Aviva Mimouni-Bloch

List of Publications by Year
in descending order

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46 papers	971 citations	516710 16 h-index	477307 29 g-index
46 all docs	46 docs citations	46 times ranked	1456 citing authors

#	Article	IF	Citations
1	Sex-Specific Long-Term Height and Body Mass Index Trajectories of Children Diagnosed with Attention-Deficit/Hyperactivity Disorder and Treated with Stimulants. Journal of Pediatrics, 2021, 238, 296-304.e4.	1.8	7
2	Association between sensory modulation and sleep difficulties in children with Attention Deficit Hyperactivity Disorder (ADHD). Sleep Medicine, 2021, 84, 107-113.	1.6	6
3	The National Autism Database of Israel: a Resource for Studying Autism Risk Factors, Biomarkers, Outcome Measures, and Treatment Efficacy. Journal of Molecular Neuroscience, 2020, 70, 1303-1312.	2.3	22
4	Effect of Methylphenidate on State Anxiety in Children With ADHD-A Single Dose, Placebo Controlled, Crossover Study. Frontiers in Behavioral Neuroscience, 2019, 13, 106.	2.0	8
5	What we can learn from naming errors of children with language impairment at preschool age. Clinical Linguistics and Phonetics, 2018, 32, 298-315.	0.9	8
6	Association between sensory modulation and daily activity function of children with attention deficit/hyperactivity disorder and children with typical development. Research in Developmental Disabilities, 2018, 83, 69-76.	2.2	32
7	Graphical Product Quality and Muscle Activity in Children With Mild Disabilities Drawing on a Horizontally or Vertically Oriented Tablet. American Journal of Occupational Therapy, 2018, 72, 7206205040p1-7206205040p7.	0.3	1
8	Methylphenidate Reduces State Anxiety During a Continuous Performance Test That Distinguishes Adult ADHD Patients From Controls. Journal of Attention Disorders, 2017, 21, 46-51.	2.6	19
9	Relationship between perceived competence and performance during real and virtual motor tasks by children with developmental coordination disorder. Disability and Rehabilitation: Assistive Technology, 2017, 12, 752-757.	2.2	10
10	Development of finger force coordination in children. Experimental Brain Research, 2017, 235, 3709-3720.	1.5	8
11	Association between sensory processing disorder and daily function of children with attention deficit/hyperactive disorder and controls. European Journal of Paediatric Neurology, 2017, 21, e171.	1.6	1
12	Does pain take holidays? Non-attendance rates at a hospital-based pain clinic are elevated during the Jewish high-holidays. Israel Journal of Health Policy Research, 2017, 6, 11.	2.6	3
13	Congenital myasthenic syndrome in Israel: Genetic and clinical characterization. Neuromuscular Disorders, 2017, 27, 136-140.	0.6	30
14	Motor Difficulties and Their Effect on Participation in School-Aged Children. Journal of Child Neurology, 2016, 31, 1290-1295.	1.4	9
15	Novel homozygous missense mutation in GAN associated with Charcot-Marie-Tooth disease type 2 in a large consanguineous family from Israel. BMC Medical Genetics, 2016, 17, 82.	2.1	15
16	Trends in Physical Medicine and Rehabilitation Publications Over the Past 16 Years. Archives of Physical Medicine and Rehabilitation, 2016, 97, 1030-1033.	0.9	24
17	Congenital Anomalies of the Central Nervous System. , 2016, , 951-961.		0
18	Self-citation rate and impact factor in pediatrics. Scientometrics, 2016, 108, 1455-1460.	3.0	8

