The 6 Minute Walk Test and Performance of Upper Limb in Ambulant Duchenne Muscular Dystrophy Boys

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Abstract

The Performance of Upper Limb (PUL) test was specifically developed for the assessment of upper limbs in Duchenne muscular dystrophy (DMD). The first published data have shown that early signs of involvement can also be found in ambulant DMD boys. The aim of this longitudinal Italian multicentric study was to evaluate the correlation between the 6 Minute Walk Test (6MWT) and the PUL in ambulant DMD boys. Both 6MWT and PUL were administered to 164 ambulant DMD boys of age between 5.0 and 16.17 years (mean 8.82).

The 6 minute walk distance (6MWD) ranged between 118 and 557 (mean: 376.38, SD: 90.59). The PUL total scores ranged between 52 and 74 (mean: 70.74, SD: 4.66). The correlation between the two measures was 0.499. The scores on the PUL largely reflect the overall impairment observed on the 6MWT but the correlation was not linear. The use of the PUL appeared to be less relevant in the very strong patients with 6MWD above 400 meters, who, with few exceptions had near full scores. In patients with lower 6MWD the severity of upper limb involvement was more variable and could not always be predicted by the 6MWD value or by the use of steroids.

Our results confirm that upper limb involvement can already be found in DMD boys even in the ambulant phase.

Funding Statement


Introduction

The 6-minute walk test (6MWT) has been recently chosen as the primary outcome measure in several international multicentric clinical trials in Duchenne muscular dystrophy (DMD). Its reliability in a multicentric setting and natural history data have recently been reported by several groups. In the last few years there has been increasing attention to the assessment of upper limb function, as it is not captured by the most commonly used motor functional scales, such as the North Star Ambulatory Assessment (NSAA), that mainly focuses on gross motor activities. The Performance of Upper Limb (PUL) is a measure of upper limb function specifically developed for DMD. The PUL provides both a total score and subscores for the three domains (shoulder, middle and distal) that in DMD are progressively involved with a proximal to distal gradient.

The first published data have shown that the PUL is suitable for multicentric studies, is reliable and can cover the whole spectrum of activities from early signs of proximal involvement in young ambulant DMD boys to distal activities in much older non ambulant DMD boys and adults. Although as part of the natural progression of DMD, upper limb weakness generally occurs at a later stage compared to lower limb involvement, initial signs of upper limb involvement were already detected in ambulant DMD boys. These preliminary findings suggested that the scale is ideal for non ambulant boys and adults who have lost ambulation and who are therefore unable to perform the 6MWT or the NSAA, but could also be used as an additional measure in clinical trials in ambulant DMD. Using the PUL in ambulant boys would not only provide additional information on the involvement of upper limbs but could also allow to follow boys who may lose ambulation during the trial and who could therefore not be followed using the 6MWT or the NSAA. Little is known of the age when initial signs of upper limb involvement occur and
of the possible correlation with walking abilities.

The aim of this longitudinal Italian multicentric study was to evaluate the correlation between the 6MWT and the PUL in ambulant DMD boys.

**Methods**

The study is part of a longitudinal multicentric cohort project involving 11 tertiary neuromuscular centers in Italy aimed at assessing outcome measures in DMD. As part of this overall project, DMD boys and young adults have been prospectively assessed using the 6MWT and, in a parallel study performed in the last two years, the PUL. In the present study we retrospectively evaluated data obtained in all the ambulant DMD boys from the previous studies who had both PUL and 6MWT.

Patients were recruited between September 2012 and March 2014.

Patient inclusion criteria at baseline were: genetically proven DMD diagnosis, patient still ambulant and able to walk independently for at least 75 meters, no severe or moderate learning difficulties or behavioral problems. Genetic and treatment information were collected and classified following the criteria used in our previous study. All patients attending the 11 participating centers who fulfilled the inclusion criteria were enrolled in the study.

Steroid use and regimen were also noted.

Details of the training for the physiotherapists involved in the study and of the interobserver reliability for PUL and 6MWT among the centers have already been reported.

