

Yael Hacoheh

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

84
papers

2,543
citations

27
h-index

49
g-index

95
ext. papers

3,567
ext. citations

6.3
avg, IF

5.26
L-index

#	Paper	IF	Citations
84	Should epileptic seizures as a first presentation be considered part of diagnostic criteria for relapsing remitting multiple sclerosis?. <i>Neuroimmunology Reports</i> , 2022 , 2, 100086		
83	The risk of infections for multiple sclerosis and neuromyelitis optica spectrum disorder disease-modifying treatments: Eighth European Committee for Treatment and Research in Multiple Sclerosis Focused Workshop Review. April 2021.. <i>Multiple Sclerosis Journal</i> , 2022 , 13524585211069068	5	1
82	Radiologically isolated aquaporin-4 antibody neuromyelitis optica spectrum disorder.. <i>Multiple Sclerosis Journal</i> , 2022 , 28, 676-679	5	2
81	085 Ten year follow-up surveillance of paediatric acquired demyelinating syndromes (ADS) in the UK. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2022 , 93, A127.1-A127	5.5	
80	Primary progressive multiple sclerosis presenting under the age of 18 years: Fact or fiction?. <i>Multiple Sclerosis Journal</i> , 2021 , 27, 309-314	5	4
79	What does first-line therapy mean for paediatric multiple sclerosis in the current era?. <i>Multiple Sclerosis Journal</i> , 2021 , 27, 1970-1976	5	7
78	Isolated central nervous system familial hemophagocytic lymphohistiocytosis (FHLH) presenting as a mimic of demyelination in children. <i>Multiple Sclerosis Journal</i> , 2021 , 13524585211053565	5	1
77	Use of Disease-Modifying Therapies in Pediatric Relapsing-Remitting Multiple Sclerosis in the United Kingdom. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021 , 8,	9.1	5
76	6-month multidisciplinary follow-up and outcomes of patients with paediatric inflammatory multisystem syndrome (PIMS-TS) at a UK tertiary paediatric hospital: a retrospective cohort study. <i>The Lancet Child and Adolescent Health</i> , 2021 , 5, 473-482	14.5	45
75	International Consensus Recommendations for the Treatment of Pediatric NMDAR Antibody Encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021 , 8,	9.1	11
74	Paediatric multiple sclerosis and antibody-associated demyelination: clinical, imaging, and biological considerations for diagnosis and care. <i>Lancet Neurology</i> , 2021 , 20, 136-149	24.1	19
73	Aquaporin-4 antibody neuromyelitis optica spectrum disorder: A paraneoplastic disease?. <i>Multiple Sclerosis Journal</i> , 2021 , 13524585211039755	5	0
72	2021 MAGNIMS-CMSC-NAIMS consensus recommendations on the use of MRI in patients with multiple sclerosis. <i>Lancet Neurology</i> , 2021 , 20, 653-670	24.1	44
71	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	14.5	28
70	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	3.8	2
69	Myelin-oligodendrocyte glycoprotein antibody-associated disease. <i>Lancet Neurology</i> , 2021 , 20, 762-772	17.2	37
68	Use and Safety of Immunotherapeutic Management of N-Methyl-d-Aspartate Receptor Antibody Encephalitis: A Meta-analysis. <i>JAMA Neurology</i> , 2021 , 78, 1333-1344	17.2	11

