

# Ming J Lim

## List of Publications by Year in descending order

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172  
papers

8,687  
citations

41627

51  
h-index

58552

86  
g-index

203  
all docs

203  
docs citations

203  
times ranked

9499  
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterizing the features and course of psychiatric symptoms in children and adolescents with autoimmune encephalitis. <i>European Archives of Psychiatry and Clinical Neuroscience</i> , 2022, 272, 477-482.	1.8	5
2	Diagnosis and management of multiple sclerosis and other relapsing demyelinating disease in childhood. <i>Archives of Disease in Childhood</i> , 2022, 107, 216-222.	1.0	2
3	Early predictors of disability of paediatric-onset AQP4-IgG-seropositive neuromyelitis optica spectrum disorders. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022, 93, 101-111.	0.9	16
4	Incidence of paediatric multiple sclerosis and other acquired demyelinating syndromes: 10-year follow-up surveillance study. <i>Developmental Medicine and Child Neurology</i> , 2022, 64, 502-508.	1.1	4
5	Neurological and cognitive outcomes after antibody-negative autoimmune encephalitis in children. <i>Developmental Medicine and Child Neurology</i> , 2022, 64, 649-653.	1.1	10
6	Clinical features, investigations, and outcomes of pediatric limbic encephalitis: A multicenter study. <i>Annals of Clinical and Translational Neurology</i> , 2022, 9, 67-78.	1.7	7
7	Patients' and carers' views and the importance of ethnicity, diversity and inclusion in research priority settings. <i>Archives of Disease in Childhood</i> , 2022, 107, 415-415.	1.0	0
8	Diagnosis and Management of Opsoclonus-Myoclonus-Ataxia Syndrome in Children. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2022, 9, .	3.1	26
9	Workshop on RanBP2/Nup358 and acute necrotizing encephalopathy. <i>Nucleus</i> , 2022, 13, 156-171.	0.6	9
10	Pediatric Ischemic Stroke: An Infrequent Complication of SARS-CoV-2. <i>Annals of Neurology</i> , 2021, 89, 657-665.	2.8	74
11	Acute flaccid myelitis: cause, diagnosis, and management. <i>Lancet, The</i> , 2021, 397, 334-346.	6.3	88
12	Systemic Inflammation Is Associated With Neurologic Involvement in Pediatric Inflammatory Multisystem Syndrome Associated With SARS-CoV-2. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	3.1	29
13	Use of Disease-Modifying Therapies in Pediatric Relapsing-Remitting Multiple Sclerosis in the United Kingdom. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	3.1	16
14	Clinical guidance for diagnosis and management of suspected Pediatric Acute-onset Neuropsychiatric Syndrome in the Nordic countries. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 3153-3160.	0.7	6
15	Check your immune privilege: Is there a role for the maternal immune system in the pathogenesis of childhood tics and obsessive-compulsive disorder?. <i>Brain, Behavior, and Immunity</i> , 2021, 95, 19-20.	2.0	0
16	International Consensus Recommendations for the Treatment of Pediatric NMDAR Antibody Encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	3.1	70
17	Vaccination in acute immune-mediated/inflammatory disorders of the central nervous system. <i>European Journal of Paediatric Neurology</i> , 2021, 34, 118-122.	0.7	1
18	Neurological manifestations of SARS-CoV-2 infection in hospitalised children and adolescents in the UK: a prospective national cohort study. <i>The Lancet Child and Adolescent Health</i> , 2021, 5, 631-641.	2.7	114

