

Stéphane Mathis

List of Publications by Year in descending order

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Version: 2024-02-01

77
papers

1,464
citations

393982

19
h-index

360668

35
g-index

78
all docs

78
docs citations

78
times ranked

2238
citing authors

#	ARTICLE	IF	CITATIONS
1	New classification of autoimmune neuropathies based on target antigens and involved domains of myelinated fibres. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022, 93, 57-67.	0.9	18
2	Neurologic manifestations of giant cell arteritis. <i>Journal of Neurology</i> , 2022, 269, 3430-3442.	1.8	2
3	Peripheral neuropathy and livedoid vasculopathy. <i>Journal of Neurology</i> , 2022, 269, 3779-3788.	1.8	6
4	Chronic Inflammatory or Chronic Inflammatory Demyelinating Polyradiculoneuropathy?. <i>Frontiers in Neurology</i> , 2022, 13, 862335.	1.1	2
5	Widening of myelin lamellae in polyneuropathy with immunoglobulin-M monoclonal gammopathy, without activity against myelin-associated glycoprotein, responsive to treatment. <i>Neuromuscular Disorders</i> , 2022, 32, 678-681.	0.3	2
6	Are Miller Fisher syndrome and CANDAs due to a paranodopathy?. <i>Journal of the Neurological Sciences</i> , 2022, 438, 120279.	0.3	5
7	The Wide Spectrum of Pathophysiologic Mechanisms of Paraproteinemic Neuropathy. <i>Neurology</i> , 2021, 96, 214-225.	1.5	11
8	The ataxic neuropathies. <i>Journal of Neurology</i> , 2021, 268, 3675-3689.	1.8	4
9	Epidemics and outbreaks of peripheral nervous system disorders: I. infectious and immune-mediated causes. <i>Journal of Neurology</i> , 2021, 268, 879-890.	1.8	2
10	Epidemics and outbreaks of peripheral nervous system disorders: II. Toxic and nutritional causes. <i>Journal of Neurology</i> , 2021, 268, 892-902.	1.8	4
11	Impact of Coronavirus Disease 2019 in a French Cohort of Myasthenia Gravis. <i>Neurology</i> , 2021, 96, e2109-e2120.	1.5	38
12	Diagnosis and treatment of CIDP: a "grand cru" of updated data. <i>European Journal of Neurology</i> , 2021, 28, 3545-3546.	1.7	3
13	CIDP and hemopathies, an underestimated association. <i>Journal of the Neurological Sciences</i> , 2021, 429, 118055.	0.3	2
14	Prognostic factor of poor outcome in anti-MAG neuropathy: clinical and electrophysiological analysis of a French Cohort. <i>Journal of Neurology</i> , 2020, 267, 561-571.	1.8	7
15	Ultrastructural Lesions of Nodoparanopathies in Peripheral Neuropathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 247-255.	0.9	21
16	Antibody- and macrophage-mediated segmental demyelination in chronic inflammatory demyelinating polyneuropathy: clinical, electrophysiological, immunological and pathological correlates. <i>European Journal of Neurology</i> , 2020, 27, 692-701.	1.7	25
17	When botany inspired pathology of the peripheral nervous system. <i>Neurology</i> , 2020, 95, 532-536.	1.5	0
18	History of acute polyradiculoneuropathy (part 1). <i>Neurology</i> , 2020, 94, 828-835.	1.5	5