The data in this study were collected as part of previous published studies on 6MWT and PUL approved by the Ethical Committee of each center and for which informed written consent was obtained. All clinical investigations were conducted according to the principles expressed in the Declaration of Helsinki.

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**6MWT**

6MWT was performed in all DMD ambulant boys older than 5 according to the ATS guidelines, modified by having two examiners, one recording time and distances and one staying close to the patient for safety issues.

**PUL**

The PUL includes 22 items with an entry item to define the starting functional level, and 21 items subdivided into shoulder level (4 items), middle level (9 items) and distal level (8 items) dimension. For weaker patients a low score on the entry item means high level items do not need to be performed. Scoring options vary across the scale between 0-1 to 0-6 according to performance. Each dimension can be scored separately with a maximum score of 16 for the shoulder level, 34 for the middle level, and 24 for the distal level. A total score can be achieved by adding the three level scores (max total score 74).

Although this was not a prospective study and there was no design of the timing of the test, as part of our routine 6MWT is always performed at the end of the rest of the clinical assessment, after the boys have a break.

Summary statistics (N, mean, SD, Range) were used. Correlations were evaluated by the Spearman rank correlation coefficients.

**Results**

One hundred and sixty-four boys fulfilled the inclusion criteria performing both measures. Their age ranged between 5.0 and 16.17 (mean 8.82, SD 2.42). One hundred and forty-eight were on steroids and 16 were not.

**6MWT**

The 6-minute walk distance (6MWD) ranged between 118 and 557 (mean: 376.38, SD: 90.59).

**PUL**

The total scores ranged between 52 and 74 (mean: 70.74, SD: 4.66).

The shoulder domain subscore ranged between 2 and 16 (mean: 13.98, SD: 3.04). The middle domain subscore ranged between 27 and 34 (mean: 33.43, SD: 1.10). The distal domain subscore ranged between 19 and 24 (mean: 23.60, SD: 0.87).

The correlation between the two measures was 0.499. Figure 1 shows the correlation between 6MWD and PUL total scores in
the individual boys. Boys with 6MWD above 400 had, with few exceptions scores >70. Between 300 and 400 the scores were more variable and those below 300 meters rarely had full scores.

Not surprisingly, the number of boys with full or near full PUL total scores in the group with 6MWT below 300 meters was much lower. The lower total scores reflected a higher number of boys who did not have full scores both in the middle and distal domain.

Boys with 6MWD between 300 and 400 meters showed a wider range of PUL total scores and not necessarily correlated with the decreasing distance on the 6MWT. Approximately 40% of them had PUL total scores < 70, mainly due to involvement of the shoulder domain. In a few cases, the scores were lower than 60, showing that even in this relatively good range of 6MWT some boys already had early signs of middle and distal involvement. Although we did not aim to systematically assess the effect of steroids on PUL as this will be more accurately assessed in an ongoing longitudinal study, we checked if the variability in the PUL scores could be justified by possible differences in steroid treatment. This was not the case as all the patients with PUL total scores below 60 were on steroids and more than half of them on a daily regime.

When we correlated the PUL with the 6MWT, we found that the correlation was not linear as patients at the stronger end of the spectrum on the 6MWT tended to plateau with full or near full PUL total scores (70 to 74 out of 74). Over 95% of the boys who were able to walk over 400 meters on the 6MWT had total scores between 70 and 74 with over 85% having scores ≥72. None of them had scores below 12/16 in the shoulder domain and, with few exceptions they all had full scores on middle and distal domains. These finding confirm that the value of 400 meters, that has recently been used in clinical trials as a cutoff point to identify patients who are unlikely to deteriorate in a short time span, can also identify DMD boys who are at the strong end of the spectrum in regard to upper limb function. In these patients the PUL has a more limited value but its use in longitudinal studies can help to establish the trajectories of changes over time.

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Discussion

The results of this study confirm that involvement of upper limb function can already be found in ambulant DMD boys. Not surprisingly the first signs of involvement are related to the shoulder domain with a % of patients not able to perform antigravity shoulder abduction and flexion movements, such as bringing the hand or progressive weights at shoulder or eye level by the side or in front. Because of the proximal to distal gradient of progression of upper limb involvement we observed less changes in the elbow or distal domain that are classically more involved after boys lose ambulation.