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19	A recent surge of fulminant and early onset subacute sclerosing panencephalitis (SSPE) in the United Kingdom: An emergence in a time of measles. <i>European Journal of Paediatric Neurology</i> , 2021, 34, 43-49.	0.7	13
20	Use and Safety of Immunotherapeutic Management of <i>N</i> -Methyl-D-Aspartate Receptor Antibody Encephalitis. <i>JAMA Neurology</i> , 2021, 78, 1333.	4.5	91
21	P080 An embedded pathway to mandibular advancement splint (MAS) construction in a tertiary hospital reduces barriers to care for low-income individuals. <i>SLEEP Advances</i> , 2021, 2, A47-A47.	0.1	0
22	Authors' reply regarding "On diagnosing and treating PANS/ PANDAS: questions from a patient support group". <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 3390-3391.	0.7	2
23	Idiopathic Hypertrophic Pachymeningitis: Does Earlier Treatment Improve Outcome?. <i>Children</i> , 2021, 8, 11.	0.6	7
24	Acute Myelopathy in Childhood. <i>Children</i> , 2021, 8, 1055.	0.6	2
25	Early predictors of epilepsy and subsequent relapse in children with acute disseminated encephalomyelitis. <i>Multiple Sclerosis Journal</i> , 2020, 26, 333-342.	1.4	37
26	Improved performance of the 2017 McDonald criteria for diagnosis of multiple sclerosis in children in a real-life cohort. <i>Multiple Sclerosis Journal</i> , 2020, 26, 1372-1380.	1.4	28
27	Radiological Cerebrospinal Posterior Reversible Encephalopathy Syndrome Mimicking Acute Disseminated Encephalomyelitis in a Neurologically Asymptomatic Child. <i>Pediatric Neurology</i> , 2020, 106, 65-67.	1.0	0
28	E.U. paediatric MOG consortium consensus: Part 5 – Treatment of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 41-53.	0.7	59
29	E.U. paediatric MOG consortium consensus: Part 4 – Outcome of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 32-40.	0.7	29
30	E.U. paediatric MOG consortium consensus: Part 1 – Classification of clinical phenotypes of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 2-13.	0.7	87
31	Catatonic features in children and adolescents with <i>N</i> -methyl-D-aspartate receptor antibody encephalitis. <i>BJPsych Open</i> , 2020, 6, .	0.3	5
32	Progress in the Management of Paediatric-Onset Multiple Sclerosis. <i>Children</i> , 2020, 7, 222.	0.6	4
33	Treatment and outcome of aquaporin-4 antibody-positive NMO. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2020, 7, .	3.1	37
34	Evolving Cognitive Dysfunction in Children with Neurologically Stable Opsoclonus-Myoclonus Syndrome. <i>Children</i> , 2020, 7, 103.	0.6	5
35	E.U. paediatric MOG consortium consensus: Part 3 – Biomarkers of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 22-31.	0.7	24
36	Acute Disseminated Encephalomyelitis: Current Perspectives. <i>Children</i> , 2020, 7, 210.	0.6	24

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37	E.U. paediatric MOG consortium consensus: Part 2 – Neuroimaging features of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. <i>European Journal of Paediatric Neurology</i> , 2020, 29, 14-21.	0.7	32
38	Treatment of MOG-IgG-associated disorder with rituximab: An international study of 121 patients. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 44, 102251.	0.9	110
39	Neutrophil-to-lymphocyte ratio correlates with disease activity in myelin oligodendrocyte glycoprotein antibody associated disease (MOGAD) in children. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 45, 102345.	0.9	13
40	Treatment of MOG antibody associated disorders: results of an international survey. <i>Journal of Neurology</i> , 2020, 267, 3565-3577.	1.8	64
41	Genomic Landscape of Reed-Sternberg Cells of Hodgkin Lymphoma from Children, Adolescents, and Young Adults. <i>Klinische Padiatrie</i> , 2020, 232, .	0.2	0
42	The Movement disorder associated with NMDAR antibody-encephalitis is complex and characteristic: an expert video-rating study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 724-726.	0.9	71
43	A framework for measurement and harmonization of pediatric multiple sclerosis etiologic research studies: The Pediatric MS Tool-Kit. <i>Multiple Sclerosis Journal</i> , 2019, 25, 1170-1177.	1.4	3
44	Is chronic lymphocytic inflammation with pontine perivascular enhancement responsive to steroids (CLIPPERS) in children the same condition as in adults?. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 490-496.	1.1	15
45	Ischemic Stroke Following Ergotamine Overdose. <i>Pediatric Neurology</i> , 2019, 101, 81-82.	1.0	4
46	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. <i>JAMA Network Open</i> , 2019, 2, e1914274.	2.8	14
47	Combined Anti-inflammatory and Neuroprotective Treatments Have the Potential to Impact Disease Phenotypes in <i>Cln3</i> <sup>+/Δ</sup> Mice. <i>Frontiers in Neurology</i> , 2019, 10, 963.	1.1	13
48	Utility and safety of plasma exchange in paediatric neuroimmune disorders. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 540-546.	1.1	8
49	Paediatric multiple sclerosis: a new era in diagnosis and treatment. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 1039-1049.	1.1	30
50	Mycophenolate mofetil in paediatric autoimmune or immune-mediated diseases of the central nervous system: clinical experience and recommendations. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 458-468.	1.1	15
51	Magnetic resonance imaging in enterovirus 71, myelin oligodendrocyte glycoprotein antibody, aquaporin 4 antibody, and multiple sclerosis-associated myelitis in children. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 1108-1116.	1.1	22
52	Childhood disintegrative disorder and autism spectrum disorder: a systematic review. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 523-534.	1.1	18
53	An increase in reports of acute flaccid paralysis (AFP) in the United Kingdom, 1 January 2018–21 January 2019: early findings. <i>Eurosurveillance</i> , 2019, 24, .	3.9	31
54	Myelin oligodendrocyte glycoprotein and aquaporin 4 antibodies are highly specific in children with acquired demyelinating syndromes. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 958-962.	1.1	105

