Ming J Lim

List of Publications by Year
in descending order

172 papers	8,687 citations	36303 51 h-index	51608 86 g-index
203 all docs	203 docs citations	203 times ranked	8990 citing authors

#	Article	IF	Citations
1	Characterizing the features and course of psychiatric symptoms in children and adolescents with autoimmune encephalitis. European Archives of Psychiatry and Clinical Neuroscience, 2022, 272, 477-482.	3.2	5
2	Diagnosis and management of multiple sclerosis and other relapsing demyelinating disease in childhood. Archives of Disease in Childhood, 2022, 107, 216-222.	1.9	2
3	Early predictors of disability of paediatric-onset AQP4-IgG-seropositive neuromyelitis optica spectrum disorders. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 101-111.	1.9	16
4	Incidence of paediatric multiple sclerosis and other acquired demyelinating syndromes: 10â€year followâ€up surveillance study. Developmental Medicine and Child Neurology, 2022, 64, 502-508.	2.1	4
5	Neurological and cognitive outcomes after antibodyâ€negative autoimmune encephalitis in children. Developmental Medicine and Child Neurology, 2022, 64, 649-653.	2.1	10
6	Clinical features, investigations, and outcomes of pediatric limbic encephalitis: A multicenter study. Annals of Clinical and Translational Neurology, 2022, 9, 67-78.	3.7	7
7	Patientsâ€™ and carersâ€™ views and the importance of ethnicity, diversity and inclusion in research priority settings. Archives of Disease in Childhood, 2022, 107, 415-415.	1.9	0
8	Diagnosis and Management of Opsoclonus-Myoclonus-Ataxia Syndrome in Children. Neurology: Neuroimmunology and NeuroInflammation, 2022, 9, .	6.0	26
9	Workshop on RanBP2/Nup358 and acute necrotizing encephalopathy. Nucleus, 2022, 13, 156-171.	2.2	9
10	Pediatric Ischemic Stroke: An Infrequent Complication of <scp>SARSâ€CoV</scp>â€2. Annals of Neurology, 2021, 89, 657-665.	5.3	74
11	Acute flaccid myelitis: cause, diagnosis, and management. Lancet, The, 2021, 397, 334-346.	13.7	88
12	Systemic Inflammation Is Associated With Neurologic Involvement in Pediatric Inflammatory Multisystem Syndrome Associated With SARS-CoV-2. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	6.0	29
13	Use of Disease-Modifying Therapies in Pediatric Relapsing-Remitting Multiple Sclerosis in the United Kingdom. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	6.0	16
14	Clinical guidance for diagnosis and management of suspected Pediatric Acuteâ€onset Neuropsychiatric Syndrome in the Nordic countries. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 3153-3160.	1.5	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?. Brain, Behavior, and Immunity, 2021, 95, 19-20.	4.1	0
16	International Consensus Recommendations for the Treatment of Pediatric NMDAR Antibody Encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	6.0	70
17	Vaccination in acute immune-mediated/inflammatory disorders of the central nervous system. European Journal of Paediatric Neurology, 2021, 34, 118-122.	1.6	1
18	Neurological manifestations of SARS-CoV-2 infection in hospitalised children and adolescents in the UK: a prospective national cohort study. The Lancet Child and Adolescent Health, 2021, 5, 631-641.	5.6	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. European Journal of Paediatric Neurology, 2021, 34, 43-49.	1.6	13
20	Use and Safety of Immunotherapeutic Management of N-Methyl-<scp>d</scp>-Aspartate Receptor Antibody Encephalitis. JAMA Neurology, 2021, 78, 1333.	9.0	91
21	P080 An embedded pathway to mandibular advancement splint (MAS) construction in a tertiary hospital reduces barriers to care for low-income individuals. SLEEP Advances, 2021, 2, A47-A47.	0.2	0
22	Authors' reply regarding "On diagnosing and treating PANS/ PANDAS: questions from a patient support group". Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 3390-3391.	1.5	2
