjects (n)	10	10
Male/female	5/5	6/4
Age (years)	60.3 ± 5.3	58.7 ± 5.2
Height (cm)	171.0 ± 6.0	171.0 ± 7.6
Weight (kg)	71.3 ± 13.9	71.4 ± 17.4
BMI (kg·m ⁻²)	24.2 ± 3.4	24.3 ± 4.0
Lung function parameters		
FEV ₁ (L)	1.85 ± 0.37	1.97 ± 0.38
FEV ₁ (%)	$\textbf{54.80} \pm \textbf{8.17}$	55.90 ± 6.70
FVC (L)	3.47 ± 0.40	3.62 ± 0.69
RV (L)	2.59 ± 0.63	2.69 ± 0.64
FEV ₁ :FVC (%)	58.5 ± 6.9	60.2 ± 5.6
TLC	7.120 ± 0.434	7.025 ± 1.267
IC	3.039 ± 0.203	3.107 ± 1.106
Exercise performance		
Work rate in watts (min/max)	105 ± 28	109 ± 43

Note: Data are presented as mean \pm 1 SD.

Abbreviations: BMI, body mass index; FEV₁, forced expiratory volume in I second; FVC, forced vital capacity; IC, inspiratory capacity; RV, residual volume; TLC, total lung capacity.

Study design

The effects of IMT were investigated by using a randomized controlled study. All measurements were performed before the IMT and within 1 week after its completion. After familiarization with testing procedures, subjects underwent pulmonary function testing. Pulmonary function testing is a symptom-limited incremental test that defines the peak work rate according to ATS standards. A constant work test at 75% of peak work rate taken was performed the following day. Ten subjects received specific inspiratory muscle training on a daily basis for 8 weeks, with the other ten patients serving as a control group.

All subjects provided their written consent for the study, which was approved by the Vienna Health Authority research ethics committee.

Baseline pulmonary function test

In order to confirm diagnosis and determine severity of the disease, standard forced expiratory spirometry (FEV₁ and FVC) and body plethysmography (residual volume, functional residual capacity, TLC) were evaluated in both groups, according to ATS⁴ before and after IMT using a computerized spirometer (Masterlab; Jaeger, Wuerzburg, Germany).

Incremental cycle ergometer testing and constant load cycle ergometer testing

At the start and the end of the study, all patients performed a symptom-limited incremental exercise test on the Ergoline 800 electronically braked cycle ergometer (Ergoline, Bitz, Germany). Incremental exercise tests consisted of a steady state resting period of 3 minutes followed by stepwise protocol increases of 20–30 watts every 2 minutes depending on the baseline fitness of the participant. The test ended at the point of peak exercise, which is defined as an inability to maintain the load reached. A constant load test on the electronically braked cycle ergometer was set at 75% of the peak work rate obtained in the pre-training incremental test.

During these tests, gas exchange, ventilatory parameters, and heart rate were recorded breath by breath (Oxycon Pro; Jaeger). Constant load test endurance time (ET) was defined as the duration of loaded pedaling. IC and IF were measured every second minute during both ergometer exercise tests and also served as parameter for dynamic hyperinflation. It is assumed that TLC remains constant during exercise in COPD.⁶

In order to reach satisfactory reproducibility of IC maneuvers, patients were given a few breath warnings before the maneuver started. Subjects were instructed to breathe in until their lungs were full and then asked to try to fill up even more (ie, to inspire to TLC) before returning to normal breathing. If efforts appeared submaximal, then IC was not accepted.

The sensation of dyspnea during exercise was measured every 2 minutes using the Borg Category Ratio scale.⁷ Measurements of IC, BORG dyspnea ratings, and ET during cycle exercise testing are known to be highly reproducible and responsive to change in patients with COPD.⁸

Inspiratory muscle training

Subjects were instructed to perform IMT sessions once per day, 7 days per week, for 8 weeks. Daily training sessions started with strength training and were followed by endurance training 15 minutes later. Patients had to perform the training seated and using a nose clip.

