Rossella Spataro

List of Publications by Year
in descending order

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50	2,918	304602 22	214721 47
papers	citations	h-index	g-index

50	50	50	4600
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Brainâ€“Computer Interfaces in Acute and Subacute Disorders of Consciousness. Journal of Clinical Neurophysiology, 2022, 39, 32-39.	0.9	9
2	A novel compound heterozygous mutation in GALC associated with adult-onset Krabbe disease: case report and literature review. Neurogenetics, 2022, , 1.	0.7	3
3	Reversible radiculomyelitis after ChAdOx1 nCoV-19 vaccination. BMJ Case Reports, 2022, 15, e247472.	0.2	4
4	Genetic investigation of amyotrophic lateral sclerosis patients in south Italy: a two-decade analysis. Neurobiology of Aging, 2021, 99, 99.e7-99.e14.	1.5	14
5	The capacity to consent to treatment in amyotrophic lateral sclerosis: a preliminary report. Journal of Neurology, 2021, 268, 219-226.	1.8	3
6	Tau protein as a diagnostic and prognostic biomarker in amyotrophic lateral sclerosis. European Journal of Neurology, 2021, 28, 1868-1875.	1.7	19
7	The unfolded protein response in amyotrophic later sclerosis: results of a phase 2 trial. Brain, 2021, 144, 2635-2647.	3.7	33
8	Genomic Portrait of a Sporadic Amyotrophic Lateral Sclerosis Case in a Large Spinocerebellar Ataxia Type 1 Family. Journal of Personalized Medicine, 2020, 10, 262.	1.1	3
9	Effects of a Vibro-Tactile P300 Based Brain-Computer Interface on the Coma Recovery Scale-Revised in Patients With Disorders of Consciousness. Frontiers in Neuroscience, 2020, 14, 294.	1.4	15
10	Effects of Repeating a Tactile Brain-Computer Interface on Patients with Disorder of Consciousness: A Hint of Recovery?*. , 2019, , .		5
11	A novel S379A TARDBP mutation associated to late-onset sporadic ALS. Neurological Sciences, 2019, 40, 2111-2118.	0.9	6
12	Shared polygenic risk and causal inferences in amyotrophic lateral sclerosis. Annals of Neurology, 2019, 85, 470-481.	2.8	118
13	Early and rapidly progressing respiratory failure in a patient with amyotrophic lateral sclerosis: when FVC% is misleading. Neurological Sciences, 2019, 40, 421-422.	0.9	2
14	CSF neurofilament proteins as diagnostic and prognostic biomarkers for amyotrophic lateral sclerosis. Journal of Neurology, 2018, 265, 510-521.	1.8	71
15	Preserved somatosensory discrimination predicts consciousness recovery in unresponsive wakefulness syndrome. Clinical Neurophysiology, 2018, 129, 1130-1136.	0.7	27
16	A Humanâ€“Humanoid Interaction Through the Use of BCI for Locked-In ALS Patients Using Neuro-Biological Feedback Fusion. IEEE Transactions on Neural Systems and Rehabilitation Engineering, 2018, 26, 487-497.	2.7	25
17	Assessment and Communication with Vibro-Tactile P300 And Motor Imagery Bcis in DOC and (C)LIS Patients. Archives of Physical Medicine and Rehabilitation, 2018, 99, e36.	0.5	0
18	The Primitive Palmomental Reflex in Amyotrophic Lateral Sclerosis. European Neurology, 2018, 79, 187-191.	0.6	2

#	Article	IF	Citations
19	Marital status is a prognostic factor in amyotrophic lateral sclerosis. Acta Neurologica Scandinavica, 2017, 136, 624-630.	1.0	7
20	ALS-Related Mutant FUS Protein Is Mislocalized to Cytoplasm and Is Recruited into Stress Granules of Fibroblasts from Asymptomatic <i>FUS </i>P525L Mutation Carriers. Neurodegenerative Diseases, 2017, 17, 292-303.	0.8	23
21	Complete Locked-in and Locked-in Patients: Command Following Assessment and Communication with Vibro-Tactile P300 and Motor Imagery Brain-Computer Interface Tools. Frontiers in Neuroscience, 2017, 11, 251.	1.4	90
22	Intraspinal stem cell transplantation for amyotrophic lateral sclerosis: Ready for efficacy clinical trials?. Cytotherapy, 2016, 18, 1471-1475.	0.3	21
23			

#	Article	IF	Citations
37	Ataxin-1 and ataxin-2 intermediate-length PolyQ expansions in amyotrophic lateral sclerosis. Neurology, 2012, 79, 2315-2320.	1.5	70
38	FUS mutations in sporadic amyotrophic lateral sclerosis: Clinical and genetic analysis. Neurobiology of Aging, 2012, 33, 837.e1-837.e5.	1.5	32
39	C9ORF72 hexanucleotide repeat expansions in the Italian sporadic ALS population. Neurobiology of Aging, 2012, 33, 1848.e15-1848.e20.	1.5	76
40	Clinical characteristics of patients with familial amyotrophic lateral sclerosis carrying the pathogenic GGGGCC hexanucleotide repeat expansion of C9ORF72. Brain, 2012, 135, 784-793.	3.7	182
41	Factors affecting the diagnostic delay in amyotrophic lateral sclerosis. Clinical Neurology and Neurosurgery, 2012, 114, 550-554.	0.6	101
42	Tracheostomy mechanical ventilation in patients with amyotrophic lateral sclerosis: Clinical features and survival analysis. Journal of the Neurological Sciences, 2012, 323, 66-70.	0.3	63
43	Percutaneous endoscopic gastrostomy in amyotrophic lateral sclerosis: Effect on survival. Journal of the Neurological Sciences, 2011, 304, 44-48.	0.3	100
44	FUS mutations in sporadic amyotrophic lateral sclerosis. Neurobiology of Aging, 2011, 32, 550.e1-550.e4.	1.5	79
45	Exome Sequencing Reveals VCP Mutations as a Cause of Familial ALS. Neuron, 2011, 69, 397.	3.8	7
46	Large Proportion of Amyotrophic Lateral Sclerosis Cases in Sardinia Due to a Single Founder Mutation of the TARDBP Gene. Archives of Neurology, 2011, 68, 594.	4.9	104
47	Sleep-wake disturbances in patients with amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 839-842.	0.9	80
48	Causes and place of death in Italian patients with amyotrophic lateral sclerosis. Acta Neurologica Scandinavica, 2010, 122, 217-223.	1.0	80
49	Exome Sequencing Reveals VCP Mutations as a Cause of Familial ALS. Neuron, 2010, 68, 857-864.	3.8	1,100
50	Two Italian kindreds with familial amyotrophic lateral sclerosis due to FUS mutation. Neurobiology of Aging, 2009, 30, 1272-1275.	1.5	128