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55	Retinal nerve fibre layer thinning is associated with worse visual outcome after optic neuritis in children with a relapsing demyelinating syndrome. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 1244-1250.	1.1	38
56	Paediatric acute disseminated encephalomyelitis followed by optic neuritis: disease course, treatment response and outcome. <i>European Journal of Neurology</i> , 2018, 25, 782-786.	1.7	45
57	“Leukodystrophy-like” phenotype in children with myelin oligodendrocyte glycoprotein antibody-associated disease. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 417-423.	1.1	81
58	Disease Course and Treatment Responses in Children With Relapsing Myelin Oligodendrocyte Glycoprotein Antibody-Associated Disease. <i>JAMA Neurology</i> , 2018, 75, 478.	4.5	306
59	Immunotherapy-responsive childhood neurodegeneration with systemic and central nervous system inflammation. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 882-888.	0.7	1
60	Pseudotumor cerebri syndrome in a patient with narcolepsy type 1. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 194-198.	0.7	1
61	Endocrinopathies in paediatric-onset neuromyelitis optica spectrum disorder with aquaporin 4 (AQP4) antibody. <i>Multiple Sclerosis Journal</i> , 2018, 24, 679-684.	1.4	9
62	NMDA-receptor antibodies alter cortical microcircuit dynamics. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E9916-E9925.	3.3	39
63	Testing combinatorial therapies for juvenile Batten disease. <i>Molecular Genetics and Metabolism</i> , 2018, 123, S33.	0.5	1
64	Glutamate receptor $\gamma 2$ serum antibodies in pediatric opsoclonus myoclonus ataxia syndrome. <i>Neurology</i> , 2018, 91, e714-e723.	1.5	43
65	The role of inflammation and hypovitaminosis-D in multiple sclerosis, schizophrenia and autism. <i>Neurology Psychiatry and Brain Research</i> , 2018, 29, 14-15.	2.0	0
66	Therapeutic plasma exchange in paediatric neurology: a critical review and proposed treatment algorithm. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 765-779.	1.1	24
67	Systematic review of immunoglobulin use in paediatric neurological and neurodevelopmental disorders. <i>Developmental Medicine and Child Neurology</i> , 2017, 59, 136-144.	1.1	24
68	Diagnostic algorithm for relapsing acquired demyelinating syndromes in children. <i>Neurology</i> , 2017, 89, 269-278.	1.5	155
69	Autoimmune encephalitis in children: clinical phenomenology, therapeutics, and emerging challenges. <i>Current Opinion in Neurology</i> , 2017, 30, 334-344.	1.8	80
70	Pseudotumor cerebri syndrome in childhood: incidence, clinical profile and risk factors in a national prospective population-based cohort study. <i>Archives of Disease in Childhood</i> , 2017, 102, 715-721.	1.0	72
71	High sensitivity and specificity in proposed clinical diagnostic criteria for anti-N-methyl-D-aspartate receptor encephalitis. <i>Developmental Medicine and Child Neurology</i> , 2017, 59, 1256-1260.	1.1	46
72	Immune-mediated neurological syndromes: Old meets new. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 805-806.	0.7	1

