Ricardo Taipa

List of Publications by Year
in descending order

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95 papers	2,466 citations	249298 26 h-index	252626 46 g-index
97 all docs	97 docs citations	97 times ranked	5080 citing authors

#	Article	IF	Citations
1	Peripheral neuropathy in Parkinsonâ€™s disease: prevalence and functional impact on gait and balance. Brain, 2023, 146, 225-236.	3.7	11
2	T2-FLAIR mismatch sign: a roadmap of pearls and pitfalls. British Journal of Radiology, 2022, 95, 20210825.	1.0	7
3	Cognitive composites for genetic frontotemporal dementia: GENFI-Cog. Alzheimer's Research and Therapy, 2022, 14, 10.	3.0	4
4	PD-L1 tumor expression is associated with poor prognosis and systemic immunosuppression in glioblastoma. Journal of Neuro-Oncology, 2022, 156, 453-464.	1.4	2
5	Examining empathy deficits across familial forms of frontotemporal dementia within the GENFI cohort. Cortex, 2022, 150, 12-28.	1.1	2
6	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	1.5	1
7	Parkinsonâ€™s disease and multiple system atrophy patient iPSC-derived oligodendrocytes exhibit alpha-synucleinâ€“induced changes in maturation and immune reactive properties. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, e2111405119.	3.3	22
8	Cadherinâ€3 is a novel oncogenic biomarker with prognostic value in glioblastoma. Molecular Oncology, 2022, 16, 2611-2631.	2.1	4
9	Brain functional network integrity sustains cognitive function despite atrophy in presymptomatic genetic frontotemporal dementia. Alzheimer's and Dementia, 2021, 17, 500-514.	0.4	36
10	Adult polyglucosan body diseaseâ€”an atypical compound heterozygous with a novel GBE1 mutation. Neurological Sciences, 2021, 42, 2955-2959.	0.9	2
11	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	2.8	42
12	Advantages of an Automated Method Compared With Manual Methods for the Quantification of Intraepidermal Nerve Fiber in Skin Biopsy. Journal of Neuropathology and Experimental Neurology, 2021, 80, 685-694.	0.9	0
13	Association Between Iron-Related Protein Lipocalin 2 and Cognitive Impairment in Cerebrospinal Fluid and Serum. Frontiers in Aging Neuroscience, 2021, 13, 663837.	1.7	3
14	Cadherin Expression and EMT: A Focus on Gliomas. Biomedicines, 2021, 9, 1328.	1.4	30
15	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	1.4	28
16	Disease-related cortical thinning in presymptomatic granulin mutation carriers. NeuroImage: Clinical, 2021, 29, 102540.	1.4	8
17	CNS Involvement in Hereditary Transthyretin Amyloidosis. Neurology, 2021, 97, 1111-1119.	1.5	30
18	Letter to the Editor on â€œCopathology in Progressive Supranuclear Palsy: Does It Matter?â€• Movement Disorders, 2020, 35, 2124-2126.	2.2	4