23	Idiopathic Hypertrophic Pachymeningitis: Does Earlier Treatment Improve Outcome?. Children, 2021, 8, 11.	1.5	7
24	Acute Myelopathy in Childhood. Children, 2021, 8, 1055.	1.5	2
25	Early predictors of epilepsy and subsequent relapse in children with acute disseminated encephalomyelitis. Multiple Sclerosis Journal, 2020, 26, 333-342.	3.0	37
26	Improved performance of the 2017 McDonald criteria for diagnosis of multiple sclerosis in children in a real-life cohort. Multiple Sclerosis Journal, 2020, 26, 1372-1380.	3.0	28
27	Radiological Cerebrospinal Posterior Reversible Encephalopathy Syndrome Mimicking Acute Disseminated Encephalomyelitis in a Neurologically Asymptomatic Child. Pediatric Neurology, 2020, 106, 65-67.	2.1	0
28	E.U. paediatric MOG consortium consensus: Part 5 â€“ Treatment of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. European Journal of Paediatric Neurology, 2020, 29, 41-53.	1.6	59
29	E.U. paediatric MOG consortium consensus: Part 4 â€“ Outcome of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. European Journal of Paediatric Neurology, 2020, 29, 32-40.	1.6	29
30	E.U. paediatric MOG consortium consensus: Part 1 â€“ Classification of clinical phenotypes of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. European Journal of Paediatric Neurology, 2020, 29, 2-13.	1.6	87
31	Catatonic features in children and adolescents with N-methyl-<scp>d</scp>-aspartate receptor antibody encephalitis. BJPsych Open, 2020, 6, .	0.7	5
32	Progress in the Management of Paediatric-Onset Multiple Sclerosis. Children, 2020, 7, 222.	1.5	4
33	Treatment and outcome of aquaporin-4 antibodyâ€“positive NMOSD. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	37
34	Evolving Cognitive Dysfunction in Children with Neurologically Stable Opsoclonusâ€“Myoclonus Syndrome. Children, 2020, 7, 103.	1.5	5
35	E.U. paediatric MOG consortium consensus: Part 3 â€“ Biomarkers of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. European Journal of Paediatric Neurology, 2020, 29, 22-31.	1.6	24
36	Acute Disseminated Encephalomyelitis: Current Perspectives. Children, 2020, 7, 210.	1.5	24

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37	E.U. paediatric MOG consortium consensus: Part 2 â€“ Neuroimaging features of paediatric myelin oligodendrocyte glycoprotein antibody-associated disorders. European Journal of Paediatric Neurology, 2020, 29, 14-21.	1.6	32
38	Treatment of MOG-IgG-associated disorder with rituximab: An international study of 121 patients. Multiple Sclerosis and Related Disorders, 2020, 44, 102251.	2.0	110
39	Neutrophil-to-lymphocyte ratio correlates with disease activity in myelin oligodendrocyte glycoprotein antibody associated disease (MOGAD) in children. Multiple Sclerosis and Related Disorders, 2020, 45, 102345.	2.0	13
40	Treatment of MOG antibody associated disorders: results of an international survey. Journal of Neurology, 2020, 267, 3565-3577.	3.6	64
41	Genomic Landscape of Reed-Sternberg Cells of Hodgkin Lymphoma from Children, Adolescents, and Young Adults. Klinische Padiatrie, 2020, 232, .	0.6	0
42	TheÂMovement disorder associated with NMDAR antibody-encephalitis is complex and characteristic: an expert video-rating study. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 724-726.	1.9	71
43	A framework for measurement and harmonization of pediatric multiple sclerosis etiologic research studies: The Pediatric MS Tool-Kit. Multiple Sclerosis Journal, 2019, 25, 1170-1177.	3.0	3
44	Is chronic lymphocytic inflammation with pontine perivascular enhancement responsive to steroids (CLIPPERS) in children the same condition as in adults?. Developmental Medicine and Child Neurology, 2019, 61, 490-496.	2.1	15