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19	History of acute polyradiculoneuropathy (part 2). <i>Neurology</i> , 2020, 94, 836-840.	1.5	5
20	Minimizing the Diagnostic Delay in Amyotrophic Lateral Sclerosis: The Role of Nonneurologist Practitioners. <i>Neurology Research International</i> , 2020, 2020, 1-8.	0.5	10
21	Peripheral nervous system involvement in vasculitis. , 2020, , 145-176.		4
22	The mysterious death of Georges Cuvier (1832): An early case of severe Guillain-Barré syndrome?. <i>Neuromuscular Disorders</i> , 2020, 30, 250-253.	0.3	0
23	ADAR1 mediated regulation of neural crest derived melanocytes and Schwann cell development. <i>Nature Communications</i> , 2020, 11, 198.	5.8	30
24	Early clinicopathologic description of nodoparanodopathy in the 19th century. <i>Neurology</i> , 2019, 93, 788-792.	1.5	3
25	Myopathy and scleromyxedema. <i>Journal of Neurology</i> , 2019, 266, 2051-2059.	1.8	4
26	The journal behind the nodes of Ranvier?. <i>Lancet Neurology</i> , The, 2019, 18, 628.	4.9	2
27	Characteristics of patients with vitamin B12-responsive neuropathy: a case series with systematic repeated electrophysiological assessment. <i>Neurological Research</i> , 2019, 41, 569-576.	0.6	24
28	Genetics of amyotrophic lateral sclerosis: A review. <i>Journal of the Neurological Sciences</i> , 2019, 399, 217-226.	0.3	182
29	Peripheral nervous system involvement in Leber's hereditary optic neuropathy. <i>Journal of the Neurological Sciences</i> , 2018, 388, 94-96.	0.3	2
30	Updating the classification of inherited neuropathies. <i>Neurology</i> , 2018, 90, e870-e876.	1.5	33
31	Acute Brachial Radiculoplexopathy and Giant Cell Arteritis. <i>Neurologist</i> , 2018, 23, 23-28.	0.4	7
32	Nerve Biopsy Is Still Useful in Some Inherited Neuropathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 88-99.	0.9	16
33	History and current difficulties in classifying inherited myopathies and muscular dystrophies. <i>Journal of the Neurological Sciences</i> , 2018, 384, 50-54.	0.3	14
34	Chronic inflammatory demyelinating polyradiculoneuropathy causing myelopathy. <i>Muscle and Nerve</i> , 2018, 57, E102-E103.	1.0	1
35	Subacute nodopathy with conduction blocks and anti-neurofascin 140/186 antibodies: an ultrastructural study. <i>Brain</i> , 2018, 141, e56-e56.	3.7	47
36	Safety of Intravenous Immunoglobulin (Tegeline®), Administered at Home in Patients with Autoimmune Disease: Results of a French Study. <i>BioMed Research International</i> , 2018, 2018, 1-10.	0.9	3

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37	Value of nerve biopsy in the management of peripheral neuropathies. Expert Review of Neurotherapeutics, 2018, 18, 589-602.	1.4	11
38	RNA-Targeted Therapies and Amyotrophic Lateral Sclerosis. Biomedicines, 2018, 6, 9.	1.4	20
39	The classification of Charcot-Marie-Tooth diseases, a never-ending story: CMT4?. Brain, 2018, 141, e70.	3.7	2
40	Posterior reversible encephalopathy syndrome and reversible cerebral vasoconstriction syndrome after bilateral carotid paraganglioma resection: A case report. Cephalalgia, 2017, 37, 89-93.	1.8	5
41	MRI and surgical lumbosacral trunk positioning palsy. Muscle and Nerve, 2017, 56, E36-E37.	1.0	0
42	Acute tibial neuropathy in an elderly. Journal of Clinical Neuroscience, 2017, 46, 58-59.	0.8	2
43	Did Jules Dejerine describe AMAN at the end of the 19th century?. Neurology, 2017, 89, 1749-1753.	1.5	4
44	Jules Dejerine and the peripheral nervous system. Neurology, 2017, 89, 611-615.	1.5	5
45	Paranodal lesions in chronic inflammatory demyelinating polyneuropathy associated with anti-Neurofascin 155 antibodies. Neuromuscular Disorders, 2017, 27, 290-293.	0.3	88
46	Sensory neuropathy in progressive motor neuronopathy <i>(pmn)</i> mice is associated with defects in microtubule polymerization and axonal transport. Brain Pathology, 2017, 27, 459-471.	2.1	16
47	Management and therapeutic perspectives in amyotrophic lateral sclerosis. Expert Review of Neurotherapeutics, 2017, 17, 263-276.	1.4	29
48	Sensory Neuronopathy Revealing Severe Vitamin B12 Deficiency in a Patient with Anorexia Nervosa: An Often-Forgotten Reversible Cause. Nutrients, 2017, 9, 281.	1.7	16
49	Current view and perspectives in amyotrophic lateral sclerosis. Neural Regeneration Research, 2017, 12, 181.	1.6	26
50	Simultaneous Combined Myositis, Inflammatory Polyneuropathy, and Overlap Myasthenic Syndrome. Case Reports in Neurological Medicine, 2016, 2016, 1-11.	0.3	1
51	Monoclonal gammopathy of undetermined significance and endoneurial IgG deposition. Medicine (United States), 2016, 95, e4807.	0.4	10
52	Congenital hypomyelinating neuropathy due to the association of a truncating mutation in PMP22 with the classical HNPP deletion. Neuromuscular Disorders, 2016, 26, 316-321.	0.3	7
53	Simultaneous Quantification of Unmyelinated Nerve Fibers in Sural Nerve and in Skin. Journal of Neuropathology and Experimental Neurology, 2016, 75, 53-60.	0.9	10
54	Natalizumab throughout pregnancy: Risk of low platelet count in the newborn at delivery. Revue Neurologique, 2016, 172, 165-166.	0.6	14