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19	McArdle's disease: Diagnostic approach after clinical symptoms of vigorous exercise intolerance in a snowboarder in Alpes. Revista Colombiana De ReumatologÃa, 2020, 27, 65-67.	0.0	0
20	Diagnosis of Aicardiâ€ŒoutiÃ¨res Syndrome in Adults: A Case Series. Movement Disorders Clinical Practice, 2020, 7, 303-307.	0.8	13
21	Social cognition impairment in genetic frontotemporal dementia within the GENFI cohort. Cortex, 2020, 133, 384-398.	1.1	26
22	Immune-mediated necrotizing myopathy associated with antibodies to the signal recognition particle: A rare cause of hyperCKaemia. European Journal of Rheumatology, 2020, 7, 143-144.	1.3	0
23	Peripheral neuropathy in systemic vasculitis and other autoimmune diseases â€“ a report of five cases emphasizing the importance of etiologic characterization. ENeurologicalSci, 2020, 21, 100272.	0.5	3
24	P.169Sarcoglycanopathies: experience of a tertiary centre. Neuromuscular Disorders, 2019, 29, S97-S98.	0.3	0
25	Serum neurofilament light chain in genetic frontotemporal dementia: a longitudinal, multicentre cohort study. Lancet Neurology, The, 2019, 18, 1103-1111.	4.9	128
26	Brain biopsy in suspected non-neoplastic neurological disease. Acta Neurochirurgica, 2019, 161, 1139-1147.	0.9	9
27	Structural and molecular correlates of cognitive aging in the rat. Scientific Reports, 2019, 9, 2005.	1.6	31
28	Clinicopathological correlations of sural nerve biopsies in TTR Val30Met familial amyloid polyneuropathy. Brain Communications, 2019, 1, fcz032.	1.5	16
29	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	1.4	27
30	Clinical features of hypertrophic pachymeningitis in a center survey. Neurological Sciences, 2019, 40, 543-551.	0.9	30
31	Functional network resilience to pathology in presymptomatic genetic frontotemporal dementia. Neurobiology of Aging, 2019, 77, 169-177.	1.5	47
32	Proinflammatory and anti-inflammatory cytokines in the CSF of patients with Alzheimer's disease and their correlation with cognitive decline. Neurobiology of Aging, 2019, 76, 125-132.	1.5	121
33	Evidence of amyloid-Î² cerebral amyloid angiopathy transmission through neurosurgery. Acta Neuropathologica, 2018, 135, 671-679.	3.9	80
34	When Decrease AÎ²1-42 in CSF May Not Mean Alzheimer's Disease. Alzheimer Disease and Associated Disorders, 2018, 32, 359-363.	0.6	0
35	Supratentorial hemangioblastomas in von Hippelâ€“Lindau wild-type patientsÂâ€“Âcase series and literature review. International Journal of Neuroscience, 2018, 128, 295-303.	0.8	13
36	Inflammatory pathology markers (activated microglia and reactive astrocytes) in early and late onset Alzheimer disease: a post mortem study. Neuropathology and Applied Neurobiology, 2018, 44, 298-313.	1.8	55

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37	CONGENITAL MYOPATHIES: NEMALINE AND TITINOPATHIES. Neuromuscular Disorders, 2018, 28, S100.	0.3	0
38	Lateâ€onset Levodopa Responsive Parkinsonism Due to Polymerase Î³ 1 Mutations. Movement Disorders Clinical Practice, 2018, 5, 645-648.	0.8	0
39	WNT6 is a novel oncogenic prognostic biomarker in human glioblastoma. Theranostics, 2018, 8, 4805-4823.	4.6	35
40	LAMA2gene mutation update: Toward a more comprehensive picture of the laminin-Î±2 variome and its related phenotypes. Human Mutation, 2018, 39, 1314-1337.	1.1	71
41	The etiology of spontaneous intracerebralhemorrhage: Insights from a neuropathological series. , 2018, 37, 16-21.		1
42	Clinical, biochemical, molecular, and histological features of 65 Portuguese patients with mitochondrial disorders. Muscle and Nerve, 2017, 56, 868-872.	1.0	6
43	Frontotemporal lobar degenerationâ€TDP with â€˜multiple system atrophy phenocopy syndromeâ€™. Neuropathology and Applied Neurobiology, 2017, 43, 533-536.	1.8	4
44	RYR1-Related Myopathies: Clinical, Histopathologic and Genetic Heterogeneity Among 17 Patients from a Portuguese Tertiary Centre. Journal of Neuromuscular Diseases, 2017, 4, 67-76.	1.1	10
45	Reply. Muscle and Nerve, 2017, 56, E49.	1.0	0
46	Inflammatory myopathy associated with myasthenia gravis with and without thymic pathology: Report of four cases and literature review. Autoimmunity Reviews, 2017, 16, 644-649.	2.5	28
47	Amyloid detection in the transverse carpal ligament of patients with hereditary ATTR V30M amyloidosis and carpal tunnel syndrome. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 73-77.	1.4	21
48	Patterns of Microglial Cell Activation in Alzheimer Disease and Frontotemporal Lobar Degeneration. Neurodegenerative Diseases, 2017, 17, 145-154.	0.8	32
49	Vascular Pathology Causing Late Onset Generalized Chorea: A Clinicoâ€Pathological Case Report. Movement Disorders Clinical Practice, 2017, 4, 819-823.	0.8	2
50	Late-onset limb-girdle myopathy with oculobulbar signs and rimmed vacuoles associated with a novel Pompe disease mutation. Neuromuscular Disorders, 2017, 27, S163.	0.3	0
51	Exonization of an Intronic LINE-1 Element Causing Becker Muscular Dystrophy as a Novel Mutational Mechanism in Dystrophin Gene. Genes, 2017, 8, 253.	1.0	25
52	Does the Interplay Between Aging and Neuroinflammation Modulate Alzheimerâ€™s Disease Clinical Phenotypes? A Clinico-Pathological Perspective. Journal of Alzheimer's Disease, 2016, 53, 403-417.	1.2	12
53	Nonprimary Cytomegalovirus Fetal Infection. Revista Brasileira De Ginecologia E Obstetricia, 2016, 38, 196-200.	0.3	1
54	DJ-1 linked parkinsonism (PARK7) is associated with Lewy body pathology. Brain, 2016, 139, 1680-1687.	3.7	89

