

Ming J Lim

List of Publications by Year in descending order

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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	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> . American Journal of Medical Genetics, Part A, 2015, 167, 296-312.	1.2	447
2	Glycine receptor antibodies in PERM and related syndromes: characteristics, clinical features and outcomes. Brain, 2014, 137, 2178-2192.	7.6	430
3	Clinical and Molecular Phenotype of Aicardi-Goutières Syndrome. American Journal of Human Genetics, 2007, 81, 713-725.	6.2	375
4	Glucose transporter-1 deficiency syndrome: the expanding clinical and genetic spectrum of a treatable disorder. Brain, 2010, 133, 655-670.	7.6	356
5	Disease Course and Treatment Responses in Children With Relapsing Myelin Oligodendrocyte Glycoprotein Antibody-Associated Disease. JAMA Neurology, 2018, 75, 478.	9.0	306
6	Utility and safety of rituximab in pediatric autoimmune and inflammatory CNS disease. Neurology, 2014, 83, 142-150.	1.1	275
7	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
8	Myelin oligodendrocyte glycoprotein antibodies are associated with a non-MS course in children. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e81.	6.0	205
9	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
10	<i>N</i> -methyl-D-aspartate receptor antibodies in post-herpes simplex virus encephalitis neurological relapse. Movement Disorders, 2014, 29, 90-96.	3.9	192
11	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
12	Prevalence and Predictors of Vitamin D Insufficiency in Children: A Great Britain Population Based Study. PLoS ONE, 2011, 6, e22179.	2.5	159
13	Diagnostic algorithm for relapsing acquired demyelinating syndromes in children. Neurology, 2017, 89, 269-278.	1.1	155
14	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.	3.3	152
15	Treatable childhood neuronopathy caused by mutations in riboflavin transporter RFVT2. Brain, 2014, 137, 44-56.	7.6	143
16	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
17	Late onset neurodegeneration in the <i>Cln3^Δ/Δ mouse model of juvenile neuronal ceroid lipofuscinosis is preceded by low level glial activation. Brain Research, 2004, 1023, 231-242.</i>	2.2	139
18	Paediatric acquired demyelinating syndromes: incidence, clinical and magnetic resonance imaging features. Multiple Sclerosis Journal, 2013, 19, 76-86.	3.0	116

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19	Infectious and Autoantibody-Associated Encephalitis: Clinical Features and Long-term Outcome. <i>Pediatrics</i> , 2015, 135, e974-e984.	2.1	115
20	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.	5.6	114
21	N-methyl-D-aspartate receptor antibody-mediated neurological disease: results of a UK-based surveillance study in children. <i>Archives of Disease in Childhood</i> , 2015, 100, 521-526.	1.9	112
22	Treatment of MOG-IgG-associated disorder with rituximab: An international study of 121 patients. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 44, 102251.	2.0	110
23	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.	2.1	105
24	Earlier treatment of NMDAR antibody encephalitis in children results in a better outcome. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2015, 2, e130.	6.0	96
25	Pediatric transverse myelitis. <i>Neurology</i> , 2016, 87, S46-52.	1.1	92
26	Meeting Review: The management of multiple sclerosis in children: a European view. <i>Multiple Sclerosis Journal</i> , 2010, 16, 1258-1267.	3.0	91
27	Use and Safety of Immunotherapeutic Management of N-Methyl-D-Aspartate Receptor Antibody Encephalitis. <i>JAMA Neurology</i> , 2021, 78, 1333.	9.0	91
28	Immune system irregularities in lysosomal storage disorders. <i>Acta Neuropathologica</i> , 2008, 115, 159-174.	7.7	90
29	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.	1.9	90
30	Acute flaccid myelitis: cause, diagnosis, and management. <i>Lancet, The</i> , 2021, 397, 334-346.	13.7	88
31	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.	1.6	87
32	NMDA receptor antibodies associated with distinct white matter syndromes. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2014, 1, e2.	6.0	85
33	“Leukodystrophy-like” phenotype in children with myelin oligodendrocyte glycoprotein antibody-associated disease. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 417-423.	2.1	81
34	Selectivity and Types of Cell Death in the Neuronal Ceroid Lipofuscinoses (NCLs). <i>Brain Pathology</i> , 2004, 14, 86-96.	4.1	80
35	Autoimmune encephalitis in children: clinical phenomenology, therapeutics, and emerging challenges. <i>Current Opinion in Neurology</i> , 2017, 30, 334-344.	3.6	80
36	Pediatric Ischemic Stroke: An Infrequent Complication of SARS-CoV-2. <i>Annals of Neurology</i> , 2021, 89, 657-665.	5.3	74