Daily inspiratory muscle strength and endurance training were performed using the same training device (Respifit STM; Mauerbach, Austria). Maximal inspiratory pressure (Pimax) and ET during resistive breathing maneuvers (tlim) served as parameters for inspiratory muscle capacity. Pimax is a reliable, noninvasive parameter for assessing the respiratory muscle strength.

Strength exercises

Patients performed ten inspiratory maneuvers against close-to-maximum resistance and had to reach at least 80% of Pimax for a minimum of 1 second in order to complete the

strength training. This subjects the respiratory muscles, and the diaphragm in particular, to maximum strain. The strength exercise involves the constriction of the air duct allowing only for a minimal amount of air to be inhaled. Each inspiratory maneuver starts from the point of maximum expiration (residual volume).

Each training (block of exercises) included ten individual exercises. If all individual exercises were completed successfully, the training was considered a success and was stored as such on the chipcard.

The length of the break between each individual exercise was 10 seconds. If the specified minimum pressure was not reached twice during the training session, training was terminated and stored as unsuccessful on the patient's chipcard. Training intensity was increased every 2 weeks by reevaluating Pimax. Fifteen efforts were performed at level functional residual capacity to determine Pimax and the highest value was used.

Endurance exercises

The exercise was performed by breathing against a predefined level of resistance. tlim, a parameter of respiratory muscle endurance, is defined as the time until the onset of muscle exhaustion of the respiratory pump (muscle fatigue) during resistive breathing against a predefined inspiratory load.

During the exercise, the patients breathed at normal breathing frequency until the specified target value could not be achieved for four subsequent breaths (sign of muscle exhaustion). The subjects had to achieve a target pressure of 60% of the previously determined Pimax value. The elapsed time in seconds from the beginning of the exercise until the onset of muscle exhaustion was determined as tlim.

Inspiratory muscle endurance training consisted of ten loaded breathing cycles against a variable flow-dependent inspiratory resistance of 1-minute duration each, separated by 20-second intervals. The intensity of endurance training was also increased every 2 weeks.

A graphical display on the training device gave visual feedback on training results to indicate whether the strength or endurance training was completed successfully. The number of correctly and incorrectly performed exercises was saved on individual memory cards, enabling the physician to check each patient's training.

Statistical analysis

Results of the baseline and descriptive statistical analysis in tables are expressed as mean ± 1 standard deviation (SD)

or median \pm 95% confidence interval (CI). Values shown in figures are described as mean ± 1 standard error of the mean (SEM) or median \pm 95% CI. The Shapiro-Wilk test was used for preliminary testing to evaluate if grouped data were normally distributed (P < 0.05). Based on general linear models, univariate analysis of variance with repeated measures (Greenhouse-Geisser correction) was used to compare the effects of IMT on: (1) inspiratory muscle capacity (Pimax, tlim); (2) incremental exercise capacity (IF, V'O₂, bf/V'_E); (3) constant work rate (ET, bf/V'_E); and (4) sensation of dyspnea (BORG Scale). Analysis of variance results between groups and between subjects (both with Bonferroni correction for pairwise comparisons) and interactions with values of P < 0.05 were considered significant. Statistical analysis and graphs were generated using the SPSS (v 17; IBM Corporation, Armonk, NY), Excel 2007 (Microsoft Corporation, Redmond, WA), and R (v 2.10.1; R Foundation for Statistical Computing, Vienna, Austria) software programs. 10

Results

Effects of IMT on inspiratory muscle capacity

The training group showed a significant increase of inspiratory muscle performance following 8 weeks of daily IMT. Pimax increased by 18% (from 7.75 \pm 0.47 kPa to 9.15 \pm 0.73 kPa, respectively; P < 0.001). ET during tlim increased from 348 \pm 54 seconds to 467 \pm 58 seconds, which means an increase of 34% (P < 0.001; Table 2).