67	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	3.8	11
66	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	3.8	8
65	Treatment of MOG-IgG-associated disorder with rituximab: An international study of 121 patients. <i>Multiple Sclerosis and Related Disorders</i> , 2020 , 44, 102251	4	46
64	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	4	3
63	Neurologic and Radiographic Findings Associated With COVID-19 Infection in Children. <i>JAMA Neurology</i> , 2020 , 77, 1440-1445	17.2	188
62	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	3.8	16
61	Treatment and outcome of aquaporin-4 antibody-positive NMOSD: A multinational pediatric study. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020 , 7,	9.1	15
60	Diagnostic algorithm for children presenting with epilepsy partialis continua. <i>Epilepsia</i> , 2020 , 61, 2224-2233	2.3	1
59	Early predictors of epilepsy and subsequent relapse in children with acute disseminated encephalomyelitis. <i>Multiple Sclerosis Journal</i> , 2020 , 26, 333-342	5	20
58	Myelin oligodendrocyte glycoprotein antibodies associated disease: how important are B cells?. <i>Developmental Medicine and Child Neurology</i> , 2020 , 62, 273	3.3	
57	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	5	16
56	Utility and safety of plasma exchange in paediatric neuroimmune disorders. <i>Developmental Medicine and Child Neurology</i> , 2019 , 61, 540-546	3.3	5
55	Treatment Approaches for MOG-Ab-Associated Demyelination in Children. <i>Current Treatment Options in Neurology</i> , 2019 , 21, 2	4.4	75
54	Aquaporin-4 IgG antibody-related disorders in patients with juvenile systemic lupus erythematosus. <i>Lupus</i> , 2019 , 28, 1243-1249	2.6	4
53	Paediatric multiple sclerosis: a new era in diagnosis and treatment. <i>Developmental Medicine and Child Neurology</i> , 2019 , 61, 1039-1049	3.3	8
52	Acquired neuromyotonia in children with CASPR2 and LGI1 antibodies. <i>Developmental Medicine and Child Neurology</i> , 2019 , 61, 1344-1347	3.3	11
51	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	5.5	48
50	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	3.3	10

49	Two Cases of Guillain-Barré Syndrome Variants Presenting With Dysautonomia. <i>Child Neurology Open</i> , 2019 , 6, 2329048X19856778	1.3	5
48	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. <i>JAMA Network Open</i> , 2019 , 2, e1914274	10.4	5
47	Paediatric MOG antibody-associated ADEM with complex movement disorder: A case report. <i>Multiple Sclerosis Journal</i> , 2019 , 25, 125-128	5	8
46	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	3.3	72
45	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	3.3	19
44	Paediatric acute disseminated encephalomyelitis followed by optic neuritis: disease course, treatment response and outcome. <i>European Journal of Neurology</i> , 2018 , 25, 782-786	6	33
43	Autoimmune neurological disorders-does the age matter?. <i>European Journal of Paediatric Neurology</i> , 2018 , 22, 341-343	3.8	2
42	GLUT-1 deficiency presenting with seizures and reversible leukoencephalopathy on MRI imaging. <i>European Journal of Paediatric Neurology</i> , 2018 , 22, 1161-1164	3.8	8
41	Leukodystrophy-like phenotype in children with myelin oligodendrocyte glycoprotein antibody-associated disease. <i>Developmental Medicine and Child Neurology</i> , 2018 , 60, 417-423	3.3	45
40	Disease Course and Treatment Responses in Children With Relapsing Myelin Oligodendrocyte Glycoprotein Antibody-Associated Disease. <i>JAMA Neurology</i> , 2018 , 75, 478-487	17.2	202
39	Immunotherapy-responsive childhood neurodegeneration with systemic and central nervous system inflammation. <i>European Journal of Paediatric Neurology</i> , 2018 , 22, 882-888	3.8	1
38	Clinical commentary on late-onset neutropenia and neurological relapse during long-term rituximab therapy in MOG antibody spectrum disorder. <i>Multiple Sclerosis Journal</i> , 2018 , 24, 1648	5	
37	Therapeutic plasma exchange in paediatric neurology: a critical review and proposed treatment algorithm. <i>Developmental Medicine and Child Neurology</i> , 2018 , 60, 765-779	3.3	15
36	Neuroimmune disorders of the central nervous system in children in the molecular era. <i>Nature Reviews Neurology</i> , 2018 , 14, 433-445	15	29
35	Endocrinopathies in paediatric-onset neuromyelitis optica spectrum disorder with aquaporin 4 (AQP4) antibody. <i>Multiple Sclerosis Journal</i> , 2018 , 24, 679-684	5	5
34	Paraneoplastic cerebellar degeneration associated with anti-ITPR1 antibodies. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2017 , 4, e326	9.1	17
33	Diagnostic algorithm for relapsing acquired demyelinating syndromes in children. <i>Neurology</i> , 2017 , 89, 269-278	6.5	117
32	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	3.3	31