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37	Magnetic Resonance Imaging Changes in Idiopathic Intracranial Hypertension in Children. <i>Journal of Child Neurology</i> , 2010, 25, 294-299.	1.4	73
38	Childhood optic neuritis clinical features and outcome. <i>Archives of Disease in Childhood</i> , 2011, 96, 860-862.	1.9	73
39	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.	1.9	72
40	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.	1.9	71
41	Immunosuppression alters disease severity in juvenile Batten disease mice. <i>Journal of Neuroimmunology</i> , 2011, 230, 169-172.	2.3	70
42	Autoantibody biomarkers in childhood-acquired demyelinating syndromes: results from a national surveillance cohort. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, 456-461.	1.9	70
43	International Consensus Recommendations for the Treatment of Pediatric NMDAR Antibody Encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	6.0	70
44	Neurological complications of pandemic influenza A H1N1 2009 infection: European case series and review. <i>European Journal of Pediatrics</i> , 2011, 170, 1007-1015.	2.7	68
45	Treatment of MOG antibody associated disorders: results of an international survey. <i>Journal of Neurology</i> , 2020, 267, 3565-3577.	3.6	64
46	Childhood presentation of COL4A1 mutations. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 569-574.	2.1	61
47	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.	1.6	59
48	IgG entry and deposition are components of the neuroimmune response in Batten disease. <i>Neurobiology of Disease</i> , 2007, 25, 239-251.	4.4	57
49	Clinical relevance of voltage-gated potassium channel complex antibodies in children. <i>Neurology</i> , 2015, 85, 967-975.	1.1	57
50	Visual failure without headache in idiopathic intracranial hypertension. <i>Archives of Disease in Childhood</i> , 2005, 90, 206-210.	1.9	56
51	Acute idiopathic transverse myelitis in children. <i>Neurology</i> , 2015, 84, 341-349.	1.1	56
52	Cerebellar defects in a mouse model of juvenile neuronal ceroid lipofuscinosis. <i>Brain Research</i> , 2009, 1266, 93-107.	2.2	55
53	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.	5.1	54
54	Fetal acetylcholine receptor inactivation syndrome. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015, 2, e57.	6.0	50