In contrast, these parameters remained unchanged in the control group.

Effects of IMT on incremental exercise capacity

Following the training period, there was a statistically significant increase of IC (from 2.98 ± 0.77 to 3.31 ± 0.89 , P < 0.001) during the incremental cycle ergometer test and therefore an increase of IF, which increased by 9.8% (from 0.41 ± 0.05 to 0.45 ± 0.05 , P < 0.001), indicating decreased dynamic hyperinflation (Figure 1; Table 3).

Exercise performance was assessed by evaluating VO_2 max and was also higher after training (increase of 2% from 1887.6 \pm 263.1 mL to 1925.0 \pm 244.3 mL). The peak work rate (watt max) remained constant.

The ratio of breathing frequency to minute ventilation (bf/V'_E) showed a significant decrease from 0.48 ± 0.11 to 0.46 ± 0.13 (P < 0.001; Figure 2; Table 3). Mean inspiratory flow increased from 2.19 ± 0.32 to 2.30 ± 0.31 (P < 0.001; Figure 3; Table 3). The level of dyspnea (BORG Scale) also showed a significant decrease (from 5.0 ± 1.0 to 4.0 ± 1.1 ; P < 0.01; Table 3).

The difference between the training group and the control group was significant for IC (P < 0.032), IF (P < 0.001), and bf/V'_E (P < 0.001) after the 8-week observation period. No changes occurred in the control group.

Effects of IMT on constant work rate

ET during the constant load test on a cycle ergometer increased significantly from 597.1 ± 80.8 seconds at baseline to 733.6 ± 74.3 seconds in the training group, which is an improvement of 22.9% (P < 0.001; Table 3).

bf/V'_E significantly improved, decreasing from 0.5 ± 0.11 to 0.48 ± 0.11 in the training group (P < 0.001; Figure 4; Table 3) and mean inspiratory flow increase from 2.16 ± 0.31 to 2.31 ± 0.30 (P < 0.001; Figure 5; Table 3).

IF showed a significant increase, from 0.43 ± 0.03 to 0.44 ± 0.03 (P < 0.001; Figure 6; Table 3), and sensation of dyspnea was significantly lower, decreasing from 7.0 ± 0.7 to 5.0 ± 0.9 (P < 0.001; Table 3).

After the 8-week observation period, differences between the training and control groups were significant for IC (P < 0.001), IF (P < 0.001), and bf/V'_F (P < 0.018).

The control group showed no significant change in ET, breathing pattern, BORG Scale, or dynamic hyperinflation.

Discussion

Both inspiratory muscle capacity and peripheral muscle force are negatively affected in patients with COPD. Skeletal muscle weakness is known to contribute to exercise

 Table 2 Respiratory muscle function parameters at the beginning and end of observation

Test	Control group		Training group		ANOVA	ANOVA		
	Baseline	Week 8	Baseline	Week 8	Pa	P b	P°	
tlim (s)	287.3 ± 48.1	254.0 ± 43.2	348.4 ± 54.4	467.4 ± 58.3	< 0.001	<0.001	< 0.001	
Pimax (kPa)	7.91 ± 0.67	$\textbf{7.57} \pm \textbf{0.73}$	7.75 ± 0.47	9.15 ± 0.73	< 0.001	0.023	< 0.001	

Notes: Data are presented as mean ± I SD unless otherwise stated. P: results of analysis of variance (ANOVA) comparison of control group versus training group with repeated measurements (within-subject effects); P: results of ANOVA comparison of control versus training group (between-group effects); P: results of ANOVA comparison of baseline versus week 8 with repeated measurements (within-subject effects).

Abbreviations: ANOVA, analysis of variance; Pimax, maximal inspiratory pressure; tlim, resistive breathing maneuvers.

