Antonia Parmeggiani

List of Publications by Year
in descending order

67	2,043	236833 25	243529 44
papers	citations	h-index	g-index

72	72	72	2720
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	EEG features and epilepsy in patients with autism. Brain and Development, 1995, 17, 169-174.	0.6	186
2	Riboflavin prophylaxis in pediatric and adolescent migraine. Journal of Headache and Pain, 2009, 10, 361-365.	2.5	150
3	Epilepsy in adolescents and young adults with autistic disorder. Brain and Development, 2000, 22, 102-106.	0.6	131
4	Epilepsy and EEG paroxysmal abnormalities in autism spectrum disorders. Brain and Development, 2010, 32, 783-789.	0.6	111
5	Landauâ€“Kleffner syndrome (LKS): long-term follow-up and links with electrical status epilepticus during sleep (ESES). Brain and Development, 1999, 21, 90-98.	0.6	108
6	Somatic and germline mosaicisms in Severe Myoclonic Epilepsy of Infancy. Biochemical and Biophysical Research Communications, 2006, 341, 489-493.	1.0	102
7	Coâ€occurring malformations of cortical development and <scp>SCN</scp>1A gene mutations. Epilepsia, 2014, 55, 1009-1019.	2.6	84
8	Intravenous high-dose immunoglobulin treatment in recent onset childhood narcolepsy with cataplexy. Journal of Neurology, 2008, 255, 1549-1554.	1.8	74
9	A Novel Null Homozygous Mutation Confirms CACNA2D2 as a Gene Mutated in Epileptic Encephalopathy. PLoS ONE, 2013, 8, e82154.	1.1	67
10	Phosphatase and Tensin Homolog (PTEN) Gene Mutations and Autism: Literature Review and a Case Report of a Patient With Cowden Syndrome, Autistic Disorder, and Epilepsy. Journal of Child Neurology, 2012, 27, 392-397.	0.7	60
11	Prognostic factors in patients with mesial temporal lobe epilepsy. Epilepsia, 2009, 50, 41-44.	2.6	51
12	Evidence of Polymorphic CYP2C19 Involvement in the Human Metabolism of N-Desmethylclobazam. Therapeutic Drug Monitoring, 2002, 24, 737-741.	1.0	49
13	Autism and Coeliac Disease. Journal of Autism and Developmental Disorders, 2008, 38, 407-408.	1.7	49
14	Cortical reflex myoclonus in rett syndrome. Annals of Neurology, 1998, 43, 472-479.	2.8	48
15	Familial Occurrence of Febrile Seizures and Epilepsy in Severe Myoclonic Epilepsy of Infancy (SMEI) Patients with SCN1A Mutations. Epilepsia, 2006, 47, 1629-1635.	2.6	48
16	Homozygous <scp>NOTCH</scp> 3 null mutation and impaired <scp>NOTCH</scp> 3 signaling in recessive earlyâ€onset arteriopathy and cavitating leukoencephalopathy. EMBO Molecular Medicine, 2015, 7, 848-858.	3.3	48
17	Maternally inherited genetic variants of CADPS 2 are present in Autism Spectrum Disorders and Intellectual Disability patients. EMBO Molecular Medicine, 2014, 6, 795-809.	3.3	42
18	Autism Spectrum Disorder from the Womb to Adulthood: Suggestions for a Paradigm Shift. Journal of Personalized Medicine, 2021, 11, 70.	1.1	40

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19	Seizure Worsening Caused by Decreased Serum Valproate During Meropenem Therapy. Journal of Child Neurology, 2005, 20, 456-457.	0.7	39
20	Neuropsychological Findings in Childhood Narcolepsy. Journal of Child Neurology, 2014, 29, 1370-1376.	0.7	34
21	Benign myoclonic epilepsy: long-term follow-up of 11 new cases. Brain and Development, 1997, 19, 473-479.	0.6	33
22	Epilepsy, Intelligence, and Psychiatric Disorders in Patients With Cerebellar Hypoplasia. Journal of Child Neurology, 2003, 18, 1-4.	0.7	33
23	Epilepsy in Patients With Pervasive Developmental Disorder Not Otherwise Specified. Journal of Child Neurology, 2007, 22, 1198-1203.	0.7	33
24	Unusual side-effects due to clobazam: a case report with genetic study of CYP2C19. Brain and Development, 2004, 26, 63-66.	0.6	32
25	Association of the mtDNA m.4171C>A/MT-ND1 mutation with both optic neuropathy and bilateral brainstem lesions. BMC Neurology, 2014, 14, 116.	0.8	28
26	Early features of autism spectrum disorder: a cross-sectional study. Italian Journal of Pediatrics, 2019, 45, 144.	1.0	26
27	Life-threatening complications of posterior reversible encephalopathy syndrome in children. European Journal of Paediatric Neurology, 2014, 18, 632-640.	0.7	24
28	Long-term follow-up of cognitive functions in patients with continuous spikeâ€“waves during sleep (CSWS). Epilepsy and Behavior, 2016, 60, 211-217.	0.9	20
29	The syndrome of polymicrogyria, thalamic hypoplasia, and epilepsy with CSWS. Neurology, 2016, 86, 1250-1259.	1.5	19
30	The characteristics and activities of child and adolescent mental health services in Italy: a regional survey. BMC Psychiatry, 2012, 12, 7.	1.1	17
31	Autosomal dominant lateral temporal epilepsy (ADLTE): Novel structural and single-nucleotide LGI1 mutations in families with predominant visual auras. Epilepsy Research, 2015, 110, 132-138.	0.8	17
32	Autism, macrocrania and epilepsy: how are they linked?. Brain and Development, 2002, 24, 296-299.	0.6	15
33	Interictal EEG findings in two cases with â€˜double cortexâ€™ syndrome. Brain and Development, 1994, 16, 320-324.	0.6	12
34	Familial Unverricht-Lundborg Disease: A Clinical, Neurophysiologic, and Genetic Study. Epilepsia, 1997, 38, 637-641.	2.6	12
35	Brain correlates of spike and wave discharges in GLUT1 deficiency syndrome. NeuroImage: Clinical, 2017, 13, 446-454.	1.4	12
36	Autism spectrum disorder and anorexia nervosa: an Italian prospective study. Italian Journal of Pediatrics, 2021, 47, 59.	1.0	12

