



# Effects of heated and humidified high flow gases during high-intensity constant-load exercise on severe COPD patients with ventilatory limitation



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## ABSTRACT

**Introduction:** High flow nasal cannula (HFNC) was shown to washout the anatomical dead space, permitting a higher fraction of minute ventilation to participate in gas pulmonary exchanges. Moreover, it is able to guarantee the desired inhaled oxygen fraction ( $\text{FiO}_2$ ) even at high level of patient's minute ventilation by minimizing the room air entrainment. The effect of HFNC has never been investigated on stable severe COPD patients in term of endurance capacity with standardised laboratory tests.

**Method/Design:** We performed, in a randomized crossover study, two constant load exercise tests at the 75% of maximum workload achieved at a previous incremental exercise test on cycle-ergometer: with (HFNC-test) and without HFNC (Control-test). Both constant load tests were fulfilled at the same inhaled oxygen fraction ( $\text{isoFiO}_2$ ).

**Results:** The endurance time significantly increased in the HFNC-test compared to the Control-test (the mean difference between the two groups was  $109 \pm 104$  s,  $p < 0.015$ ). At iso-time, HFNC-test showed a better oxygen saturation ( $95 \pm 3\%$  vs  $89 \pm 3\%$ , respectively,  $p < 0.005$ ) either in the subgroup of patients who used supplemental oxygen and in the subgroup who did not. Moreover, a significantly lower dyspnea (median of 5.5 vs 10, respectively,  $p = 0.002$ ) and leg fatigue score (median of 5 vs 9.5,  $p = 0.002$ ) was recorded at iso-time during HFNC-test.

**Conclusion:** HFNC may improve the exercise performance in severe COPD patients with ventilatory limitation. This effect is associated to an improvement of  $\text{SaO}_2$  and perceived symptoms at iso-time. In a Pulmonary Rehabilitation program HFNC may allow a given high intensity load to be sustained for a longer time with less symptoms.

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## 1. Introduction

Chronic obstructive pulmonary disease (COPD) patients often complain about exercise limitation as a main symptom, due to a severe dyspnea and/or muscular weakness [1]. Current literature clearly shows the benefit of pulmonary rehabilitation in symptomatic COPD [2,3]. However, these patients are frequently unable to sustain a workload high enough to obtain full benefits from the training program [2]. The most important mechanism underlying the exercise-induced dyspnea is an imbalance between ventilatory

capacity and ventilatory demand [2]. The development of hypoxia [4] and the increase of dead space [5] explain to a large extent the out-of-proportion increase in ventilation during effort, reaching prematurely the ventilatory reserve. Recent studies performed on heated and humidified high flow gases delivered through nasal cannula (HFNC), showed to (i) decrease respiratory rate, (ii) increase tidal volume (iii) and reduce the work of breathing [6] in patients with different forms of respiratory failure. The main mechanism of action is the dead space washing-out, permitting a higher fraction of minute ventilation to take part in gas exchange. Moreover, HFNC is able to guarantee the desired inhaled oxygen fraction ( $\text{FiO}_2$ ) even in patients with very high inspiratory flow rates (like during exercise) better than any other oxygen delivery system [6]. We postulated that, in severe COPD patients with exercise

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limitation, the HFNC could improve the efficiency of ventilation, leading to an increase in their exercise performance.

To assess this hypothesis, we performed a randomized crossover study evaluating the effect of HFNC on the endurance time of stable severe COPD patients.

## 2. Materials and methods

### 2.1. Participants

This was a single-centre, pilot study with a randomized crossover design performed on 12 clinically stable severe COPD patients admitted to our Pulmonary Rehabilitation Unit for an inpatient exercise training program. Patients with Forced Expiratory Volume in 1 s (FEV1) < 50% of predicted value and a 6-min walked distance < 75% of the predicted value [7] associated to a ventilatory limitation were included in the study. Ventilatory limitation was assessed by measuring the Minute Ventilation with a pneumotachograph (Spiropalm®, Cosmed-Italy) during the 6-min walking test and was defined as the ratio of peak-Minute Ventilation and Maximal Voluntary Ventilation (peakVe/MVV) > 70%, or MVV-peakVe < 11 L/min [8]. Exclusion criteria were: left heart failure, pulmonary disorder other than COPD, neuromuscular disease, osteoarticular limitation, recent COPD exacerbation (last three months), recent change in medication status (within 1 month) and anaemia. All enrolled patients signed a written informed consent form. The study protocol was approved by the medical ethics committee of the Fondazione Salvatore Maugeri di Pavia (protocol number 1086).