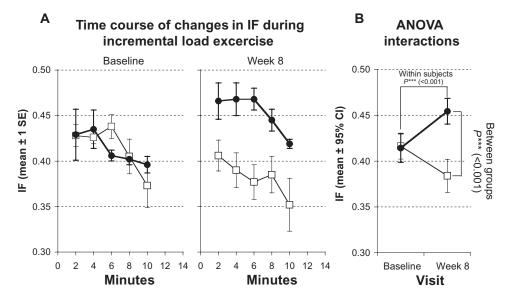


Figure I Measurements of inspiratory capacity (IF) during incremental load test. Solid symbols ●: subjects with inspiratory muscle training; open symbols □: control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means ± I standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

Note: Values represented as mean ± 95% confidence interval (CI).

limitation.^{11,12} Respiratory muscle training has been shown to positively influence the sensations of dyspnea, exercise tolerance, and quality of life,¹³ but the exact mechanisms remain unclear.

The efficiency of IMT is known to be influenced by training strategy and patient characteristics.¹⁴ The most effective method for training the inspiratory muscles has yet to be defined.

The training device used was equipped with a visual feedback system allowing the patients to control the success of the training. This proved to be important in motivating the patients, as IMT is considered to be cumbersome, boring, and without immediate effects. In addition, this feedback system enables the patient to breathe with a constant inspiratory flow during tlim. Our training protocol contained both strength and endurance components. The endurance training

Table 3 Comparison of spirometric data at the beginning and end of observation

Test	Control group		Training group		ANOVA		
	Baseline	Week 8	Baseline	Week 8	P a	P b	Pc
Constant load test							
IC	2.95 ± 0.42	2.46 ± 0.41	3.06 ± 0.47	3.11 \pm 0.58	< 0.001	< 0.001	< 0.001
IF (IC/TLC)	0.41 ± 0.05	0.34 ± 0.05	0.43 ± 0.03	0.44 ± 0.03	< 0.001	< 0.001	< 0.001
bf/V' _F	0.52 ± 0.09	0.54 ± 0.09	0.50 ± 0.11	$\textbf{0.48} \pm \textbf{0.11}$	< 0.001	0.018	0.649
Vt/ti (L/sec)	2.22 ± 0.27	2.19 ± 0.27	2.16 ± 0.31	2.31 ± 0.30	< 0.001	0.184	< 0.001
$V'O_{2}$ (mL·min ⁻¹)	1644.8 ± 208.9	1617.6 ± 190.1	1779.7 ± 268.8	1805.3 ± 276.8	< 0.001	0.148	0.901
BORG (±95% CI)	4.0 ± 0.9	6.0 ± 1.0	7.0 ± 0.7	5.0 ± 0.9	< 0.001	0.215	0.124
ET (s)	606.0 ± 126.4	573.4 ± 118.5	597.1 ± 80.8	733.6 ± 74.3	< 0.001	0.100	0.002
Incremental load test							
IC	2.98 ± 0.35	2.74 ± 0.44	2.98 ± 0.77	3.31 ± 0.89	< 0.001	0.032	0.125
IF (IC/TLC)	0.42 ± 0.05	$\textbf{0.38} \pm \textbf{0.06}$	0.41 ± 0.05	0.45 ± 0.05	< 0.001	0.001	0.268
bf/V' _E	$\textbf{0.57} \pm \textbf{0.14}$	$\textbf{0.59} \pm \textbf{0.14}$	$\textbf{0.48} \pm \textbf{0.11}$	$\textbf{0.46} \pm \textbf{0.13}$	< 0.001	< 0.001	0.780
Vt/ti (l/s)	2.21 ± 0.33	2.20 ± 0.33	2.19 ± 0.32	2.30 ± 0.31	< 0.001	0.675	< 0.001
$V'O_{2}$ (mL·min ⁻¹)	1753.6 ± 226.2	1712.9 ± 209.1	1887.6 ± 263.1	1925.0 ± 244.3	< 0.001	0.118	0.845
BORG (±95% CI)	4.0 ± 0.9	5.0 ± 1.2	5.0 ± 1.0	4.0 ± 1.1	< 0.001	0.738	0.282

Notes: Data are presented as mean ± 1 SD unless otherwise stated. P¹: results of analysis of variance (ANOVA) comparison of control group versus training group with repeated measurements (within-subject effects); P²: results of ANOVA comparison of control versus training group (between-group effects); P³: results of ANOVA comparison of baseline versus week 8 with repeated measurements (within-subject effects).