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73	Myelin oligodendrocyte glycoprotein antibody (MOG-ab) associated demyelination presenting with an opsoclonus myoclonus like syndrome. <i>European Journal of Paediatric Neurology</i> , 2017, 21, e117.	0.7	0
74	Focal status epilepticus and progressive dyskinesia: A novel phenotype for glycine receptor antibody-mediated neurological disease in children. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 414-417.	0.7	16
75	An unusual neuroimaging finding and response to immunotherapy in a child with genetically confirmed vanishing white matter disease. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 410-413.	0.7	7
76	ABO236...Association of lyve-1 protein in exosome with disease activity as a new candidate biomarker for rheumatoid arthritis. , 2017, , .		0
77	A multicentre randomised controlled TRIal of IntraVENous immunoglobulin compared with standard therapy for the treatment of transverse myelitis in adults and children (STRIVE). <i>Health Technology Assessment</i> , 2017, 21, 1-50.	1.3	20
78	Successful Treatment of Hepatitis C in Renal Transplant Recipients With Direct-Acting Antiviral Agents. <i>American Journal of Transplantation</i> , 2016, 16, 1588-1595.	2.6	201
79	Paediatric brainstem encephalitis associated with glial and neuronal autoantibodies. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 836-841.	1.1	29
80	Anti-methylaspartate (NMDA) receptor antibodies encephalitis mimicking an autistic regression. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 1092-1094.	1.1	34
81	Immunoglobulin in the treatment of Encephalitis (IgNiTE): protocol for a multicentre randomised controlled trial. <i>BMJ Open</i> , 2016, 6, e012356.	0.8	21
82	Autoimmune neurologic disorders in children. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2016, 133, 485-510.	1.0	9
83	Teaching Neuro Images : Neuroradiologic evolution of Leigh disease. <i>Neurology</i> , 2016, 87, e159-e160.	1.5	0
84	The origins and progression of CNS autoimmunity. <i>Neurology</i> , 2016, 87, 560-561.	1.5	0
85	Inflammatory Biomarkers in Childhood Arterial Ischemic Stroke. <i>Stroke</i> , 2016, 47, 2221-2228.	1.0	38
86	Intravenous immunoglobulin in paediatric neurology: evaluating effective usage and outcomes. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 1105-1106.	1.1	0
87	Postencephalitic epilepsy and drug-resistant epilepsy after infectious and antibody-associated encephalitis in childhood: Clinical and etiologic risk factors. <i>Epilepsia</i> , 2016, 57, e7-e11.	2.6	54
88	Acute flaccid weakness with myelopathy and peripheral nerve involvement in 2 children: Recent characterization of a previously observed phenomenon. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 948-952.	0.7	1
89	Pediatric transverse myelitis. <i>Neurology</i> , 2016, 87, S46-52.	1.5	92
90	Neuroimaging in encephalitis: analysis of imaging findings and interobserver agreement. <i>Clinical Radiology</i> , 2016, 71, 1050-1058.	0.5	49