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37	Neuropsychological profile in Italian children with neurofibromatosis type 1 (NF1) and their relationships with neuroradiological data: Preliminary results. European Journal of Paediatric Neurology, 2018, 22, 822-830.	0.7	11
38	The Role of the Noradrenergic System in Eating Disorders: A Systematic Review. International Journal of Molecular Sciences, 2021, 22, 11086.	1.8	11
39	Neurological impairment in congenital bilateral ptosis with ophthalmoplegia. Brain and Development, 1992, 14, 107-109.	0.6	10
40	Sneddon syndrome, arylsulfatase A pseudodeficiency and impairment of cerebral white matter. Brain and Development, 2000, 22, 390-393.	0.6	10
41	Exacerbation of epileptic seizures by carbamazepine: Report of 10 cases. Seizure: the Journal of the British Epilepsy Association, 1998, 7, 479-483.	0.9	9
42	Epilepsy and trimethylaminuria: A new case report and literature review. Brain and Development, 2011, 33, 593-596.	0.6	9
43	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	0.8	9
44	Challenges in the clinical interpretation of small de novo copy number variants in neurodevelopmental disorders. Gene, 2019, 706, 162-171.	1.0	9
45	Should Echolalia Be Considered a Phonic Stereotypy? A Narrative Review. Brain Sciences, 2021, 11, 862.	1.1	9
46	Association among Autistic Traits, Treatment Intensity and Outcomes in Adolescents with Anorexia Nervosa: Preliminary Results. Journal of Clinical Medicine, 2021, 10, 3605.	1.0	9
47	Quantification of diffuse and focal delta activity in hypsarrhythmia. Brain and Development, 1990, 12, 310-315.	0.6	8
48	Posterior Fossa Malformations and Epilepsy. Journal of Child Neurology, 1999, 14, 113-117.	0.7	8
49	Epilepsy in Chromosomal Abnormalities: An Italian Sample. Journal of Child Neurology, 2005, 20, 419-423.	0.7	7
50	Inpatient treatment of anorexia nervosa with adjunctive valproate: a case series of 14 young and adolescent patients. Eating and Weight Disorders, 2022, 27, 1209-1215.	1.2	7
51	Timing of Psychopharmacological and Nutritional Interventions in the Inpatient Treatment of Anorexia Nervosa: An Observational Study. Brain Sciences, 2021, 11, 1242.	1.1	6
52	Cerebellar Hypoplasia, Continuous Spike-waves During Sleep, and Neuropsychological and Behavioral Disorders. Journal of Child Neurology, 2008, 23, 1472-1476.	0.7	5
53	Prognostic factors of drugâ€resistant epilepsy in childhood: An Italian study. Pediatrics International, 2015, 57, 1143-1148.	0.2	5
54	Pervasive refusal syndrome or anorexia nervosa: a case report with a successful behavioural treatment. Eating and Weight Disorders, 2021, 26, 2089-2093.	1.2	5

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55	Gastrointestinal Disorders and Autism. , 2014, , 2035-2046.		5
56	Low-Dose Olanzapine in the Treatment of Adolescents with Anorexia Nervosa: An Observational Naturalistic Caseâ€“Control Study. Journal of Child and Adolescent Psychopharmacology, 2022, 32, 304-310.	0.7	5
57	Eating Disorders in Infants and Toddlers. , 2022, , 5-13.		3
58	Neuropsychological implications of adjunctive levetiracetam in childhood epilepsy. Journal of Pediatric Neurosciences, 2014, 9, 115.	0.2	3
59	The effect of chronic neuroglycopenia on resting state networks in GLUT1 syndrome across the lifespan. Human Brain Mapping, 2020, 41, 453-466.	1.9	2
60	Linguistic feature of anorexia nervosa: a prospective caseâ€“control pilot study. Eating and Weight Disorders, 2022, 27, 1367-1375.	1.2	2
61	Neuropsychological impairment in early-onset hydrocephalus and epilepsy with continuous spike-waves during slow-wave sleep: A case report and literature review. Journal of Pediatric Neurosciences, 2013, 8, 141.	0.2	2
62	Specific Learning Disorders and Eating Disorders: an Italian retrospective study. Italian Journal of Pediatrics, 2022, 48, .	1.0	2
63	Methyl-CpG-binding Protein 2 (MECP2) Gene Mutations in an Italian Sample of Patients with Pervasive Developmental Disorder and Mental Retardation. Journal of Child Neurology, 2009, 24, 772-774.	0.7	1
64	Quadriceps muscle strength in Duchenne muscular dystrophy and effect of corticosteroid treatment. Acta Myologica, 2020, 39, 200-206.	1.5	1
65	Treatment response in children and adolescents with anorexia nervosa: a naturalistic, caseâ€“control study. Eating and Weight Disorders, 0, , .	1.2	1
66	Complex brain malformation and drug resistant epilepsy. Brain and Development, 1997, 19, 66-70.	0.6	0
67	Disability and inclusive education in an Italian Region: analysis of the data for the school year 2012-2013. Minerva Pediatrics, 2017, , .	0.2	0