

- Baiardini, I., Minetti, C., Bonifacino, S., Porcu, A., Klersy, C., Petralia, P., Balestracci, S., Tarchino, F., Parodi, S., Canonica, G. W., & Braido, F. (2011). Quality of life in Duchenne muscular dystrophy: The subjective impact on children and parents. *Journal of Child Neurology*, 26(6), 707-713. doi:10.1177/0883073810389043
- Chen, J. (2008). Mediators affecting family function in families of children with Duchenne muscular dystrophy. *The Kaohsiung Journal of Medical Sciences*, 24(10), 514-522. doi:10.1016/S1607-551X(09)70010-5
- Chen, J., & Clark, M. (2010). Family resources and parental health in families of children with Duchenne muscular dystrophy. *Journal of Nursing Research*, 18(4), 239-248. doi:10.1097/JNR.0b013e3181fbe37b
- Mah, J. K., Thannhauser, J. E., Kolski, H., & Dewey, D. (2008). Parental stress and quality of life in children with neuromuscular disease. *Pediatric Neurology*, 39(2), 102-107. doi:10.1016/j.pediatrneurol.2008.04.011
- Parsons, E. P., Clarke, A. J., Hood, K., Lycett, E., & Bradley, D. M. (2002). Newborn screening for Duchenne muscular dystrophy: A psychosocial study. *Archives of Disease in Childhood-Fetal and Neonatal Edition*, 86(2), F91-F95. doi:10.1136/fn.86.2.F91
- Plumridge, G., Metcalfe, A., Coad, J., & Gill, P. (2010). Family communication about genetic risk information: Particular issues for Duchenne muscular dystrophy. *American Journal Medical Genetic*, 152A(5), 1225–1232. doi:10.1002/ajmg.a.33364
- Read, J., Kinali, M., Muntoni, F., & Garralda, M. E. (2010). Psychosocial adjustment in siblings of young people with Duschenne muscular dystrophy. *European Journal of Paediatric Neurology*, 14(4), 340-348. doi:10.1016/j.ejpn.2009.09.011
- Reid, D. T., & Renwick, R. M. (2001). Relating familial stress to the psychosocial adjustment of adolescents with Duchenne muscular dystrophy. *International Journal of Rehabilitation Research*, 24(2), 83-93. doi:10.1097/00004356-200106000-00001
- Samson, A., Tomiak, E., Dimillo, J., Lavigne, R., Miles, S., Choquette, M., Chakraborty, P., & Jacob, P. (2009). The lived experience of hope among parents of a child with Duchenne muscular dystrophy: Perceiving the human being beyond the illness. *Chronic Illness*, 5(2), 103-114. doi:10.1177/1742395309104343
- Tomiak, E. M., Samson, A., Miles, S. A., Choquette, M. C., Chakraborty, P. K., & Jacob, P. J. (2007). Gender-specific differences in the psychosocial adjustment of parents of a child with duchenne muscular dystrophy (DMD): Two points of view for a shared experience. *Qualitative Research Journal*, 7(2), 2-21. doi: 110.3316/QRJ0702002
- Yilmaz, O., Yildirim, S. A., Oksuz, C., Atay, S., & Turan, E. (2010). Mothers' depression and health-related quality of life in neuromuscular disease: Role of functional independence level of the children. *Pediatrics International*, 52(4), 648-652. doi:10.1111/j.1442-200X.2010.03094.x