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55	Neuroimaging in encephalitis: analysis of imaging findings and interobserver agreement. <i>Clinical Radiology</i> , 2016, 71, 1050-1058.	1.1	49
56	Severe acute disseminated encephalomyelitis: a paediatric intensive care population-based study. <i>Multiple Sclerosis Journal</i> , 2011, 17, 1258-1261.	3.0	46
57	The tympanic membrane displacement analyser for monitoring intracranial pressure in children. <i>Child's Nervous System</i> , 2013, 29, 927-933.	1.1	46
58	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.	2.1	46
59	Paediatric acute disseminated encephalomyelitis followed by optic neuritis: disease course, treatment response and outcome. <i>European Journal of Neurology</i> , 2018, 25, 782-786.	3.3	45
60	Glutamate receptor $\gamma 2$ serum antibodies in pediatric opsoclonus myoclonus ataxia syndrome. <i>Neurology</i> , 2018, 91, e714-e723.	1.1	43
61	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.	7.1	39
62	Inflammatory Biomarkers in Childhood Arterial Ischemic Stroke. <i>Stroke</i> , 2016, 47, 2221-2228.	2.0	38
63	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.	2.1	38
64	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.	3.2	37
65	Early predictors of epilepsy and subsequent relapse in children with acute disseminated encephalomyelitis. <i>Multiple Sclerosis Journal</i> , 2020, 26, 333-342.	3.0	37
66	Treatment and outcome of aquaporin-4 antibody-positive NMOSD. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020, 7, .	6.0	37
67	N-methyl-D-aspartate (NMDA) receptor antibodies encephalitis mimicking an autistic regression. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 1092-1094.	2.1	34
68	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.	1.6	32
69	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, .	7.0	31
70	N-methyl-D-aspartate receptor antibody-associated movement disorder without encephalopathy. <i>Developmental Medicine and Child Neurology</i> , 2014, 56, 190-193.	2.1	30
71	Paediatric multiple sclerosis: a new era in diagnosis and treatment. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 1039-1049.	2.1	30
72	Paediatric brainstem encephalitis associated with glial and neuronal autoantibodies. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 836-841.	2.1	29

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73	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.	1.6	29
74	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, .	6.0	29
75	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.	3.0	28
76	Autoimmune Encephalopathies. <i>Pediatric Clinics of North America</i> , 2015, 62, 667-685.	1.8	27
77	Diagnosis and Management of Opsoclonus-Myoclonus-Ataxia Syndrome in Children. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2022, 9, .	6.0	26
78	Osmotic demyelination syndrome associated with hypophosphataemia: 2 cases and a review of literature. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2013, 102, e164-8.	1.5	25
79	Outcome of children with acetylcholine receptor (AChR) antibody positive juvenile myasthenia gravis following thymectomy. <i>Neuromuscular Disorders</i> , 2014, 24, 25-30.	0.6	24
80	Systematic review of immunoglobulin use in paediatric neurological and neurodevelopmental disorders. <i>Developmental Medicine and Child Neurology</i> , 2017, 59, 136-144.	2.1	24
81	Therapeutic plasma exchange in paediatric neurology: a critical review and proposed treatment algorithm. <i>Developmental Medicine and Child Neurology</i> , 2018, 60, 765-779.	2.1	24
82	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.	1.6	24
83	Acute Disseminated Encephalomyelitis: Current Perspectives. <i>Children</i> , 2020, 7, 210.	1.5	24
84	A study on clinical and radiological features and outcome in patients with posterior reversible encephalopathy syndrome (PRES). <i>European Journal of Pediatrics</i> , 2014, 173, 1225-1231.	2.7	23
85	Pediatric Herpes Simplex Virus Encephalitis Complicated by N-Methyl-D-aspartate Receptor Antibody Encephalitis. <i>Journal of the Pediatric Infectious Diseases Society</i> , 2015, 4, e17-e21.	1.3	22
86	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.	2.1	22
87	Reversible Vigabatrin-Induced Life-Threatening Encephalopathy. <i>JAMA Neurology</i> , 2014, 71, 108.	9.0	21
88	Immuno-globulin treatment of Encephalitis (IgNiTE): protocol for a multicentre randomised controlled trial. <i>BMJ Open</i> , 2016, 6, e012356.	1.9	21
89	Paediatric multiple sclerosis: examining utility of the McDonald 2010 criteria. <i>Multiple Sclerosis Journal</i> , 2012, 18, 679-682.	3.0	20
90	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.	2.8	20

