Results.— FSH patients displayed a lower peak torque than controls (−41%). During exercise, deoxygenated hemoglobin (HHb) and blood volume were significantly lower in the FSH patients. The initial muscle deoxygenation and functional impairment (walking endurance) were correlated with the peak torque.

Discussion.— The findings in this study suggest that FSH subjects present an impairment in their capacity to deliver or to use oxygen and would be the consequences of the deconditioning syndrome.

Further reading
http://dx.doi.org/10.1016/j.rehab.2014.03.331

CO04-005-e
Rasch analysis of the motor function measure in patients with congenital muscle dystrophy and congenital myopathy

C. Vuilleurt∗, P. Rippert, V. Kinet∗, A. Renders†, M. Jain‡, M. Waite‡, A. Glanzman†, F. Girardot*, D. Hamrouni*, J. Iwas*, S. Quijano-Roy**, C. Berard**, I. Poirot†, C. Bonnemann‡

∗Service L’escale, Hôpital Femme-Mère–Enfant Aile Al, Bron, France
‡Hospices Civils de Lyon, Pôle Information Médicale, Évaluation, Recherche, 69003 Lyon, France
§Cliniques Universitaires Saint-Luc, Centre de Référence des maladies neuromusculaires, Université Catholique de Louvain, 1200 Bruxelles, Belgium
¶Clinical Research Center, National Institutes of Health, Bethesda, MD 20814, USA
†The Children’s Hospital of Philadelphia, Physical Therapy Department, Philadelphia, PA 19104-4399, USA
‡CHU de Montpellier, Hôpital Arnaud-de-Villeneuve, 34000 Montpellier, France
§CNRS UMR 5558, Laboratoire de Biométrie et Biologie Évolutive, Équipe Biostatistique Santé, 69310 Pierre-Bénite, France
¶Hôpital Raymond-Poincarre, Garches, France
**National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, MD 20814, USA

∗Corresponding author.

Keywords: Dystrophie musculaire congénitale; Myopathie congénitale; Évaluation de la fonction motrice; Mesure des résultats; Analyse Rasch

Objectives.— Valid outcome measures are necessary to monitor the treatment effects in patients with congenital disorders of muscle.

Methods.— In 19 departments in France, Belgium, and USA, 289 patients aged 5- to 77-years-old were enrolled. A Rasch analysis examined the robustness of the motor function measure across the disease spectrum. The three domains (standing position and transfers, axial and proximal motor function, and distal motor function) were examined using RUMM 2030 software with a partial credit model.

Results.— The original 32-item MFM did not fit the Rasch model expectations enough in neither of its domains. Switching from a four- to a three-category response-scale in 18 items restored response order in 16. Various additional checks suggested the removal of seven items. The resulting 25-item MFM demonstrated a good fit to the Rasch model. Domain 1 was well-targeted to the whole severity spectrum whereas domains 2 and 3 were better targeted to severe cases. The reliability coefficients MFM-25 suggested sufficient ability for each summed score to distinguish between patient groups (0.9, 0.8, and 0.7 for domains 1, 2, and 3, respectively).

Discussion.— The Rasch-scaled MFM-25 can be assumed to be a linear scale in each of its three domains.

http://dx.doi.org/10.1016/j.rehab.2014.03.332