45	Ischemic Stroke Following Ergotamine Overdose. Pediatric Neurology, 2019, 101, 81-82.	2.1	4
46	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. JAMA Network Open, 2019, 2, e1914274.	5.9	14
47	Combined Anti-inflammatory and Neuroprotective Treatments Have the Potential to Impact Disease Phenotypes in Cln3â^'/â^' Mice. Frontiers in Neurology, 2019, 10, 963.	2.4	13
48	Utility and safety of plasma exchange in paediatric neuroimmune disorders. Developmental Medicine and Child Neurology, 2019, 61, 540-546.	2.1	8
49	Paediatric multiple sclerosis: a new era in diagnosis and treatment. Developmental Medicine and Child Neurology, 2019, 61, 1039-1049.	2.1	30
50	Mycophenolate mofetil in paediatric autoimmune or immuneâ€mediated diseases of the central nervous system: clinical experience and recommendations. Developmental Medicine and Child Neurology, 2019, 61, 458-468.	2.1	15
51	Magnetic resonance imaging in enterovirusâ€71, myelin oligodendrocyte glycoprotein antibody, aquaporinâ€4 antibody, and multiple sclerosisâ€associated myelitis in children. Developmental Medicine and Child Neurology, 2019, 61, 1108-1116.	2.1	22
52	Childhood disintegrative disorder and autism spectrum disorder: aÂsystematic review. Developmental Medicine and Child Neurology, 2019, 61, 523-534.	2.1	18
53	An increase in reports of acute flaccid paralysis (AFP) in the United Kingdom, 1 January 2018â€“21 January 2019: early findings. Eurosurveillance, 2019, 24, .	7.0	31
54	Myelin oligodendrocyte glycoprotein and aquaporinâ€4 antibodies are highly specific in children with acquired demyelinating syndromes. Developmental Medicine and Child Neurology, 2018, 60, 958-962.	2.1	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. Developmental Medicine and Child Neurology, 2018, 60, 1244-1250.	2.1	38
56	Paediatric acute disseminated encephalomyelitis followed by optic neuritis: disease course, treatment response and outcome. European Journal of Neurology, 2018, 25, 782-786.	3.3	45
57	â€˜Leukodystrophyâ€like phenotype in children with myelin oligodendrocyte glycoprotein antibodyâ€associated disease. Developmental Medicine and Child Neurology, 2018, 60, 417-423.	2.1	81
58	Disease Course and Treatment Responses in Children With Relapsing Myelin Oligodendrocyte Glycoprotein Antibodyâ€“Associated Disease. JAMA Neurology, 2018, 75, 478.	9.0	306
59	Immunotherapy-responsive childhood neurodegeneration with systemic and central nervous system inflammation. European Journal of Paediatric Neurology, 2018, 22, 882-888.	1.6	1
60	Pseudotumor cerebri syndrome in a patient with narcolepsy type 1. European Journal of Paediatric Neurology, 2018, 22, 194-198.	1.6	1
61	Endocrinopathies in paediatric-onset neuromyelitis optica spectrum disorder with aquaporin 4 (AQP4) antibody. Multiple Sclerosis Journal, 2018, 24, 679-684.	3.0	9
62	NMDA-receptor antibodies alter cortical microcircuit dynamics. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E9916-E9925.	7.1	39
63	Testing combinatorial therapies for juvenile Batten disease. Molecular Genetics and Metabolism, 2018, 123, S33.	1.1	1
64	Glutamate receptor Î2 serum antibodies in pediatric opsoclonus myoclonus ataxia syndrome. Neurology, 2018, 91, e714-e723.	1.1	43
65	The role of inflammation and hypovitaminosis-D in multiple sclerosis, schizophrenia and autism. Neurology Psychiatry and Brain Research, 2018, 29, 14-15.	2.0	0
66	Therapeutic plasma exchange in paediatric neurology: a critical review and proposed treatment algorithm. Developmental Medicine and Child Neurology, 2018, 60, 765-779.	2.1	24
67	Systematic review of immunoglobulin use in paediatric neurological and neurodevelopmental disorders. Developmental Medicine and Child Neurology, 2017, 59, 136-144.	2.1	24
68	Diagnostic algorithm for relapsing acquired demyelinating syndromes in children. Neurology, 2017, 89, 269-278.	1.1	155