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91	Sensory Processing Difficulties in Opsoclonus-Myoclonus Syndrome. <i>Journal of Child Neurology</i> , 2016, 31, 965-970.	0.7	5
92	<i>N</i> -methyl-D-aspartate receptor antibody encephalitis: how do we evaluate symptomatic treatment?. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 325-326.	1.1	1
93	Acute onset blindness: a case of optic neuritis and review of childhood optic neuritis. <i>BMJ Case Reports</i> , 2016, 2016, bcr2016214929.	0.2	2
94	Fetal acetylcholine receptor inactivation syndrome. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015, 2, e57.	3.1	50
95	Protocol for a multicentre randomised controlled Trial of Intravenous immunoglobulin versus standard therapy for the treatment of transverse myelitis in adults and children (STRIVE). <i>BMJ Open</i> , 2015, 5, e008312-e008312.	0.8	18
96	Acute idiopathic transverse myelitis in children. <i>Neurology</i> , 2015, 84, 341-349.	1.5	56
97	Autoimmune epilepsy: the search for a definition. <i>Developmental Medicine and Child Neurology</i> , 2015, 57, 402-403.	1.1	3
98	Characterization of human disease phenotypes associated with mutations in <i>TREX1</i> , <i>RNASEH2A</i> , <i>RNASEH2B</i> , <i>RNASEH2C</i> , <i>SAMHD1</i> , <i>ADAR</i> , and <i>IFIH1</i> . <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 296-312.	0.7	447
99	Fifteen minute consultation: Managing neonatal and childhood herpes encephalitis: Table 1. <i>Archives of Disease in Childhood: Education and Practice Edition</i> , 2015, 100, 58-63.	0.3	10
100	Paediatric neuromyelitis optica: clinical, MRI of the brain and prognostic features: Table 1. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 470-472.	0.9	90
101	RANBP2 mutation and acute necrotizing encephalopathy: 2 cases and a literature review of the expanding clinico-radiological phenotype. <i>European Journal of Paediatric Neurology</i> , 2015, 19, 106-113.	0.7	184
102	Clinical and radiological features of recurrent demyelination following acute disseminated encephalomyelitis (ADEM). <i>Multiple Sclerosis and Related Disorders</i> , 2015, 4, 451-456.	0.9	9
103	Earlier treatment of NMDAR antibody encephalitis in children results in a better outcome. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015, 2, e130.	3.1	96
104	Autoimmune Encephalopathies. <i>Pediatric Clinics of North America</i> , 2015, 62, 667-685.	0.9	27
105	Fifteen-minute consultation: autoimmune encephalitis. <i>Archives of Disease in Childhood: Education and Practice Edition</i> , 2015, 100, 282-287.	0.3	3
106	Myelin oligodendrocyte glycoprotein antibodies are associated with a non-MS course in children. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015, 2, e81.	3.1	205
107	Infectious and Autoantibody-Associated Encephalitis: Clinical Features and Long-term Outcome. <i>Pediatrics</i> , 2015, 135, e974-e984.	1.0	115
108	<i>N</i> -methyl-D-aspartate receptor antibody encephalitis: how much treatment is enough?. <i>Developmental Medicine and Child Neurology</i> , 2015, 57, 14-15.	1.1	4

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109	N-methyl-D-aspartate receptor antibody-mediated neurological disease: results of a UK-based surveillance study in children. Archives of Disease in Childhood, 2015, 100, 521-526.	1.0	112
110	Clinical relevance of voltage-gated potassium channel- $\alpha$ complex antibodies in children. Neurology, 2015, 85, 967-975.	1.5	57
111	Pediatric Herpes Simplex Virus Encephalitis Complicated by N-Methyl-D-aspartate Receptor Antibody Encephalitis. Journal of the Pediatric Infectious Diseases Society, 2015, 4, e17-e21.	0.6	22
112	Autoimmune encephalitis following haematopoietic stem cell transplant: a new clinical entity or a previously unrecognised one?. Translational Pediatrics, 2015, 4, 327-30.	0.5	1
113	Autoantibody biomarkers in childhood-acquired demyelinating syndromes: results from a national surveillance cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 456-461.	0.9	70
114	NMDA receptor antibodies associated with distinct white matter syndromes. Neurology: Neuroimmunology and NeuroInflammation, 2014, 1, e2.	3.1	85
115	Treatable childhood neuronopathy caused by mutations in riboflavin transporter RFVT2. Brain, 2014, 137, 44-56.	3.7	143
116	CSF albumin and immunoglobulin analyses in childhood neurologic disorders. Neurology: Neuroimmunology and NeuroInflammation, 2014, 1, e10.	3.1	9
117	Reversible Vigabatrin-Induced Life-Threatening Encephalopathy. JAMA Neurology, 2014, 71, 108.	4.5	21
118	N-methyl-D-aspartate receptor antibody-associated movement disorder without encephalopathy. Developmental Medicine and Child Neurology, 2014, 56, 190-193.	1.1	30
119	Idiopathic intracranial hypertension: new insights, new definitions but the same old problems. Developmental Medicine and Child Neurology, 2014, 56, 707-708.	1.1	0
120	Neurological Manifestations of Influenza Infection in Children and Adults: Results of a National British Surveillance Study. Clinical Infectious Diseases, 2014, 58, 775-784.	2.9	143
121	Utility and safety of rituximab in pediatric autoimmune and inflammatory CNS disease. Neurology, 2014, 83, 142-150.	1.5	275
122	Glycine receptor antibodies in PERM and related syndromes: characteristics, clinical features and outcomes. Brain, 2014, 137, 2178-2192.	3.7	430
123	N-methyl-D-aspartate receptor antibodies in post-herpes simplex virus encephalitis neurological relapse. Movement Disorders, 2014, 29, 90-96.	2.2	192
124	A study on clinical and radiological features and outcome in patients with posterior reversible encephalopathy syndrome (PRES). European Journal of Pediatrics, 2014, 173, 1225-1231.	1.3	23
125	Outcome of children with acetylcholine receptor (AChR) antibody positive juvenile myasthenia gravis following thymectomy. Neuromuscular Disorders, 2014, 24, 25-30.	0.3	24
126	Limbic Encephalitis Associated With Elevated Antithyroid Antibodies. Journal of Child Neurology, 2014, 29, 769-773.	0.7	12