Abbreviations: bf, breathing frequency; BORG, Borg Category Ratio scale (I–10); Cl, confidence interval; ET, endurance time; IC, inspiratory capacity; IF, insipratory fraction; TLC, total lung capacity; tlim, V'_p, minute ventilation; V'O_p, peak oxygen uptake; Vt/ti, mean inspiratory flow.

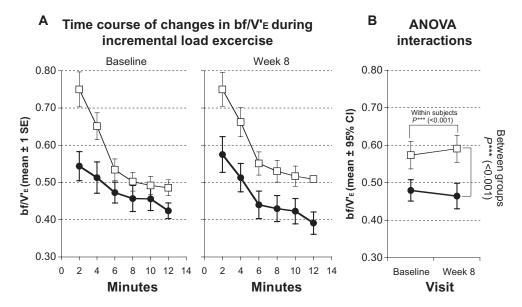


Figure 2 Measurements of breathing frequency (bf/V_E^r) during incremental load test. Solid symbols \bullet : subjects with inspiratory muscle training; open symbols \square : control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means \pm 1 standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

Notes: Values represented as mean \pm 95% confidence interval (CI).

was performed with at least 60% of Pimax because training with low pressures has been shown not to improve inspiratory muscle function.¹⁵

Dynamic hyperinflation appears to be the main cause of exertional breathlessness and exercise performance in patients with COPD. Hyperinflation causes profound dyspnea and activity limitations, which have very significant long term consequences. ¹⁶ The primary goal for treatment of

COPD patients should therefore be the deflation of the lungs, thus achieving a greater inspiratory capacity and relieving shortness of breath.

Pharmacological therapy such as long-acting bronchodilators has been documented to improve dynamic small airway function, leading to lung deflation and overall improvement in exercise performance.^{17,18} In addition, a number of non-pharmacological therapies can also reduce dynamic

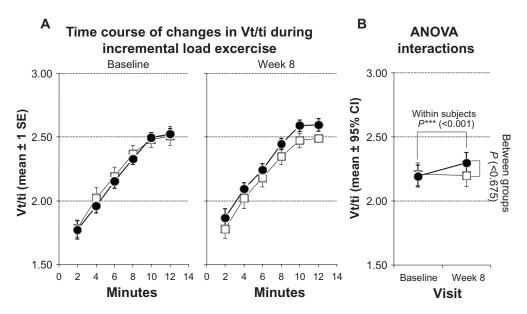


Figure 3 Measurements of mean inspiratory flow (Vt/ti) during incremental load test. Solid symbols ●: subjects with inspiratory muscle training; open symbols □: control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means ± I standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

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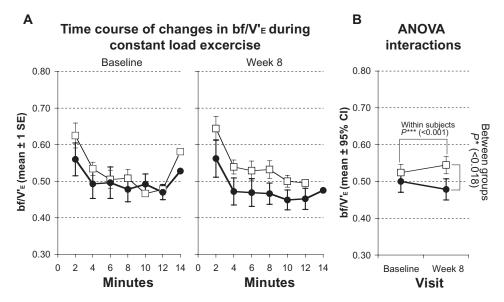


Figure 4 Measurements of breathing frequency (bf/V_{ϵ}) during constant load test. Solid symbols \bullet : subjects with inspiratory muscle training; open symbols \square : control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means ± 1 standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

