

Helen Stewart

List of Publications by Year in descending order

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7
papers

829
citations

1307594

7
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

1300
citing authors

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