### 2.2. Study design

The study was performed in three separate days. On the first day, the patient underwent a symptom-limited incremental exercise test on cycle-ergometer (Ergoselect100P, Cosmed-Italy) in order to assess his/her maximal exercise capacity and to determine the FiO<sub>2</sub> to be used during the two following constant load tests. If oxygen saturation (SaO<sub>2</sub>) during the incremental test reached values lower than 85%, the incremental test was repeated with a Venturi mask at a FiO<sub>2</sub> able to maintain a mean SaO<sub>2</sub> during the test higher than 88%. In the following two days, patients performed, in a random order, two constant-load, symptom-limited exercise tests at 75% of maximum workload achieved with the incremental test, with (HFNC-test) and without (Control-test) HFNC. If a Venturi mask was added during the incremental test, the same FiO<sub>2</sub> was kept in both constant-load tests. To minimize the bias due to the unblinded design, the two constant-load tests were performed randomly during the two consecutive days. Moreover, in patients who did not need additional oxygen, Control-test was actually performed with a Venturi mask connected to compressed air in place of oxygen. Because the Venturi mask is not routinely used to give oxygen during exercise in our department, all patients were instructed to receive two different tools (Venturi mask and HFNC) both potentially able to increase their exercise performance. HFNC was administered through nasal cannula (Optiflow™, Fisher&Paykel-NewZealand) using the AIRVO2® (Fisher&Paykel-NewZealand), a turbine-based system able to generate heated and humidified gas flow up to 60 L/min. Adding an oxygen flow-rate to the system, it is possible to ensure FiO<sub>2</sub> from 0.21 (if no oxygen was added) to 1.0 (if the oxygen flow added and the set flow are equal). The system allows to set a Temperature and a Flow and to measure the supplied FiO<sub>2</sub> through an inbuilt ultrasonic analyser. As the expected effects of dead space washout and enhancement of pharyngeal oxygen concentration are both flow-dependent, the Flow was set at the highest value tolerated by the patients up to 60 L/min. In the 24 h between each test, the patient was asked to

refrain from exercise and smoking and to avoid caffeine. The endurance time for each constant-load test was recorded (Tlim); furthermore peripheral SaO<sub>2</sub>, heart rate (HR), blood pressure (BP), 0–10 Borg scale for dyspnea (Borg-D) and leg fatigue (Borg-F) were recorded every minute until the end of the test.

### 2.3. Statistical analysis

Results are presented as mean ± SD or median (range). Paired *t*-test for parametrical data or Wilcoxon test for non-parametrical data was used to compare each variable in the two constant-load tests, HFNC-test and Control-test, at isotime, defined as the time the shortest test ended. A *p*-value < 0.05 was considered statistically significant.

## 3. Results

We enrolled 12 consecutive COPD patients. Anthropometric and functional data are reported in Table 1. There were no dropouts from the study. No adverse events were detected during the incremental test and during the two constant-load tests. Mean workload imposed during the constant-load tests was 44 ± 17 W. Four out of 12 patients performed the tests on ambient air, while the other 8 needed the addition of oxygen with a mean FiO<sub>2</sub> 0.44 ± 0.11. The mean flow used during HFNC was 58.7 L/min (range 55–60 L/min).

As showed in Fig. 1, Tlim was significantly higher during the HFNC-test (mean difference of 109 ± 104 s, *p* < 0.015) compared to the Control-test, resulting in a mean increase of 41 ± 36%. At isotime Borg-D and Borg-F were significantly reduced (Fig. 2), while mean SaO<sub>2</sub> significantly increased during HFNC test (*p* < 0.005) (Fig. 3A). Mean SaO<sub>2</sub> was significantly higher during the HFNC-test, both in the subgroup of patients who used oxygen (Fig. 3B) and in the subgroup that did not (89 ± 3% vs 92 ± 3% at isotime, during Control-test and HFNC-test respectively, Fig. 3 C).

## 4. Discussion

In this pilot study we demonstrated that HFNC was able to significantly improve the duration of exercise in advanced COPD patients with exercise limitation, by improving SaO<sub>2</sub> and reducing symptoms. Surely, due to the lack of measurement of tidal volume and respiratory frequency, this study was not able to explain the mechanism underlying this improvement. However we can speculate that the main mechanism may be the already described reduction of ineffective ventilation due to dead space, that is washed-out by the high flows [9]. In fact, it was demonstrated that exercise intolerance in COPD is partially due to high dead space causing a compensatory and out-of proportion increase in minute

**Table 1**  
Anthropometric and functional data.

Age, yr	70 ± 8
Male:female, n	10:2
FEV1/FVC, %	40 ± 10
FEV1, % predicted	35 ± 12
PaO <sub>2</sub> , mmHg	73 ± 13
PaCO <sub>2</sub> , mmHg	41.7 ± 5.3
peakVe/MVV, %	87 ± 12
MVV-peakVe, L/min	3 ± 4

Data are presented as mean ± SD. FEV1: forced expiratory volume in 1 s; FVC: forced vital capacity; PaO<sub>2</sub>: arterial oxygen tension; PaCO<sub>2</sub>: arterial carbon dioxide tension; peakVe: maximal minute ventilation at the end of six minute walking test; MVV: maximal voluntary ventilation.

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