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55	Guillain-Barré Syndrome (42 Cases) Occurring During a Zika Virus Outbreak in French Polynesia. <i>Medicine (United States)</i> , 2016, 95, e3257.	0.4	92
56	Reasons Charcot-Marie-Tooth disease due to mutations in the <i>MME</i> gene should not be named ARMT2T. <i>Annals of Neurology</i> , 2016, 80, 477-477.	2.8	4
57	Contactin-Associated Protein 1 (<i>CNTNAP1</i>) Mutations Induce Characteristic Lesions of the Paranodal Region. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 1155-1159.	0.9	31
58	Long-term outcome of basilar stenosis in Erdheim-Chester disease. <i>Medicine (United States)</i> , 2016, 95, e4813.	0.4	11
59	Therapeutic options and management of polyneuropathy associated with anti-MAG antibodies. <i>Expert Review of Neurotherapeutics</i> , 2016, 16, 1111-1119.	1.4	10
60	Classifications of neurogenetic diseases: An increasingly complex problem. <i>Revue Neurologique</i> , 2016, 172, 339-349.	0.6	22
61	Intraventricular Silicone Oil. <i>Medicine (United States)</i> , 2016, 95, e2359.	0.4	21
62	Too many numbers and complexity: time to update the classifications of neurogenetic disorders?. <i>Journal of Medical Genetics</i> , 2016, 53, 647-650.	1.5	10
63	Heterogeneity of Polyneuropathy Associated with Anti-MAG Antibodies. <i>Journal of Immunology Research</i> , 2015, 2015, 1-9.	0.9	54
64	Therapeutic options in Charcot-Marie-Tooth diseases. <i>Expert Review of Neurotherapeutics</i> , 2015, 15, 355-366.	1.4	16
65	Charcot-Marie-Tooth diseases: an update and some new proposals for the classification. <i>Journal of Medical Genetics</i> , 2015, 52, 681-690.	1.5	80
66	Multiple simultaneous intracranial hemorrhages due to hornet stings. <i>Clinical Neurology and Neurosurgery</i> , 2015, 128, 53-55.	0.6	4
67	Hereditary motor and sensory neuropathies or Charcot-Marie-Tooth diseases: An update. <i>Journal of the Neurological Sciences</i> , 2014, 347, 14-22.	0.3	69
68	Testing the validity of a set of diagnostic criteria for sensory neuronopathies: a francophone collaborative study. <i>Journal of Neurology</i> , 2014, 261, 2093-2100.	1.8	22
69	Apathy in Parkinson's Disease: An Electrophysiological Study. <i>Neurology Research International</i> , 2014, 2014, 1-9.	0.5	18
70	Paroxysmal Sneezing at the Onset of Syncope and Transient Ischemic Attack Revealing a Papillary Cardiac Fibroelastoma. <i>Case Reports in Neurological Medicine</i> , 2014, 2014, 1-3.	0.3	4
71	Motor neuronopathy in Chediak-Higashi syndrome. <i>Journal of the Neurological Sciences</i> , 2014, 344, 203-207.	0.3	13
72	A Rare Cause of Stroke in Young Adults: Occlusion of the Middle Cerebral Artery by a Meningioma Postpartum. <i>Case Reports in Neurological Medicine</i> , 2013, 2013, 1-4.	0.3	6

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73	Bilateral Femoral Neuropathy After Massive Toxic Ingestion in a Suicide Attempt. <i>Neurologist</i> , 2012, 18, 70-72.	0.4	2
74	Cerebral abscesses in hereditary haemorrhagic telangiectasia: A clinical and microbiological evaluation. <i>Clinical Neurology and Neurosurgery</i> , 2012, 114, 235-240.	0.6	66
75	POEMS syndrome with prominent acute axonal lesions. <i>Journal of the Neurological Sciences</i> , 2012, 313, 185-188.	0.3	9
76	Amyloid neuropathy mimicking chronic inflammatory demyelinating polyneuropathy. <i>Muscle and Nerve</i> , 2012, 45, 26-31.	1.0	74
77	The ovarioleukodystrophy. <i>Clinical Neurology and Neurosurgery</i> , 2008, 110, 1035-1037.	0.6	16