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91	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.	2.1	18
92	Cutaneous signs are important in the diagnosis of the rare neoplasia syndrome Carney complex. <i>European Journal of Pediatrics</i> , 2009, 168, 1401-1404.	2.7	18
93	Encephalopathy and <i>SCN1A</i> mutations. <i>Epilepsia</i> , 2011, 52, e26-30.	5.1	18
94	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.	1.9	18
95	Childhood disintegrative disorder and autism spectrum disorder: a systematic review. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 523-534.	2.1	18
96	Guillain-Barré syndrome associated with <i>CASPR2</i> antibodies: two paediatric cases. <i>Journal of the Peripheral Nervous System</i> , 2014, 19, 246-249.	3.1	17
97	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.	1.6	16
98	Use of Disease-Modifying Therapies in Pediatric Relapsing-Remitting Multiple Sclerosis in the United Kingdom. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2021, 8, .	6.0	16
99	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.	1.9	16
100	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.	2.1	15
101	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.	2.1	15
102	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. <i>JAMA Network Open</i> , 2019, 2, e1914274.	5.9	14
103	CSF diversion in refractory idiopathic intracranial hypertension: single-centre experience and review of efficacy. <i>Child's Nervous System</i> , 2012, 29, 263-7.	1.1	13
104	Combined Anti-inflammatory and Neuroprotective Treatments Have the Potential to Impact Disease Phenotypes in <i>Cln3</i> ^{fl/fl} Mice. <i>Frontiers in Neurology</i> , 2019, 10, 963.	2.4	13
105	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.	2.0	13
106	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.	1.6	13
107	The effects of carbon dioxide on measuring cerebral spinal fluid pressure. <i>Child's Nervous System</i> , 2009, 25, 783-784.	1.1	12
108	Acute life threatening cerebellitis presenting with no apparent cerebellar signs. <i>Clinical Neurology and Neurosurgery</i> , 2011, 113, 928-930.	1.4	12

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109	Treating inflammation in childhood neurodegenerative disorders. <i>Developmental Medicine and Child Neurology</i> , 2011, 53, 298-304.	2.1	12
110	Limbic Encephalitis Associated With Elevated Antithyroid Antibodies. <i>Journal of Child Neurology</i> , 2014, 29, 769-773.	1.4	12
111	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.	0.5	10
112	Neurological and cognitive outcomes after antibody-negative autoimmune encephalitis in children. <i>Developmental Medicine and Child Neurology</i> , 2022, 64, 649-653.	2.1	10
113	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.	3.0	9
114	A case of squamous cell carcinoma in an ileoanal pouch. <i>Colorectal Disease</i> , 2011, 13, e314-e315.	1.4	9
115	Paediatric UK demyelinating disease longitudinal study (PUDDLs). <i>BMC Pediatrics</i> , 2011, 11, 68.	1.7	9
116	Intracranial hypertension presenting with severe visual failure, without concurrent headache, in a child with nephrotic syndrome. <i>BMC Pediatrics</i> , 2013, 13, 167.	1.7	9
117	CSF albumin and immunoglobulin analyses in childhood neurologic disorders. <i>Neurology: Neuroimmunology and Neuroinflammation</i> , 2014, 1, e10.	6.0	9
118	Clinical and radiological features of recurrent demyelination following acute disseminated encephalomyelitis (ADEM). <i>Multiple Sclerosis and Related Disorders</i> , 2015, 4, 451-456.	2.0	9
119	Autoimmune neurologic disorders in children. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2016, 133, 485-510.	1.8	9
120	Endocrinopathies in paediatric-onset neuromyelitis optica spectrum disorder with aquaporin 4 (AQP4) antibody. <i>Multiple Sclerosis Journal</i> , 2018, 24, 679-684.	3.0	9
121	Workshop on RanBP2/Nup358 and acute necrotizing encephalopathy. <i>Nucleus</i> , 2022, 13, 156-171.	2.2	9
122	A clinical and radiological phenotype of voltage-gated potassium channel complex antibody-mediated disorder presenting with seizures and basal ganglia changes. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 1157-1159.	2.1	8
123	Beneficial use of steroids in hereditary neuropathy with liability to pressure palsy. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 183-186.	2.1	8
124	Utility and safety of plasma exchange in paediatric neuroimmune disorders. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 540-546.	2.1	8
125	Basilar artery dolichoectasia in childhood: evidence of vascular compromise. <i>Child's Nervous System</i> , 2011, 27, 193-196.	1.1	7
126	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.	1.6	7

