

# Mutations of optineurin in amyotrophic lateral sclerosis

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Identifying HLA supertypes by learning distance functions. <i>Bioinformatics</i> , 2007, 23, e148-e155.	1.8	45
2	Amyotrophic lateral sclerosis: current practice and future treatments. <i>Current Opinion in Neurology</i> , 2010, 23, 524-529.	1.8	20
3	Current World Literature. <i>Current Opinion in Neurology</i> , 2010, 23, 541-552.	1.8	0
4	Discovering the connection between familial and sporadic amyotrophic lateral sclerosis: pathology trumps genetics. <i>Future Neurology</i> , 2010, 5, 625-628.	0.9	2
5	Towards a unifying, systems biology understanding of large-scale cellular death and destruction caused by poorly liganded iron: Parkinsonâ€™s, Huntingtonâ€™s, Alzheimerâ€™s, prions, bactericides, chemical toxicology and others as examples. <i>Archives of Toxicology</i> , 2010, 84, 825-889.	1.9	330
6	Optineurin, a multifunctional protein involved in glaucoma, amyotrophic lateral sclerosis and antiviral signalling. <i>Journal of Biosciences</i> , 2010, 35, 501-505.	0.5	12
7	TDP-43 and FUS in amyotrophic lateral sclerosis and frontotemporal dementia. <i>Lancet Neurology</i> , The, 2010, 9, 995-1007.	4.9	816
8	A quantitatively-modeled homozygosity mapping algorithm, qHomozygosityMapping, utilizing whole genome single nucleotide polymorphism genotyping data. <i>BMC Bioinformatics</i> , 2010, 11, S5.	1.2	9
9	Quantification of cystatin C in cerebrospinal fluid from various neurological disorders and correlation with G73A polymorphism in CST3. <i>Brain Research</i> , 2010, 1361, 140-145.	1.1	26
10	Characterization of the Properties of a Novel Mutation in VAPB in Familial Amyotrophic Lateral Sclerosis. <i>Journal of Biological Chemistry</i> , 2010, 285, 40266-40281.	1.6	136
11	Systemic transplantation of c-kit+ cells exerts a therapeutic effect in a model of amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2010, 19, 3782-3796.	1.4	66
12	Does Huntingtin play a role in selective macroautophagy?. <i>Cell Cycle</i> , 2010, 9, 3401-3413.	1.3	68
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14	Optic nerve geneticsâ€™ more than meets the eye. <i>Nature Reviews Neurology</i> , 2010, 6, 357-358.	4.9	7
15	Segmental copy-number gain within the region of isopenentenyl diphosphate isomerase genes in sporadic amyotrophic lateral sclerosis. <i>Biochemical and Biophysical Research Communications</i> , 2010, 402, 438-442.	1.0	17
16	Granins as disease-biomarkers: translational potential for psychiatric and neurological disorders. <i>Neuroscience</i> , 2010, 170, 289-297.	1.1	55
17	Mutations in UBQLN2 cause dominant X-linked juvenile and adult-onset ALS and ALS/dementia. <i>Nature</i> , 2011, 477, 211-215.	13.7	1,016
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19	Brainstem and spinal cord motor neuron involvement with optineurin inclusions in proximal-dominant hereditary motor and sensory neuropathy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2011, 82, 1402-1403.	0.9	17
20	Dysregulation of axonal transport and motorneuron diseases. <i>Biology of the Cell</i> , 2011, 103, 87-107.	0.7	29
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73	Genetics of Adult Glaucoma. <i>International Ophthalmology Clinics</i> , 2011, 51, 37-51.	0.3	6
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