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127	Guillain-Barré syndrome associated with <scp>CASPR2</scp> antibodies: two paediatric cases. Journal of the Peripheral Nervous System, 2014, 19, 246-249.	1.4	17
128	Osmotic demyelination syndrome associated with hypophosphataemia: 2 cases and a review of literature. Acta Paediatrica, International Journal of Paediatrics, 2013, 102, e164-8.	0.7	25
129	The tympanic membrane displacement analyser for monitoring intracranial pressure in children. Child's Nervous System, 2013, 29, 927-933.	0.6	46
130	Intracranial hypertension presenting with severe visual failure, without concurrent headache, in a child with nephrotic syndrome. BMC Pediatrics, 2013, 13, 167.	0.7	9
131	Paediatric acquired demyelinating syndromes: incidence, clinical and magnetic resonance imaging features. Multiple Sclerosis Journal, 2013, 19, 76-86.	1.4	116
132	Acute disseminated encephalomyelitis associated with positive voltage gated potassium channel complex antibody. Multiple Sclerosis and Related Disorders, 2013, 2, 147-150.	0.9	2
133	A glimpse at the cerebrospinal fluid immunoglobulins in neurological conditions. Does it help the clinician?. Developmental Medicine and Child Neurology, 2013, 55, 10-12.	1.1	5
134	Paediatric autoimmune encephalopathies: clinical features, laboratory investigations and outcomes in patients with or without antibodies to known central nervous system autoantigens. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 748-755.	0.9	217
135	More movements in neuroimmunology. Brain, 2012, 135, 3201-3202.	3.7	3
136	CSF diversion in refractory idiopathic intracranial hypertension: single-centre experience and review of efficacy. Child's Nervous System, 2012, 29, 263-7.	0.6	13
137	Paediatric multiple sclerosis: examining utility of the McDonald 2010 criteria. Multiple Sclerosis Journal, 2012, 18, 679-682.	1.4	20
138	Childhood presentation of <i>COL4A1</i> mutations. Developmental Medicine and Child Neurology, 2012, 54, 569-574.	1.1	61
139	A clinico-radiological phenotype of voltage-gated potassium channel complex antibody-mediated disorder presenting with seizures and basal ganglia changes. Developmental Medicine and Child Neurology, 2012, 54, 1157-1159.	1.1	8
140	Beneficial use of steroids in hereditary neuropathy with liability to pressure palsy. Developmental Medicine and Child Neurology, 2012, 54, 183-186.	1.1	8
141	Management of suspected viral encephalitis in children - Association of British Neurologists and British Paediatric Allergy, Immunology and Infection Group National Guidelines. Journal of Infection, 2012, 64, 449-477.	1.7	152
142	Childhood optic neuritis clinical features and outcome. Archives of Disease in Childhood, 2011, 96, 860-862.	1.0	73
143	Acute life threatening cerebellitis presenting with no apparent cerebellar signs. Clinical Neurology and Neurosurgery, 2011, 113, 928-930.	0.6	12
144	Prevalence of mycoplasma encephalitis. Lancet Infectious Diseases, The, 2011, 11, 425-426.	4.6	4

