

# Monitoring lung function at home in teens with Duchenne muscular dystrophy

December 17 2018

---



Hand-held device and mouthpiece used in this study. Credit: DELOS Study Group

Heart and lung complications are responsible for much of the morbidity and mortality associated with Duchenne muscular dystrophy (DMD).

Although regular monitoring of pulmonary function is recommended in order to detect deterioration, compliance with routine testing, such as hospital-based spirometry, is frequently poor. A new study in the *Journal of Neuromuscular Diseases* describes the at-home use of a child-compatible, hand-held device (HHD) that makes monitoring pulmonary function in teens more convenient and provides the data needed for better disease management.

"This is the first study that demonstrates that the use of a hand-held device to assess [pulmonary function](#) is feasible in adolescent patients with DMD. Pulmonary function data obtained by this method are reliable as shown by comparison to standard spirometry results and participating patients showed a high level of adherence in the use of the device. This new approach to collect pulmonary function data, together with several analyses methods, could be used in the planning of clinical intervention trials with respiratory function endpoints or to improve routine clinical care in patients with DMD," explained Prof. Gunnar M. Buyse, MD, Ph.D., Department of Paediatric Neurology, University Hospitals Leuven (Belgium).

DMD is a debilitating neuromuscular disease that is usually diagnosed in children between the ages of 3 and 5 years. During the early teenage years, patients with DMD find their mobility impaired and pulmonary function begins to deteriorate. Clinical symptoms include the inability to cough efficiently, poor airway clearance, recurrent respiratory infections and, eventually, respiratory failure necessitating ventilator support.

"Safeguarding pulmonary function, therefore, represents a significant unmet need," said Dr. Buyse.

This study was part of a multinational, randomized, placebo-controlled Phase III study (DELOS) investigating whether the drug idebenone could improve or delay the loss of respiratory function in adolescents 10-18 years old with DMD. One aim of the trial was to compare assessment of

pulmonary function with a hand-held device to spirometry performed in a clinic or hospital.

The results showed that weekly use of the hand-held device was well accepted by adolescent trial participants over the course of a year, with average adherence exceeding 75%. Comparison of values for peak expiratory flow expressed as percent of predicted (PEF%<sub>p</sub>) collected with the hand-held device with data obtained on the same day with standard spirometry were highly correlated indicating the reliability of this method. Both assessment methods showed that idebenone treatment slowed the decline in pulmonary function compared to placebo.

Explanations for the limited compliance to hospital-based pulmonary function testing could be that physicians do not recommend testing, or patients are not followed in specialized neuromuscular centers where pulmonary function tests are available and recommended. Use of a hand-held device would offer an appropriate addition to hospital-based spirometry assessment with the advantage of allowing more frequent collection of pulmonary function data. Another possibility is that teenagers with limited mobility may be resistant to traveling to clinical centers.

Advantages of using home-based pulmonary function monitoring include the ability to obtain more frequent assessments, sometimes on a daily basis, which could facilitate early detection of pulmonary function decline. The study also showed that short observation periods (between 2 and 4 weeks) were sufficient to distinguish clinical changes in PEF%<sub>p</sub>. Finally, this method may prove to be very useful for clinical testing of new therapies for DMD.

**More information:** Gunnar M. Buyse et al, Home-Based Monitoring of Pulmonary Function in Patients with Duchenne Muscular Dystroph, *Journal of Neuromuscular Diseases* (2018). [DOI: 10.3233/JND-180338](https://doi.org/10.3233/JND-180338)

Provided by IOS Press

Citation: Monitoring lung function at home in teens with Duchenne muscular dystrophy (2018, December 17) retrieved 26 April 2024 from <https://medicalxpress.com/news/2018-12-lung-function-home-teens-duchenne.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.