Publications for Megan Brewer

2016

Drew, A., Cutrupi, A., Brewer, M., Nicholson, G., Kennerson, M. (2016). A 1.35Aï¿½Mb DNA fragment is inserted into the DHMN1 locus on chromosome 7q34ï¿½q36.2. *Human Genetics*, 135(11), 1269-1278. [More Information]

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Drew, A., Zhu, D., Kidambi, A., Ly, C., Tey, S., Brewer, M., Ahmad-Annuar, A., Nicholson, G., Kennerson, M. (2015). Improved inherited peripheral neuropathy genetic diagnosis by whole-exome sequencing. *Molecular Genetics & Genomic Medicine*, 3(2), 143-154. [More Information]

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Chaudhry, R., Kidambi, A., Brewer, M., Antonellis, A., Mathews, K., Nicholson, G., Kennerson, M. (2013). Re-analysis of an original CMTX3 family using exome sequencing identifies a known BSCL2 mutation. *Muscle and Nerve*, 47(6), 922-924. [More Information]

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Nicholson, G., Kennerson, M., Brewer, M., Garbern, J., Shy, M. (2009). Genotypes & Sensory Phenotypes in 2 New X-Linked Neuropathies (CMTX3 and dSMAX) and Dominant CMT/HMN Overlap Syndromes. In Carmen Espinos, Vicente Felipo, Francesc Palau (Eds.), *Inherited Neuromuscular Diseases*, (pp. 201-206). Dordrecht: Springer Science+Business Media. [More Information]