Note: Values represented as mean \pm 95% confidence interval (CI).

hyperinflation such as oxygen therapy,¹⁹ heliox,^{20,21} lung volume reduction surgery,^{22,23} and general exercise training.²⁴

The most important finding of our study is the observed increase in exercise capacity and the significant reduction in dynamic hyperinflation, as reflected by the significant increase in IF. This is the first study showing positive effects of IMT on dynamic hyperinflation in patients with COPD. Therefore, apart from the aforementioned pharmacological

and non-pharmacological treatment modalities, IMT can be seen as an additional therapy modality to reduce dynamic hyperinflation.

IF can predict mortality in patients with COPD²⁵ and we hypothesized that IMT positively influences the survival rate of COPD patients. One mechanism for the reduced dynamic hyperinflation induced by IMT may be the optimization of breathing patterns during exercise.

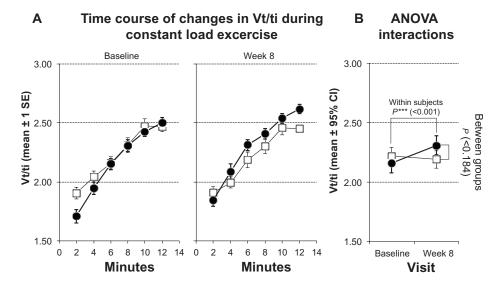


Figure 5 Measurements of mean inspiratory flow (Vt/ti) during constant load test. Solid symbols ●: subjects with inspiratory muscle training; open symbols □: control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means ± I standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

Note: Values represented as mean ± 95% confidence interval (CI).

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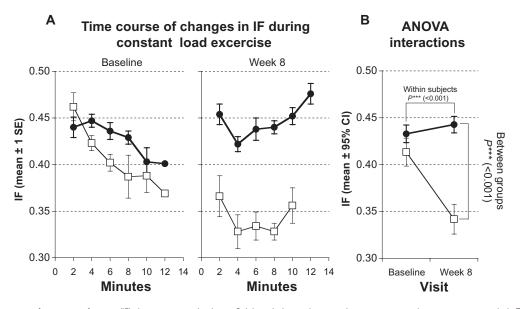


Figure 6 Measurements of inspiratory fraction (IF) during constant load test. Solid symbols ●: subjects with inspiratory muscle training; open symbols □: control group. (A) Time courses of changes at baseline and after 8 weeks. Values represented as stepwise means ± I standard error (SE) of the mean every 2 min; (B) results of analysis of variance (ANOVA) (repeated measurement design) representing interactions of within-subject and between-group effects.

Note: Values represented as mean ± 95% confidence interval (CI).

Significant increase in the size of type 2 muscle fiber following IMT in patients with COPD were observed, ²⁶ and enhancement of velocity of inspiratory muscle shortening ²⁷ was demonstrated. We hypothesize that IMT shortens inspiratory time. This allows more time for exhalation and relaxation and may reduce dynamic hyperinflation, facilitating lung emptying. This study showed that increased muscle capacity led to improvements in breathing patterns, with a significant decrease of the fraction breathing frequency (bf) to minute ventilation (Ve) and an increase of mean inspiratory flow both during incremental and constant load exercises. This pattern of breathing, in conjunction with greater respiratory muscle strength and endurance, seems to be the most important reason for the observed reduction in breathlessness during both kinds of exercise tests.

In conclusion, inspiratory muscle strength and endurance training leads to an improvement of exercise capacity and respiratory muscle endurance performance and reduces dynamic hyperinflation and breathlessness during exercise.

The results of this study may enhance the understanding of the mechanisms leading to improved exercise capacity when using IMT. Further studies are needed to verify the potential role of IMT during exercise rehabilitation programs in COPD patients.

Disclosure

The authors report no conflicts of interest in this work. This study has been registered at ClinicalTrials.gov (NCT00469313).

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