69	Autoimmune encephalitis in children: clinical phenomenology, therapeutics, and emerging challenges. Current Opinion in Neurology, 2017, 30, 334-344.	3.6	80
70	Pseudotumor cerebri syndrome in childhood: incidence, clinical profile and risk factors in a national prospective population-based cohort study. Archives of Disease in Childhood, 2017, 102, 715-721.	1.9	72
71	High sensitivity and specificity in proposed clinical diagnostic criteria for antiâ€Nâ€methylâ€<scp>D</scp>â€aspartate receptor encephalitis. Developmental Medicine and Child Neurology, 2017, 59, 1256-1260.	2.1	46
72	Immune-mediated neurological syndromes: Old meets new. European Journal of Paediatric Neurology, 2017, 21, 805-806.	1.6	1

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73	Myelin oligodendrocyte glycoprotein antibody (MOG-ab) associated demyelination presenting with an opsoclonus myoclonus like syndrome. European Journal of Paediatric Neurology, 2017, 21, e117.	1.6	0
74	Focal status epilepticus and progressive dyskinesia: A novel phenotype for glycine receptor antibody-mediated neurological disease in children. European Journal of Paediatric Neurology, 2017, 21, 414-417.	1.6	16
75	An unusual neuroimaging finding and response to immunotherapy in a child with genetically confirmed vanishing white matter disease. European Journal of Paediatric Neurology, 2017, 21, 410-413.	1.6	7
76	AB0236â€…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). Health Technology Assessment, 2017, 21, 1-50.	2.8	20
78	Successful Treatment of Hepatitis C in Renal Transplant Recipients With Direct-Acting Antiviral Agents. American Journal of Transplantation, 2016, 16, 1588-1595.	4.7	201
79	Paediatric brainstem encephalitis associated with glial and neuronal autoantibodies. Developmental Medicine and Child Neurology, 2016, 58, 836-841.	2.1	29
80	Nâ€methylâ€<scp>d</scp>âaspartate (<scp>NMDA</scp>) receptor antibodies encephalitis mimicking an autistic regression. Developmental Medicine and Child Neurology, 2016, 58, 1092-1094.	2.1	34
81	I mmuno g lobuli N i n the T reatment of E ncephalitis (IgNiTE): protocol for a multicentre randomised controlled trial. BMJ Open, 2016, 6, e012356.	1.9	21
82	Autoimmune neurologic disorders in children. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2016, 133, 485-510.	1.8	9
83	Teaching Neuro Images : Neuroradiologic evolution of Leigh disease. Neurology, 2016, 87, e159-e160.	1.1	0
84	The origins and progression of CNS autoimmunity. Neurology, 2016, 87, 560-561.	1.1	0
85	Inflammatory Biomarkers in Childhood Arterial Ischemic Stroke. Stroke, 2016, 47, 2221-2228.	2.0	38
86	Intravenous immunoglobulin in paediatric neurology: evaluating effective usage and outcomes. Developmental Medicine and Child Neurology, 2016, 58, 1105-1106.	2.1	0
87	Postencephalitic epilepsy and drugâ€resistant epilepsy after infectious and antibodyâ€associated encephalitis in childhood: Clinical and etiologic risk factors. Epilepsia, 2016, 57, e7-e11.	5.1	54
88	Acute flaccid weakness with myelopathy and peripheral nerve involvement in 2 children: Recent characterization of a previously observed phenomenon. European Journal of Paediatric Neurology, 2016, 20, 948-952.	1.6	1
89	Pediatric transverse myelitis. Neurology, 2016, 87, S46-52.	1.1	92
90	Neuroimaging in encephalitis: analysis of imaging findings and interobserver agreement. Clinical Radiology, 2016, 71, 1050-1058.	1.1	49

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91	Sensory Processing Difficulties in Opsoclonus-Myoclonus Syndrome. Journal of Child Neurology, 2016, 31, 965-970.	1.4	5
92	Nâ€methylâ€<scp>d</scp>â€aspartate receptor antibody encephalitis: how do we evaluate symptomatic treatment?. Developmental Medicine and Child Neurology, 2016, 58, 325-326.	2.1	1