31	Abnormal white matter development in children with multiple sclerosis and monophasic acquired demyelination. <i>Brain</i> , 2017 , 140, 1172-1174	11.2	4
30	Acute disseminated encephalomyelitis 2016 , 372-375		
29	Salbutamol-responsive fetal acetylcholine receptor inactivation syndrome. <i>Neurology</i> , 2016 , 86, 692-4	6.5	7
28	Paediatric brainstem encephalitis associated with glial and neuronal autoantibodies. <i>Developmental Medicine and Child Neurology</i> , 2016 , 58, 836-41	3.3	25
27	N-methyl-d-aspartate (NMDA) receptor antibodies encephalitis mimicking an autistic regression. <i>Developmental Medicine and Child Neurology</i> , 2016 , 58, 1092-4	3.3	25
26	Multiple sclerosis in children 2016 , 361-364		
25	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	6.4	30
24	Clinical and radiological features of recurrent demyelination following acute disseminated encephalomyelitis (ADEM). <i>Multiple Sclerosis and Related Disorders</i> , 2015 , 4, 451-456	4	6
23	Earlier treatment of NMDAR antibody encephalitis in children results in a better outcome. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015 , 2, e130	9.1	73
22	Autoimmune encephalopathies. <i>Pediatric Clinics of North America</i> , 2015 , 62, 667-85	3.6	24
21	Myelin oligodendrocyte glycoprotein antibodies are associated with a non-MS course in children. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015 , 2, e81	9.1	144
20	Infectious and autoantibody-associated encephalitis: clinical features and long-term outcome. <i>Pediatrics</i> , 2015 , 135, e974-84	7.4	99
19	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, 521-6	2.2	96
18	Neuromyelitis optica in a child with Aicardi-Goutières syndrome. <i>Neurology</i> , 2015 , 85, 381-3	6.5	17
17	Clinical relevance of voltage-gated potassium channel complex antibodies in children. <i>Neurology</i> , 2015 , 85, 967-75	6.5	53
16	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, e17-21	4.8	18
15	Fetal acetylcholine receptor inactivation syndrome: A myopathy due to maternal antibodies. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2015 , 2, e57	9.1	34
14	N-methyl-D-aspartate receptor antibody-associated movement disorder without encephalopathy. <i>Developmental Medicine and Child Neurology</i> , 2014 , 56, 190-3	3.3	27

13	N-methyl-D-aspartate receptor antibodies in post-herpes simplex virus encephalitis neurological relapse. <i>Movement Disorders</i> , 2014 , 29, 90-6	7	160
12	Limbic encephalitis associated with elevated antithyroid antibodies. <i>Journal of Child Neurology</i> , 2014 , 29, 769-73	2.5	8
11	Guillain-Barré syndrome associated with CASPR2 antibodies: two paediatric cases. <i>Journal of the Peripheral Nervous System</i> , 2014 , 19, 246-9	4.7	14
10	Autoantibody biomarkers in childhood-acquired demyelinating syndromes: results from a national surveillance cohort. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014 , 85, 456-61	5.5	61
9	NMDA receptor antibodies associated with distinct white matter syndromes. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2014 , 1, e2	9.1	69
8	CSF albumin and immunoglobulin analyses in childhood neurologic disorders. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2014 , 1, e10	9.1	7
7	Acute disseminated encephalomyelitis associated with positive voltage gated potassium channel complex antibody. <i>Multiple Sclerosis and Related Disorders</i> , 2013 , 2, 147-50	4	2
6	Paediatric autoimmune encephalopathies: clinical features, laboratory investigations and outcomes in patients with or without antibodies to known central nervous system autoantigens. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2013 , 84, 748-55	5.5	164
5	Guillain-Barré-like syndrome associated with lung adenocarcinoma and CASPR2 antibodies. <i>Muscle and Nerve</i> , 2013 , 48, 836-7	3.4	13
4	Dominantly inherited nonprogressive cerebellar hypoplasia identified in utero: no doubt. <i>Journal of Child Neurology</i> , 2013 , 28, 279-80	2.5	
3	A clinico-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, 1157-9	3.3	7
2	Dominantly inherited nonprogressive cerebellar hypoplasia identified in utero. <i>Journal of Child Neurology</i> , 2012 , 27, 1000-3	2.5	3
1	Acute life threatening cerebellitis presenting with no apparent cerebellar signs. <i>Clinical Neurology and Neurosurgery</i> , 2011 , 113, 928-30	2	11