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145	A case of squamous cell carcinoma in an ileoanal pouch. <i>Colorectal Disease</i> , 2011, 13, e314-e315.	0.7	9
146	Secondary frosted branch angiitis in Neuro-Behçet's disease with serous macular detachment. <i>Pediatrics International</i> , 2011, 53, 285-286.	0.2	6
147	Treating inflammation in childhood neurodegenerative disorders. <i>Developmental Medicine and Child Neurology</i> , 2011, 53, 298-304.	1.1	12
148	Encephalopathy and <i>SCN1A</i> mutations. <i>Epilepsia</i> , 2011, 52, e26-30.	2.6	18
149	Immunosuppression alters disease severity in juvenile Batten disease mice. <i>Journal of Neuroimmunology</i> , 2011, 230, 169-172.	1.1	70
150	Neurological complications of pandemic influenza A H1N1 2009 infection: European case series and review. <i>European Journal of Pediatrics</i> , 2011, 170, 1007-1015.	1.3	68
151	Basilar artery dolichoectasia in childhood: evidence of vascular compromise. <i>Child's Nervous System</i> , 2011, 27, 193-196.	0.6	7
152	Paediatric UK demyelinating disease longitudinal study (PUDDLs). <i>BMC Pediatrics</i> , 2011, 11, 68.	0.7	9
153	Childhood N-Methyl-D-Aspartic Acid Receptor (NMDAR) Antibody Mediated Encephalitis. <i>Neuropediatrics</i> , 2011, 42, 177-178.	0.3	1
154	Severe acute disseminated encephalomyelitis: a paediatric intensive care population-based study. <i>Multiple Sclerosis Journal</i> , 2011, 17, 1258-1261.	1.4	46
155	Prevalence and Predictors of Vitamin D Insufficiency in Children: A Great Britain Population Based Study. <i>PLoS ONE</i> , 2011, 6, e22179.	1.1	159
156	Glucose transporter-1 deficiency syndrome: the expanding clinical and genetic spectrum of a treatable disorder. <i>Brain</i> , 2010, 133, 655-670.	3.7	356
157	Magnetic Resonance Imaging Changes in Idiopathic Intracranial Hypertension in Children. <i>Journal of Child Neurology</i> , 2010, 25, 294-299.	0.7	73
158	Meeting Review: The management of multiple sclerosis in children: a European view. <i>Multiple Sclerosis Journal</i> , 2010, 16, 1258-1267.	1.4	91
159	Use of therapeutic drug monitoring in the long-term valaciclovir therapy of relapsing herpes simplex virus encephalitis in children. <i>Journal of Antimicrobial Chemotherapy</i> , 2009, 64, 1340-1341.	1.3	9
160	Thalamic infarct presenting as apparent life-threatening event in infants. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2009, 98, 2002-2005.	0.7	5
161	Cerebellar defects in a mouse model of juvenile neuronal ceroid lipofuscinosis. <i>Brain Research</i> , 2009, 1266, 93-107.	1.1	55
162	Cutaneous signs are important in the diagnosis of the rare neoplasia syndrome Carney complex. <i>European Journal of Pediatrics</i> , 2009, 168, 1401-1404.	1.3	18

#	ARTICLE	IF	CITATIONS
163	The effects of carbon dioxide on measuring cerebral spinal fluid pressure. <i>Child's Nervous System</i> , 2009, 25, 783-784.	0.6	12
164	Immune system irregularities in lysosomal storage disorders. <i>Acta Neuropathologica</i> , 2008, 115, 159-174.	3.9	90
165	Clinical and Molecular Phenotype of Aicardi-Goutières Syndrome. <i>American Journal of Human Genetics</i> , 2007, 81, 713-725.	2.6	375
166	Massive <i>SCA7</i> expansion detected in a 7-month-old male with hypotonia, cardiomegaly, and renal compromise. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 140-143.	1.1	18
167	IgG entry and deposition are components of the neuroimmune response in Batten disease. <i>Neurobiology of Disease</i> , 2007, 25, 239-251.	2.1	57
168	Distinct patterns of serum immunoreactivity as evidence for multiple brain-directed autoantibodies in juvenile neuronal ceroid lipofuscinosis. <i>Neuropathology and Applied Neurobiology</i> , 2006, 32, 469-482.	1.8	37
169	Visual failure without headache in idiopathic intracranial hypertension. <i>Archives of Disease in Childhood</i> , 2005, 90, 206-210.	1.0	56
170	Selectivity and Types of Cell Death in the Neuronal Ceroid Lipofuscinoses (NCLs). <i>Brain Pathology</i> , 2004, 14, 86-96.	2.1	80
171	Late onset neurodegeneration in the <i>Cln3<sup>Δ</sup></i> mouse model of juvenile neuronal ceroid lipofuscinosis is preceded by low level glial activation. <i>Brain Research</i> , 2004, 1023, 231-242.	1.1	139
172	Neurological and Psychiatric Manifestations of COVID-19 in UK Children: A Prospective National Cohort Study. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0