93	Acute onset blindness: a case of optic neuritis and review of childhood optic neuritis. BMJ Case Reports, 2016, 2016, bcr2016214929.	0.5	2
94	Fetal acetylcholine receptor inactivation syndrome. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e57.	6.0	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). BMJ Open, 2015, 5, e008312-e008312.	1.9	18
96	Acute idiopathic transverse myelitis in children. Neurology, 2015, 84, 341-349.	1.1	56
97	Autoimmune epilepsy: the search for a definition. Developmental Medicine and Child Neurology, 2015, 57, 402-403.	2.1	3
98	Characterization of human disease phenotypes associated with mutations in TREX1, RNASEH2A, RNASEH2B, RNASEH2C, SAMHD1, ADAR, and IFIH1. American Journal of Medical Genetics, Part A, 2015, 167, 296-312.	1.2	447
99	Fifteen minute consultation: Managing neonatal and childhood herpes encephalitis: TableÂ1. Archives of Disease in Childhood: Education and Practice Edition, 2015, 100, 58-63.	0.5	10
100	Paediatric neuromyelitis optica: clinical, MRI of the brain and prognostic features: TableÂ1. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 470-472.	1.9	90
101	RANBP2 mutation and acute necrotizing encephalopathy: 2 cases and a literature review of the expanding clinico-radiological phenotype. European Journal of Paediatric Neurology, 2015, 19, 106-113.	1.6	184
102	Clinical and radiological features of recurrent demyelination following acute disseminated encephalomyelitis (ADEM). Multiple Sclerosis and Related Disorders, 2015, 4, 451-456.	2.0	9
103	Earlier treatment of NMDAR antibody encephalitis in children results in a better outcome. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e130.	6.0	96
104	Autoimmune Encephalopathies. Pediatric Clinics of North America, 2015, 62, 667-685.	1.8	27
105	Fifteen-minute consultation: autoimmune encephalitis. Archives of Disease in Childhood: Education and Practice Edition, 2015, 100, 282-287.	0.5	3
106	Myelin oligodendrocyte glycoprotein antibodies are associated with a non-MS course in children. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e81.	6.0	205
107	Infectious and Autoantibody-Associated Encephalitis: Clinical Features and Long-term Outcome. Pediatrics, 2015, 135, e974-e984.	2.1	115
108	Nâ€methylâ€<scp>d</scp>â€aspartate receptor antibody encephalitis: how much treatment is enough?. Developmental Medicine and Child Neurology, 2015, 57, 14-15.	2.1	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.	1.9	112
110	Clinical relevance of voltage-gated potassium channelâ€“complex antibodies in children. Neurology, 2015, 85, 967-975.	1.1	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.	1.3	22
112	Autoimmune encephalitis following haematopoietic stem cell transplant: a new clinical entity or a previously unrecognised one?. Translational Pediatrics, 2015, 4, 327-30.	1.2	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.	1.9	70
114	NMDA receptor antibodies associated with distinct white matter syndromes. Neurology: Neuroimmunology and NeuroInflammation, 2014, 1, e2.	6.0	85
115	Treatable childhood neuronopathy caused by mutations in riboflavin transporter RFVT2. Brain, 2014, 137, 44-56.	7.6	143
116	CSF albumin and immunoglobulin analyses in childhood neurologic disorders. Neurology: Neuroimmunology and NeuroInflammation, 2014, 1, e10.	6.0	9
117	Reversible Vigabatrin-Induced Life-Threatening Encephalopathy. JAMA Neurology, 2014, 71, 108.	9.0	21
118	Nâ€methylâ€Dâ€aspartate receptor antibodyâ€associated movement disorder without encephalopathy. Developmental Medicine and Child Neurology, 2014, 56, 190-193.	2.1	30
119	Idiopathic intracranial hypertension: new insights, new definitions but the same old problems. Developmental Medicine and Child Neurology, 2014, 56, 707-708.	2.1	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.	5.8	143
