## Helen Stewart

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/12022190/publications.pdf

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#	Article	IF	CITATIONS
1	A Transgenic Rat Model of Charcot-Marie-Tooth Disease. Neuron, 1996, 16, 1049-1060.	8.1	346
2	Mutations in NOTCH2 cause Hajdu-Cheney syndrome, a disorder of severe and progressive bone loss. Nature Genetics, 2011, 43, 303-305.	21.4	291
3	Serpentine fibula polycystic kidney syndrome is part of the phenotypic spectrum of Hajdu–Cheney syndrome. European Journal of Human Genetics, 2012, 20, 122-124.	2.8	60
4	A large-scale mutation search reveals genetic heterogeneity in 3M syndrome. European Journal of Human Genetics, 2009, 17, 395-400.	2.8	48
5	<i>OBSL1</i> mutations in 3-M syndrome are associated with a modulation of <i>IGFBP2</i> and <i>IGFBP5</i> expression levels. Human Mutation, 2010, 31, 20-26.	2.5	34
6	De Novo Mutations in DENR Disrupt Neuronal Development and Link Congenital Neurological Disorders to Faulty mRNA Translation Re-initiation. Cell Reports, 2016, 15, 2251-2265.	6.4	30
7	The "CMT Rat― Peripheral Neuropathy and Dysmyelination Caused by Transgenic Overexpression of PMP22. Annals of the New York Academy of Sciences, 1999, 883, 254-261.	3.8	20