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Fig. 1: Correlation of 6MWT (in meters) and PUL total scores and subscores in the three main domains.

**Conclusions**
In conclusion, our results confirm that upper limb involvement can already be found in DMD boys even in the ambulant phase. The scores on the PUL largely reflect the overall impairment observed on the 6MWT but the correlation was not linear. The use of the PUL appears to be less relevant in the very strong patients with 6MWT above 400 meters, that are often excluded from clinical trials. In patients with lower 6MWT early signs of involvement are not only limited to shoulder involvement but also, in some cases, to the loss of the ability to fully perform some activities in the middle and distal domain. The severity of upper limb involvement was variable and could not always be predicted by the overall weakness and fatigability measured by the 6MWT or by the use of steroids. The concomitant use of the PUL appears to be even more relevant in the patients with 6MWT results below 350 meters, and especially in those below 230 meters who are more likely to lose ambulation within two years\(^6\). In these boys the PUL will provide the possibility to monitor possible functional changes also during and after loss of ambulation. Longitudinal studies are in process in both ambulant and non ambulant DMD boys and adults in order to better understand the changes of the PUL scores over time and the possible effect of different variables, such as age and steroids on the progression of upper limb involvement.

**Competing Interests**

The authors have declared that no competing interests exist. They report the following disclosures: Dr. Pane, E. Mazzone, Dr. Sivo, Dr. Fanelli, Dr. De Sanctis, Dr. D’Amico, Dr. Messina, Dr. Battini, Dr. Bianco, Dr. Scutiferi, Dr. Petillo, Dr. Frosini, Dr. Scalise, Dr. Vitale, Dr. Bruno, Dr. Pedemonte report no disclosures. Dr. Mongini has served on a scientific advisory board for Teledhond Italy; has received funding for travel from Gendrome Corporation; and has received research support from AIFA (Italian Government Drug Agency) and Teledhond Italy. Dr. Pegoraro has served on a scientific advisory board for BioMarin Pharmaceutical Inc.; has received funding for travel from Genzyme Corporation; and has received speaker honoraria from MedaPharma; and receives research support from Wellstone and Teledhond Italy. Dr. Brustia, Dr. Gardani, Dr. Berardinelli, Dr. Lanzillotta, Dr. Viggiano, Dr. Cavallaro, Dr. Sframeli, Dr. Bello, Dr. Barp, Dr. Busato, Dr. Bonfiglio, Dr. Rolle, Dr. Colia, Dr. Bonetti, Dr. Palermo, Dr. Graziano, Dr. D’Angelo, Dr. Pini, Dr. Corlatti, Dr. Gorni, Dr. Baranello, Dr. Antonaci report no disclosures. Dr. Bertini is site PI for the PTC extension study of Ataluren in DMD, for the GSK study on exon skipping. Dr. Brustia, Dr. Gardani, Dr. Berardinelli, Dr. Lanzillotta, Dr. Viggiano, Dr. Cavallaro, Dr. Sframeli, Dr. Bello, Dr. Barp, Dr. Busato, Dr. Bonfiglio, Dr. Rolle, Dr. Colia, Dr. Bonetti, Dr. Palermo, Dr. Graziano, Dr. D’Angelo, Dr. Pini, Dr. Corlatti, Dr. Gorni, Dr. Baranello, Dr. Antonaci report no disclosures. Dr. Bertini is site PI for the PTC extension study of Ataluren in DMD, for the GSK study on exon skipping. He also receives funds from the Italian Teledhond, the Italian Ministry of Health and SMA Europe for observational studies on outcome measures. Dr. Politano reports no disclosure. Dr. Mercuri is site PI for the PTC extension study of Ataluren in DMD, for the GSK Prosensa and Sarepta studies on exon skipping. He also receives funds from the Italian Teledhond and SMA Europe. He has acted as advisory board for Acceleron Pharma, Shire and PTC Therapeutics, Inc, Prosensa.

**Authorship Notes**

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**References**