121	Utility and safety of rituximab in pediatric autoimmune and inflammatory CNS disease. Neurology, 2014, 83, 142-150.	1.1	275
122	Glycine receptor antibodies in PERM and related syndromes: characteristics, clinical features and outcomes. Brain, 2014, 137, 2178-2192.	7.6	430
123	Nâ€methylâ€Dâ€aspartate receptor antibodies in postâ€herpes simplex virus encephalitis neurological relapse. Movement Disorders, 2014, 29, 90-96.	3.9	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.	2.7	23
125	Outcome of children with acetylcholine receptor (AChR) antibody positive juvenile myasthenia gravis following thymectomy. Neuromuscular Disorders, 2014, 24, 25-30.	0.6	24
126	Limbic Encephalitis Associated With Elevated Antithyroid Antibodies. Journal of Child Neurology, 2014, 29, 769-773.	1.4	12

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127	Guillainâ€BarrÃ© syndrome associated with <scp>CASPR2</scp> antibodies: two paediatric cases. Journal of the Peripheral Nervous System, 2014, 19, 246-249.	3.1	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.	1.5	25
129	The tympanic membrane displacement analyser for monitoring intracranial pressure in children. Child's Nervous System, 2013, 29, 927-933.	1.1	46
130	Intracranial hypertension presenting with severe visual failure, without concurrent headache, in a child with nephrotic syndrome. BMC Pediatrics, 2013, 13, 167.	1.7	9
131	Paediatric acquired demyelinating syndromes: incidence, clinical and magnetic resonance imaging features. Multiple Sclerosis Journal, 2013, 19, 76-86.	3.0	116
132	Acute disseminated encephalomyelitis associated with positive voltage gated potassium channel complex antibody. Multiple Sclerosis and Related Disorders, 2013, 2, 147-150.	2.0	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.	2.1	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.	1.9	217
135	More movements in neuroimmunology. Brain, 2012, 135, 3201-3202.	7.6	3
136	CSF diversion in refractory idiopathic intracranial hypertension: single-centre experience and review of efficacy. Child's Nervous System, 2012, 29, 263-7.	1.1	13
137	Paediatric multiple sclerosis: examining utility of the McDonald 2010 criteria. Multiple Sclerosis Journal, 2012, 18, 679-682.	3.0	20
138	Childhood presentation of COL4A1 mutations. Developmental Medicine and Child Neurology, 2012, 54, 569-574.	2.1	61
139	A clinicoâ€radiological phenotype of voltageâ€gated potassium channel complex antibodyâ€mediated disorder presenting with seizures and basal ganglia changes. Developmental Medicine and Child Neurology, 2012, 54, 1157-1159.	2.1	8
140	Beneficial use of steroids in hereditary neuropathy with liability to pressure palsy. Developmental Medicine and Child Neurology, 2012, 54, 183-186.	2.1	8
141	Management of suspected viral encephalitis in children â€“ Association of British Neurologists and British Paediatric Allergy, Immunology and Infection Group National Guidelines. Journal of Infection, 2012, 64, 449-477.	3.3	152
142	Childhood optic neuritis clinical features and outcome. Archives of Disease in Childhood, 2011, 96, 860-862.	1.9	73
143	Acute life threatening cerebellitis presenting with no apparent cerebellar signs. Clinical Neurology and Neurosurgery, 2011, 113, 928-930.	1.4	12
144	Prevalence of mycoplasma encephalitis. Lancet Infectious Diseases, The, 2011, 11, 425-426.	9.1	4

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145	A case of squamous cell carcinoma in an ileoanal pouch. Colorectal Disease, 2011, 13, e314-e315.	1.4	9
146	Secondary frosted branch angiitis in Neuroâ€BehÃ§et's disease with serous macular detachment. Pediatrics International, 2011, 53, 285-286.	0.5	6
