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Need for Portable Oxygen Titration Using 6-Minute Walk Tests[☆]



Necesidad de titular el oxígeno portátil mediante pruebas de marcha de 6 minutos

To the Editor:

Oxygen therapy improves survival, quality of life, and exercise capacity in patients with chronic obstructive pulmonary disease (COPD) and severe respiratory failure at rest.^{1–3} Portable oxygen (O_2) devices facilitate compliance with oxygen therapy and help avoid restrictions in physical activity. The SEPAR guidelines for oxygen therapy recommend that O_2 flow be adjusted during stress testing to achieve a mean arterial oxyhemoglobin saturation (SpO_2) of $\geq 90\%$.⁴ The 6-minute walk test (6MWT) is the method most widely used.⁵ Often, however, O_2 is inappropriately adjusted for exercise, since it is a laborious process and can sometimes be contraindicated.^{6,7} Some countries recommend using the same O_2 flow rate as that indicated at rest or recommend increasing O_2 by an additional 1 l.⁸

The aim of this study was to determine if the prescribed oxygen flow after titrating portable oxygen therapy for the 6MWT is similar to the flow that would be indicated if an additional liter were added to the prescribed O_2 .

We prospectively included all patients with chronic respiratory failure seen in the oxygen therapy clinic between October 2015 and September 2018 who were prescribed a continuous flow portable O_2 device. They were in a stable phase, met criteria for home O_2 therapy, had the autonomy to carry out activities outside the home, and were capable of performing a 6MWT.⁶ Patients who were prescribed a device with a valve were not included in the study.

O_2 was initially adjusted to the at rest rate following SEPAR recommendations.⁴ At least 1 6MWT was then performed, using a WristOx₂ pulse oximeter, Model 3150, with continuous flow O_2 using the device that we considered most appropriate, depending on the estimated flow requirement and the patient's mobility and preferences. The 6MWTs were performed following SEPAR recommendations.⁶ If mean $SpO_2 \geq 90\%$ was not achieved in the first test, the test was repeated after a minimum rest period of 30 min, increasing flow by 1 l/min until the objective was achieved. We compared the flow rate after adjustment for the 6MWT with the flow that would be prescribed if 1 l was added to the O_2 at-rest flow rate.

The SPSS package version 20.0 was used for the statistical analysis. A descriptive analysis of patient characteristics was performed, and the Student's *t*-test was used for comparison of means. A *p* value < 0.05 was considered statistically significant.

A total of 165 patients, 113 (68.5%) of whom were men, mean age 70.9 (SD 9.31) years, were included. Mean O_2 flow prescribed for the portable device was 3.64 (SD 0.95) l/m. Seventy-seven patients (46.7%) used continuous flow concentrators and 88 (53.3%) had liquid O_2 backpacks. After titration for the 6MWT, the prescribed O_2 was only the same if 1 l had been added to the resting O_2 rate in 49 patients (29.7%) (*p* < 0.0001). We increased the O_2 flow rate in 88 patients (53.3%) and reduced it in 28 (17%). Table 1 shows the diseases causing chronic respiratory failure and the relationship between both methods for prescribing portable O_2 flow. The prescriptions coincided in 36% of the COPD patients, but in only 17.5% of the interstitial diseases, and in 11% of the patients with pulmonary hypertension, in whom desaturation with exertion is greater. Twenty-one patients refused the liquid O_2 backpack, despite requiring more than 3 l/min. In 10 patients, desaturation experienced during the 6MWT could not be corrected.

The results show the superiority of titration by 6MWT over the alternative of adding 1 l of O_2 to the resting flow rate for correcting desaturation during activities of daily living. This is because with the latter, 53.3% of patients (72.5% and 66.7%, in the case of diffuse interstitial pulmonary disease and pulmonary hypertension, respectively) continue to desaturate during exertion. Patients with chronic respiratory failure who are stable often present prolonged periods of hypoxemia that are associated with reduced exercise tolerance and an increased rate of complications, such as pulmonary hypertension, right heart failure, and polycythemia.⁹ Arterial blood gas at rest is not useful for adjusting portable O_2 flow.¹⁰ Stress tests, in contrast, allow us to assess the effectiveness of therapeutic interventions.^{5,11} The most widely used is the 6MWT,⁷ in its different modalities,^{6,12} though cycle ergometers have also been used to titrate O_2 .¹³

Other factors to bear in mind are the mobility profile of each patient, their preferences, and the mobility permitted by each of the O_2 sources.¹⁴ Thus, 21 patients (12.7%) refused to change their device to a liquid O_2 backpack, despite needing to increase their flow by more than 3 l/min O_2 , because that would limit their autonomy.

At the present time, no portable devices are available that meet the needs of the more severe patients, as the liquid O_2 backpack can only provide a flow of up to 5 l/min. In fact, despite having liquid O_2 backpacks providing 5 l/min, 10 patients in our study experienced desaturation during the 6MWT that could not be corrected, with a mean sustained SpO_2 of < 90%.