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55	Duchenne muscular dystrophy: Clinical, genetic and pathological changes in preclinical and early stages. Neuromuscular Disorders, 2016, 26, S124.	0.3	0
56	Regulation of WNT6 by HOXA9 in glioblastoma: functional and clinical relevance. European Journal of Cancer, 2016, 61, S45-S46.	1.3	1
57	Ryanodine-related myopathies: Clinical, histopathologic and genetic heterogeneity among 16 patients from a Portuguese tertiary centre. Neuromuscular Disorders, 2016, 26, S135-S136.	0.3	0
58	The expanding phenotype of LAMA2-related muscular dystrophies: Four additional cases diagnosed during adulthood. Neuromuscular Disorders, 2016, 26, S190.	0.3	1
59	Post-mortem assessment in vascular dementia: advances and aspirations. BMC Medicine, 2016, 14, 129.	2.3	99
60	CADASIL: MRI may be normal in the fourth decade of life â€“ a case report. Cephalalgia, 2016, 36, 1082-1085.	1.8	8
61	New massive parallel sequencing approach improves the genetic characterization of congenital myopathies. Journal of Human Genetics, 2016, 61, 497-505.	1.1	15
62	Appendectomy may delay <scp>P</scp>arkinson's disease Onset. Movement Disorders, 2015, 30, 1404-1407.	2.2	46
63	Neuropatia da DoenÃ§a de Hansen: Um DiagnÃ³stico a Considerar na InvestigaÃ§Ã£o da Neuropatia PerifÃ©rica. Acta Medica Portuguesa, 2015, 28, 329.	0.2	0
64	New splicing mutation in the choline kinase beta (CHKB) gene causing a muscular dystrophy detected by whole-exome sequencing. Journal of Human Genetics, 2015, 60, 305-312.	1.1	33
65	PARK2 presenting as a disabling peripheral axonal neuropathy. Neurological Sciences, 2015, 36, 341-343.	0.9	3
66	Chronic Lymphocytic Inflammation With Pontine Perivascular Enhancement Responsive to Steroids (CLIPPERS). Journal of Neuropathology and Experimental Neurology, 2015, 74, 186-190.	0.9	19
67	Chondroma of the Cerebral Falx: Case Report of a Very Rare Intracranial Tumor. Brazilian Neurosurgery, 2015, 34, 144-147.	0.0	1
68	CNS involvement in V30M transthyretin amyloidosis: clinical, neuropathological and biochemical findings. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 159-167.	0.9	97
69	Hansen Neuropathy: Still a Possible Diagnosis in the Investigation of a Peripheral Neuropathy. Acta Medica Portuguesa, 2015, 28, 329-32.	0.2	0
70	Acute Ischemic Stroke Secondary to Glioblastoma. Neuroradiology Journal, 2014, 27, 85-90.	0.6	22
71	Freezing of gait â€“ First motor manifestation in late infantile variant neuronal ceroid lipofuscinosis. Parkinsonism and Related Disorders, 2014, 20, 243-244.	1.1	7
72	Ryanodine Myopathies Without Central Coresâ€”Clinical, Histopathologic, and Genetic Description of Three Cases. Pediatric Neurology, 2014, 51, 275-278.	1.0	9