147	Treating inflammation in childhood neurodegenerative disorders. Developmental Medicine and Child Neurology, 2011, 53, 298-304.	2.1	12
148	Encephalopathy and SCN1A mutations. Epilepsia, 2011, 52, e26-30.	5.1	18
149	Immunosuppression alters disease severity in juvenile Batten disease mice. Journal of Neuroimmunology, 2011, 230, 169-172.	2.3	70
150	Neurological complications of pandemic influenza A H1N1 2009 infection: European case series and review. European Journal of Pediatrics, 2011, 170, 1007-1015.	2.7	68
151	Basilar artery dolichoectasia in childhood: evidence of vascular compromise. Child's Nervous System, 2011, 27, 193-196.	1.1	7
152	Paediatric UK demyelinating disease longitudinal study (PUDDLS). BMC Pediatrics, 2011, 11, 68.	1.7	9
153	Childhood N-Methyl-D-Aspartic Acid Receptor (NMDAR) Antibody Mediated Encephalitis. Neuropediatrics, 2011, 42, 177-178.	0.6	1
154	Severe acute disseminated encephalomyelitis: a paediatric intensive care population-based study. Multiple Sclerosis Journal, 2011, 17, 1258-1261.	3.0	46
155	Prevalence and Predictors of Vitamin D Insufficiency in Children: A Great Britain Population Based Study. PLoS ONE, 2011, 6, e22179.	2.5	159
156	Glucose transporter-1 deficiency syndrome: the expanding clinical and genetic spectrum of a treatable disorder. Brain, 2010, 133, 655-670.	7.6	356
157	Magnetic Resonance Imaging Changes in Idiopathic Intracranial Hypertension in Children. Journal of Child Neurology, 2010, 25, 294-299.	1.4	73
158	Meeting Review: The management of multiple sclerosis in children: a European view. Multiple Sclerosis Journal, 2010, 16, 1258-1267.	3.0	91
159	Use of therapeutic drug monitoring in the long-term valaciclovir therapy of relapsing herpes simplex virus encephalitis in children. Journal of Antimicrobial Chemotherapy, 2009, 64, 1340-1341.	3.0	9
160	Thalamic infarct presenting as apparent lifeâ€threatening event in infants. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 2002-2005.	1.5	5
161	Cerebellar defects in a mouse model of juvenile neuronal ceroid lipofuscinosis. Brain Research, 2009, 1266, 93-107.	2.2	55
162	Cutaneous signs are important in the diagnosis of the rare neoplasia syndrome Carney complex. European Journal of Pediatrics, 2009, 168, 1401-1404.	2.7	18

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163	The effects of carbon dioxide on measuring cerebral spinal fluid pressure. Child's Nervous System, 2009, 25, 783-784.	1.1	12
164	Immune system irregularities in lysosomal storage disorders. Acta Neuropathologica, 2008, 115, 159-174.	7.7	90
165	Clinical and Molecular Phenotype of Aicardi-GoutiÃ¨res Syndrome. American Journal of Human Genetics, 2007, 81, 713-725.	6.2	375
166	Massive SCA7 expansion detected in a 7â€monthâ€old male with hypotonia, cardiomegaly, and renal compromise. Developmental Medicine and Child Neurology, 2007, 49, 140-143.	2.1	18
167	IgG entry and deposition are components of the neuroimmune response in Batten disease. Neurobiology of Disease, 2007, 25, 239-251.	4.4	57
168	Distinct patterns of serum immunoreactivity as evidence for multiple brain-directed autoantibodies in juvenile neuronal ceroid lipofuscinosis. Neuropathology and Applied Neurobiology, 2006, 32, 469-482.	3.2	37
169	Visual failure without headache in idiopathic intracranial hypertension. Archives of Disease in Childhood, 2005, 90, 206-210.	1.9	56
170	Selectivity and Types of Cell Death in the Neuronal Ceroid Lipofuscinoses (NCLs). Brain Pathology, 2004, 14, 86-96.	4.1	80
171	Late onset neurodegeneration in the Cln3â^'/â^' mouse model of juvenile neuronal ceroid lipofuscinosis is preceded by low level glial activation. Brain Research, 2004, 1023, 231-242.	2.2	139
172	Neurological and Psychiatric Manifestations of COVID-19 in UK Children: A Prospective National Cohort Study. SSRN Electronic Journal, 0, , .	0.4	0