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19	Extensive Mongolian Spots and Lysosomal Storage Diseases. Journal of Pediatrics, 2016, 170, 333-333.e1.	1.8	12
20	Methylphenidate mediated change in prosody is specific to the performance of a cognitive task in female adult ADHD patients. World Journal of Biological Psychiatry, 2015, 16, 635-639.	2.6	2
21	A de-novo interstitial microduplication involving 2p16.1-p15 and mirroring 2p16.1-p15 microdeletion syndrome: Clinical and molecular analysis. European Journal of Paediatric Neurology, 2015, 19, 711-715.	1.6	16
22	Evaluating Computer Screen Time and Its Possible Link to Psychopathology in the Context of Age: A Cross-Sectional Study of Parents and Children. PLoS ONE, 2015, 10, e0140542.	2.5	12
23	Thiamine Deficiency in Infancy: Long-Term Follow-Up. Pediatric Neurology, 2014, 51, 311-316.	2.1	45
24	Breastfeeding May Protect from Developing Attention-Deficit/Hyperactivity Disorder. Breastfeeding Medicine, 2013, 8, 363-367.	1.7	77
25	Hypomyelination and congenital cataract: Identification of novel mutations in two unrelated families. European Journal of Paediatric Neurology, 2013, 17, 108-111.	1.6	7
26	The mental health consequences of student â€œHolocaust memorial journeysâ€• Australasian Psychiatry, 2013, 21, 326-328.	0.7	2
27	Can Computerized Cognitive Tests Assist in the Clinical Diagnosis of Attention-Deficit Hyperactivity Disorder?. Journal of Neuropsychiatry and Clinical Neurosciences, 2012, 24, 111-114.	1.8	8
28	Auditory System Dysfunction due to Infantile Thiamine Deficiency: Long-Term Auditory Sequelae. Audiology and Neuro-Otology, 2012, 17, 309-320.	1.3	26
29	eIF2Î³ Mutation that Disrupts eIF2 Complex Integrity Links Intellectual Disability to Impaired Translation Initiation. Molecular Cell, 2012, 48, 641-646.	9.7	63
30	An Emerging 1q21.1 Deletion-Associated Neurodevelopmental Phenotype. Journal of Child Neurology, 2011, 26, 113-116.	1.4	19
31	Infantile Cerebral and Cerebellar Atrophy Is Associated with a Mutation in the MED17 Subunit of the Transcription Preinitiation Mediator Complex. American Journal of Human Genetics, 2010, 87, 667-670.	6.2	58
32	Neuropsychiatric Manifestations in Langerhansâ€™ Cell Histiocytosis Disease: A Case Report and Review of the Literature. Journal of Child Neurology, 2010, 25, 884-887.	1.4	11
33	Familial Vasovagal Syncope Associated With Migraine. Pediatric Neurology, 2009, 40, 27-30.	2.1	16
34	Late Infantile Neuronal Ceroid Lipofuscinosis: A New Mutation in Arabs. Pediatric Neurology, 2009, 41, 297-300.	2.1	8
35	Lesions of the Corpus Callosum in Children With Neurofibromatosis 1. Pediatric Neurology, 2008, 38, 406-410.	2.1	9
36	Follow-up of preschool children with severe emotional and behavioral symptoms. Israel Journal of Psychiatry and Related Sciences, 2006, 43, 16-20.	0.5	1

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37	Identification of a locus for nongoitrous congenital hypothyroidism on chromosome 15q25.3-26.1. Human Genetics, 2005, 118, 348-355.	3.8	20
38	Autosomal Dominant Resistance to Thyrotropin as a Distinct Entity in Five Multigenerational Kindreds: Clinical Characterization and Exclusion of Candidate Loci. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4025-4034.	3.6	27
39	Sleep-Wake Patterns in Children With Intrauterine Growth Retardation. Journal of Child Neurology, 2002, 17, 872-876.	1.4	26
40	Myoclonic Seizures as a Main Manifestation of Epstein-Barr Virus Infection. Journal of Child Neurology, 2002, 17, 446-447.	1.4	6
41	Winging of the scapula in a child with hereditary multiple exostoses. Pediatric Neurology, 2002, 26, 74-76.	2.1	18
42	Electroconvulsive Therapy in Adolescents: Similarities to and Differences From Adults. Journal of the American Academy of Child and Adolescent Psychiatry, 2001, 40, 1332-1336.	0.5	52
43	Psychogenic seizures: A review. International Journal of Adolescent Medicine and Health, 2000, 12, 71-86.	1.3	3
44	Phenotypeâ€“genotype correlation in familial Mediterranean fever: evidence for an association between Met694Val and amyloidosis. European Journal of Human Genetics, 1999, 7, 287-292.	2.8	194
45	Familial hypothyroidism with autosomal dominant inheritance.. Archives of Disease in Childhood, 1996, 75, 245-246.	1.9	14
46	Clonidine-Induced Hyperglycemia in a Young Diabetic Girl. Annals of Pharmacotherapy, 1993, 27, 980-980.	1.9	8