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127	Idiopathic Hypertrophic Pachymeningitis: Does Earlier Treatment Improve Outcome?. Children, 2021, 8, 11.	1.5	7
128	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
129	Secondary frosted branch angiitis in NeuroBehçet's disease with serous macular detachment. Pediatrics International, 2011, 53, 285-286.	0.5	6
130	Clinical guidance for diagnosis and management of suspected Pediatric Acute-Onset Neuropsychiatric Syndrome in the Nordic countries. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 3153-3160.	1.5	6
131	Thalamic infarct presenting as apparent life-threatening event in infants. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 2002-2005.	1.5	5
132	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
133	Sensory Processing Difficulties in Opsoclonus-Myoclonus Syndrome. Journal of Child Neurology, 2016, 31, 965-970.	1.4	5
134	Catatonic features in children and adolescents with <i>N</i> -methyl-D-aspartate receptor antibody encephalitis. BJPsych Open, 2020, 6, .	0.7	5
135	Evolving Cognitive Dysfunction in Children with Neurologically Stable Opsoclonus-Myoclonus Syndrome. Children, 2020, 7, 103.	1.5	5
136	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
137	Prevalence of mycoplasma encephalitis. Lancet Infectious Diseases, The, 2011, 11, 425-426.	9.1	4
138	<i>N</i> -methyl-D-aspartate receptor antibody encephalitis: how much treatment is enough?. Developmental Medicine and Child Neurology, 2015, 57, 14-15.	2.1	4
139	Ischemic Stroke Following Ergotamine Overdose. Pediatric Neurology, 2019, 101, 81-82.	2.1	4
140	Progress in the Management of Paediatric-Onset Multiple Sclerosis. Children, 2020, 7, 222.	1.5	4
141	Incidence of paediatric multiple sclerosis and other acquired demyelinating syndromes: 10-year follow-up surveillance study. Developmental Medicine and Child Neurology, 2022, 64, 502-508.	2.1	4
142	More movements in neuroimmunology. Brain, 2012, 135, 3201-3202.	7.6	3
143	Autoimmune epilepsy: the search for a definition. Developmental Medicine and Child Neurology, 2015, 57, 402-403.	2.1	3
144	Fifteen-minute consultation: autoimmune encephalitis. Archives of Disease in Childhood: Education and Practice Edition, 2015, 100, 282-287.	0.5	3

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145	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.	3.0	3
146	Acute disseminated encephalomyelitis associated with positive voltage gated potassium channel complex antibody. <i>Multiple Sclerosis and Related Disorders</i> , 2013, 2, 147-150.	2.0	2
147	Diagnosis and management of multiple sclerosis and other relapsing demyelinating disease in childhood. <i>Archives of Disease in Childhood</i> , 2022, 107, 216-222.	1.9	2
148	Acute onset blindness: a case of optic neuritis and review of childhood optic neuritis. <i>BMJ Case Reports</i> , 2016, 2016, bcr2016214929.	0.5	2
149	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.	1.5	2
150	Acute Myelopathy in Childhood. <i>Children</i> , 2021, 8, 1055.	1.5	2
151	Childhood N-Methyl-D-Aspartic Acid Receptor (NMDAR) Antibody Mediated Encephalitis. <i>Neuropediatrics</i> , 2011, 42, 177-178.	0.6	1
152	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.	1.6	1
153	N-methyl-D-aspartate receptor antibody encephalitis: how do we evaluate symptomatic treatment?. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 325-326.	2.1	1
154	Immune-mediated neurological syndromes: Old meets new. <i>European Journal of Paediatric Neurology</i> , 2017, 21, 805-806.	1.6	1
155	Immunotherapy-responsive childhood neurodegeneration with systemic and central nervous system inflammation. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 882-888.	1.6	1
156	Pseudotumor cerebri syndrome in a patient with narcolepsy type 1. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 194-198.	1.6	1
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