Publications for Michael Nafisinia

2017

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Nafisinia, M., Sobreira, N., Riley, L., Gold, W., Uhlenberg, B., Weib, C., Boehm, C., Prelog, K., Ouvrier, R., Christodoulou, J. (2017). Mutations in RARS cause a hypomyelination disorder akin to Pelizaeus-Merzbacher disease. *European Journal of Human Genetics*, 25(10), 1134-1141. [More Information]

Nafisinia, M., Guo, Y., Dang, X., Li, J., Chen, Y., Zhang, J., Lake, N., Gold, W., Riley, L., Thornburn, D., Christodoulou, J., et al (2017). Whole Exome Sequencing Identifies the Genetic Basis of Late-Onset Leigh Syndrome in a Patient with MRI but Little Biochemical Evidence of a Mitochondrial Disorder. *JIMD Reports*, 32, 117-124. [More Information]

2013

Wolff, J., Nafisinia, M., Sutovsky, P., Ballard, J. (2013). Paternal transmission of mitochondrial DNA as an integral part of mitochondrial inheritance in metapopulations of Drosophila simulans. *Heredity*, 110(1), 57-62. [More Information]