In the future, an alternative to the current titration procedures may be to individualize the provision of home oxygen supply to each patient by integrating sensors in portable O_2 devices that would measure SpO_2 in real time and automatically adjust the flow of O_2 , according to patient needs.¹⁵

In summary, it currently seems necessary to titrate the portable O_2 flow with a stress test if we want to adequately correct desaturation during exercise. Even so, this method has its limitations and is not the only factor to be taken into account when prescribing portable O_2 .

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Table 1

Underlying Diseases of Patients and Comparison of the Prescribed Oxygen Flow, After Titrating Portable Oxygen Flow Using Both Methods.

	n (%)	Prescription of the Same O ₂ Flow n (%)	Need to Increase O ₂ Flow n (%)	Need to Reduce O ₂ Flow n (%)
Total patients	165	49 (29.7)	88 (53.3)	28 (17)
COPD	86 (52.1)	31 (36)	42 (49)	13 (15)
Diffuse ILD	40 (24.2)	7 (17.5)	29 (72.5)	4 (10)
Pulmonary hypertension	9 (5.5)	1 (11.1)	6 (66.7)	2 (22.2)
Asthma	8 (4.8)	2 (25)	3 (37.5)	3 (37.5)
Heart disease	5 (3)	1 (20)	3 (60)	1 (20)
Cancer	4 (2.4)	3 (75)	1 (25)	0
Bronchiectasis	3 (1.8)	2 (66.6)	1 (33.3)	0
Lung disease of unknown etiology	5 (3)	2 (40)	3 (60)	0
Kyphoscoliosis	1 (0.6)	0	1 (100)	0
Thoracic surgery	1 (0.6)	0	1 (100)	0
Pulmonary embolism	1 (0.6)	1 (100)	0	0
Sleep apnea-hypopnea syndrome	1 (0.6)	0	0	1 (100)
Obesity hypoventilation syndrome	1 (0.6)	0	0	1 (100)

COPD: chronic obstructive pulmonary disease; IPD: interstitial lung disease.

References

- Long term domiciliary oxygen therapy in chronic hypoxic cor pulmonale complicating chronic bronchitis and emphysema. Report of the Medical Research Council Working Party. Lancet. 1981;1:681–6.
 - Miravitles M, Soler-Cataluña JJ, Calle M, Molina J, Almagro P, Quintano JA, et al. Guía española de la enfermedad pulmonar obstructiva crónica (GesE-POC) 2017. Tratamiento farmacológico en fase estable. Arch Bronconeumol. 2017;53:324–35.
 - Nocturnal Oxygen Therapy Trial Group. Continuous or nocturnal oxygen therapy in hypoxemic chronic obstructive lung disease: a clinical trial. Ann Intern Med. 1980;93:391–8.
 - Ortega-Ruiz F, Díaz-Lobato S, Galdiz-Iturri G, García-Rio F, Güell-Rous R, Morante-Velez F, et al. Oxigenoterapia continua domiciliaria. Arch Bronconeumol. 2014;50:185–200.
 - Morante F, Güell R, Mayos M. Eficacia de la prueba de 6 minutos de marcha en la valoración de la oxigenoterapia de deambulación. Arch Bronconeumol. 2005;41:595–600.
 - Vilaró J. Prueba de marcha de 6 minutos. In: Comité científico SEPAR, editor. Manuales SEPAR de procedimientos. Barcelona: Ed. P. Pernmayer; 2004. p. 100–13.
 - Wijkstra PJ, Guyatt GH, Ambrosino N, Celli BR, Güell R, Muir JF, et al. International approaches to the prescription of long-term oxygen therapy. Eur Respir J. 2001;18:909–13.
 - Celli BR, MacNee W, ATS/ERS Task Force. Standards for the diagnosis and treatment of patients with COPD: a summary of the ATS/ERS position paper. Eur Respir J. 2004;23:932–46.
 - Singh SJ, Puhan MA, Andrianopoulos V, Hernandes NA, Mitchell KE, Hill CJ, et al. An official systematic review of the European Respiratory Society/American Thoracic Society: measurement properties of field walking tests in chronic respiratory disease. Eur Respir J. 2014;44:1447–78.
 - Zhu Z, Barnette RK, Fussell KM, Rodriguez RM, Canonico A, Light RW. Continuous oxygen monitoring – a better way to prescribe long-term oxygen therapy. Respir Med. 2005;99:1386–92.
 - Puente-Maestú L, García de Pedro J, Benedetti PA. The future of exercise tolerance testing. Arch Bronconeumol. 2018;54:405–6.
 - Giovacchini CX, Mathews AM, Lawlor BR, MacIntyre NR. Titration oxygen requirements during exercise evaluation of a standardized single walk test protocol. Chest. 2018;153:922–8.
 - Galera R, Casitas R, Martínez E, Lores V, Rojo B, Carpio C, et al. Exercise oxygen flow titration methods in COPD patients with respiratory failure. Respir Med. 2012;106:1544–50.
 - Díaz-Lobato S, García-González JL, Mayoralas-Alises S. Controversias en oxigenoterapia continua domiciliaria. Arch Bronconeumol. 2015;51:31–7.
 - Mayoralas-Alises S, Carratalá JM, Díaz-Lobato S. Nuevas perspectivas en la titulación de la oxigenoterapia: ¿es la titulación automática el futuro? Arch Bronconeumol. 2019;55:319–27.
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