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73	Multiple cerebral infarcts and intravascular central nervous system lymphoma: A rare but potentially treatable association. Journal of the Neurological Sciences, 2013, 325, 183-185.	0.3	12
74	Novel TTC19 mutation in a family with severe psychiatric manifestations and complex III deficiency. Neurogenetics, 2013, 14, 153-160.	0.7	42
75	Genetic Analysis of Inherited Leukodystrophies. JAMA Neurology, 2013, 70, 875.	4.5	147
76	Expanding the MTM1 mutational spectrum: novel variants including the first multi-exonic duplication and development of a locus-specific database. European Journal of Human Genetics, 2013, 21, 540-549.	1.4	29
77	Hereditary Neuropathy with Liability to Pressure Palsy: A Recurrent and Bilateral Foot Drop Case Report. Case Reports in Pediatrics, 2013, 2013, 1-5.	0.2	2
78	Clinico-Pathological Correlations of the Most Common Neurodegenerative Dementias. Frontiers in Neurology, 2012, 3, 68.	1.1	31
79	Clinical, Neuropathological, and Genetic Characteristics of the Novel IVS9+1delG GRN Mutation in a Patient with Frontotemporal Dementia. Journal of Alzheimer's Disease, 2012, 30, 83-90.	1.2	9
80	Sporadic hemiplegic migraine with normal imaging as the initial manifestation of CADASIL. Cephalalgia, 2012, 32, 255-257.	1.8	8
81	Permanent dysphagia in familial amyloid polyneuropathy (ATTRVal30Met). Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 110-112.	1.4	5
82	TTC7B Emerges as a Novel Risk Factor for Ischemic Stroke Through the Convergence of Several Genome-Wide Approaches. Journal of Cerebral Blood Flow and Metabolism, 2012, 32, 1061-1072.	2.4	86
83	Task-specific contribution of the human striatum to perceptualâ€“motor skill learning. Journal of Clinical and Experimental Neuropsychology, 2011, 33, 51-62.	0.8	19
84	Medulloblastoma and gliomatosis cerebri: rare brain tumors in multiple sclerosis patients. Neurological Sciences, 2011, 32, 893-897.	0.9	7
85	Acute hemorrhagic leukoencephalitis with severe brainstem and spinal cord involvement: MRI features with neuropathological confirmation. Journal of Magnetic Resonance Imaging, 2011, 33, 957-961.	1.9	31
86	Gliomatosis Cerebri Diagnostic Challenge. Neurologist, 2011, 17, 269-272.	0.4	3
87	Primary antiphospholipid antibody syndrome presenting with encephalopathy, psychosis and seizures. Lupus, 2011, 20, 1433-1435.	0.8	7
88	Kalirin: a novel genetic risk factor for ischemic stroke. Human Genetics, 2010, 127, 513-523.	1.8	51
89	Adultâ€onset dystonia in Aicardiâ€GoutiÃ¨res syndrome. Movement Disorders, 2010, 25, 791-793.	2.2	2
90	A case of haemophagocytic syndrome presenting with oculogyric crises. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 469-471.	0.9	1

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91	Mitochondrial haplogroup H1 is protective for ischemic stroke in Portuguese patients. BMC Medical Genetics, 2008, 9, 57.	2.1	42
92	Streptococcus suis meningitis: First case report from Portugal. Journal of Infection, 2008, 56, 482-483.	1.7	11
93	Specific Configuration of Dendritic Degeneration in Pyramidal Neurons of the Medial Prefrontal Cortex Induced by Differing Corticosteroid Regimens. Cerebral Cortex, 2007, 17, 1998-2006.	1.6	146
94	Morphological Correlates of Corticosteroid-Induced Changes in Prefrontal Cortex-Dependent Behaviors. Journal of Neuroscience, 2005, 25, 7792-7800.	1.7	242
95	X-Linked Myotubular Myopathy: A Novel Mutation Expanding the Genotypic Spectrum of a Phenotypically Heterogeneous Myopathy. Journal of Pediatric Genetics, 0, , .	0.3	0