Three years follow up of 6-minute walk test and North Star Ambulatory Assessment in 30 ambulant DMD boys since the beginning of steroid treatment

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OBJECTIVE

Analysis and follow up of 6-minute walk test results and NSAA values while introducing the tests as obligatory outcome measures in neuromuscular patients.

Methods: 6-minute walk test, North Star Ambulatory Assessment were performed by ambulant DMD boys aged 5 to 12 years, at baseline and 1, 2 and 3 years after initiating steroid treatment.

Patients	m distance	m distance	Mean distance	(Std. Deviation)	р	Trends of stagnation, slow or fast minimal meaningful changes in di
						concerning age at the beginning a
30	255	525	352,83	56,92	Reneated measures	
30	232	550	1359,07	72,45	ANOVA ⁺ : F(1.271, 36,865) = 9,213,	
30	202	530	√354,30	92,57		Lowering the NSAA score from 0,
30	165	510	↓325,77	101,53	p < 0.002*	beginning of follow up and from y
.	1 tients 30 30 30 30 30	itients m distance 30 255 30 232 30 202 30 165	itientsmmdistancedistance30255302323020230165510	itients m m m distance distance distance distance distance 30 255 525 352,83 30 232 550 1359,07 30 202 530 ↓354,30 30 165 510 ↓325,77	itientsm distancem distancem distancedistanceDeviation)30255525352,8356,92302325501359,0772,4530202530↓354,3092,5730165510↓325,77101,53	itientsmmmdistanceDeviationdistancedistanceDeviationp 30 255525352,8356,92 30 232550 \uparrow 359,0772,45Repeated measures 30 202530 \downarrow 354,3092,5736,865) = 9,213, 30 165510 \downarrow 325,77101,53p < 0.002*

* p<0,05

with a Greenhouse-Geisser correction



REFERENCES

Mazzone E et al. North Star Ambulatory Assessment, 6-minute walk test and timed items in ambulant boys with with Duchenne muscular dystrophy Neuromuscular Disorders 20 (2010) 712–716 Mayhew A, Cano S, Scott E, et al. Moving towards meaningful measurement: rasch analysis of the North Star Ambulatory Assessment in Duchenne muscular dystrophy. Dev Med Child Neurol2011;53:535–42 Detecting meaningful change using the North Star Ambulatory Assessment in Duchenne muscular dystrophy Mayhew AG, Cano SJ, Scot E, Eagle M, Bushby K, Manzur A, Muntoni F, North Star Clinical Network for Neuromuscular Disease. Dev Med Child Neurol2013 Nov;55(11):1046-52

MATERIALS AND METHODS

RESULTS

t progression of the disease analysing listance passed at 6MWT (<30m/year) and initially passed distance.	NSAA DMD boys	Mean points	Patients	Std. Deviation	Minimum	Maximum	p
5 points to 4 points per year from the younger to older age.	zero	↓27,07	30	5,15	18	34	Fridman test:
	1 year	↓26,73	30	5,11	18	34	N=30 Chi-Square
	2 years	↓25,37	30	5,33	16	34	(3)=75,35
	3 years	↓23,37	30	5,18	15	34	p=0,00001*

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CONCLUSIONS

Our longitudinal follow up data presented the slower progression of the disease at younger age group and in the group with better initial functional findings. The exact mutational analysis has to be considered.

CONTACT