Anne-Sophie Lebre

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Novel <i>NDUFS4</i> gene mutation in an atypical late-onset mitochondrial form of multifocal dystonia. Neurology: Genetics, 2017, 3, e205.	0.9	7
2	Quantitative multiplex PCR of short fluorescent fragments for the detection of large intragenic POLG rearrangements in a large French cohort. European Journal of Human Genetics, 2014, 22, 542-550.	1.4	27
3	Amyloid precursor-like protein 2 cleavage contributes to neuronal intranuclear inclusions and cytotoxicity in spinocerebellar ataxia-7 (SCA7). Neurobiology of Disease, 2011, 41, 33-42.	2.1	6
4	NDUFS4 mutations cause Leigh syndrome with predominant brainstem involvement. Molecular Genetics and Metabolism, 2009, 97, 185-189.	0.5	54
5	Amyotrophic lateral sclerosis with neuronal intranuclear protein inclusions. Acta Neuropathologica, 2004, 108, 81-87.	3.9	49
6	PML nuclear bodies and neuronal intranuclear inclusion in polyglutamine diseases. Neurobiology of Disease, 2003, 13, 230-237.	2.1	57
7	Spinocerebellar Ataxia 7 (SCA7). , 2003, , 85-94.		0
8	Two populations of neuronal intranuclear inclusions in SCA7 differ in size and promyelocytic leukaemia protein content. Brain, 2002, 125, 1534-1543.	3.7	61
9	Molecular and Clinical Correlations in Spinocerebellar Ataxia 2: A Study of 32 Families. Human Molecular Genetics, 1997, 6, 709-715.	1.4	270