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	Linguistic feature of anorexia nervosa: a prospective caseâ€“control pilot study. Eating and Weight Disorders, 2022, 27, 1367-1375.	1.2	2
2	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
3	Eating Disorders in Infants and Toddlers. , 2022, , 5-13.		3
4	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
5	Specific Learning Disorders and Eating Disorders: an Italian retrospective study. Italian Journal of Pediatrics, 2022, 48, .	1.0	2
6	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
7	Autism spectrum disorder and anorexia nervosa: an Italian prospective study. Italian Journal of Pediatrics, 2021, 47, 59.	1.0	12
8	Should Echolalia Be Considered a Phonic Stereotypy? A Narrative Review. Brain Sciences, 2021, 11, 862.	1.1	9
9	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
10	Timing of Psychopharmacological and Nutritional Interventions in the Inpatient Treatment of Anorexia Nervosa: An Observational Study. Brain Sciences, 2021, 11, 1242.	1.1	6
11	Autism Spectrum Disorder from the Womb to Adulthood: Suggestions for a Paradigm Shift. Journal of Personalized Medicine, 2021, 11, 70.	1.1	40
12	The Role of the Noradrenergic System in Eating Disorders: A Systematic Review. International Journal of Molecular Sciences, 2021, 22, 11086.	1.8	11
13	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
14	Quadriceps muscle strength in Duchenne muscular dystrophy and effect of corticosteroid treatment. Acta Myologica, 2020, 39, 200-206.	1.5	1
15	Challenges in the clinical interpretation of small de novo copy number variants in neurodevelopmental disorders. Gene, 2019, 706, 162-171.	1.0	9
16	Early features of autism spectrum disorder: a cross-sectional study. Italian Journal of Pediatrics, 2019, 45, 144.	1.0	26
17	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
18	Brain correlates of spike and wave discharges in GLUT1 deficiency syndrome. NeuroImage: Clinical, 2017, 13, 446-454.	1.4	12

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19	Disability and inclusive education in an Italian Region: analysis of the data for the school year 2012-2013. Minerva Pediatrics, 2017, , .	0.2	0
20	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
21	The syndrome of polymicrogyria, thalamic hypoplasia, and epilepsy with CSWS. Neurology, 2016, 86, 1250-1259.	1.5	19
22	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
23	Prognostic factors of drugâ€resistant epilepsy in childhood: An Italian study. Pediatrics International, 2015, 57, 1143-1148.	0.2	5
24	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
25	Neuropsychological Findings in Childhood Narcolepsy. Journal of Child Neurology, 2014, 29, 1370-1376.	0.7	34
26	Coâ€occurring malformations of cortical development and <scp>SCN</scp>1A gene mutations. Epilepsia, 2014, 55, 1009-1019.	2.6	84
27	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
28	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
29	Life-threatening complications of posterior reversible encephalopathy syndrome in children. European Journal of Paediatric Neurology, 2014, 18, 632-640.	0.7	24
30	Gastrointestinal Disorders and Autism. , 2014, , 2035-2046.		5
31	Neuropsychological implications of adjunctive levetiracetam in childhood epilepsy. Journal of Pediatric Neurosciences, 2014, 9, 115.	0.2	3
32	A Novel Null Homozygous Mutation Confirms CACNA2D2 as a Gene Mutated in Epileptic Encephalopathy. PLoS ONE, 2013, 8, e82154.	1.1	67
33	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
34	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
35	The characteristics and activities of child and adolescent mental health services in Italy: a regional survey. BMC Psychiatry, 2012, 12, 7.	1.1	17
36	Epilepsy and trimethylaminuria: A new case report and literature review. Brain and Development, 2011, 33, 593-596.	0.6	9

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37	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	0.8	9
38	Epilepsy and EEG paroxysmal abnormalities in autism spectrum disorders. Brain and Development, 2010, 32, 783-789.	0.6	111
39	Riboflavin prophylaxis in pediatric and adolescent migraine. Journal of Headache and Pain, 2009, 10, 361-365.	2.5	150
40	Prognostic factors in patients with mesial temporal lobe epilepsy. Epilepsia, 2009, 50, 41-44.	2.6	51
41	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
42	Autism and Coeliac Disease. Journal of Autism and Developmental Disorders, 2008, 38, 407-408.	1.7	49
43	Intravenous high-dose immunoglobulin treatment in recent onset childhood narcolepsy with cataplexy. Journal of Neurology, 2008, 255, 1549-1554.	1.8	74
44	Cerebellar Hypoplasia, Continuous Spike-waves During Sleep, and Neuropsychological and Behavioral Disorders. Journal of Child Neurology, 2008, 23, 1472-1476.	0.7	5
45	Epilepsy in Patients With Pervasive Developmental Disorder Not Otherwise Specified. Journal of Child Neurology, 2007, 22, 1198-1203.	0.7	33
46	Somatic and germline mosaicisms in Severe Myoclonic Epilepsy of Infancy. Biochemical and Biophysical Research Communications, 2006, 341, 489-493.	1.0	102
47	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
48	Epilepsy in Chromosomal Abnormalities: An Italian Sample. Journal of Child Neurology, 2005, 20, 419-423.	0.7	7
49	Seizure Worsening Caused by Decreased Serum Valproate During Meropenem Therapy. Journal of Child Neurology, 2005, 20, 456-457.	0.7	39
50	Unusual side-effects due to clobazam: a case report with genetic study of CYP2C19. Brain and Development, 2004, 26, 63-66.	0.6	32
51	Epilepsy, Intelligence, and Psychiatric Disorders in Patients With Cerebellar Hypoplasia. Journal of Child Neurology, 2003, 18, 1-4.	0.7	33
52	Evidence of Polymorphic CYP2C19 Involvement in the Human Metabolism of N-Desmethylclobazam. Therapeutic Drug Monitoring, 2002, 24, 737-741.	1.0	49
53	Autism, macrocrania and epilepsy: how are they linked?. Brain and Development, 2002, 24, 296-299.	0.6	15
54	Sneddon syndrome, arylsulfatase A pseudodeficiency and impairment of cerebral white matter. Brain and Development, 2000, 22, 390-393.	0.6	10

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55	Epilepsy in adolescents and young adults with autistic disorder. Brain and Development, 2000, 22, 102-106.	0.6	131
56	Posterior Fossa Malformations and Epilepsy. Journal of Child Neurology, 1999, 14, 113-117.	0.7	8
57	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
58	Cortical reflex myoclonus in rett syndrome. Annals of Neurology, 1998, 43, 472-479.	2.8	48
59	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
60	Complex brain malformation and drug resistant epilepsy. Brain and Development, 1997, 19, 66-70.	0.6	0
61	Benign myoclonic epilepsy: long-term follow-up of 11 new cases. Brain and Development, 1997, 19, 473-479.	0.6	33
62	Familial Unverricht-Lundborg Disease: A Clinical, Neurophysiologic, and Genetic Study. Epilepsia, 1997, 38, 637-641.	2.6	12
63	EEG features and epilepsy in patients with autism. Brain and Development, 1995, 17, 169-174.	0.6	186
64	Interictal EEG findings in two cases with â€˜double cortexâ€™ syndrome. Brain and Development, 1994, 16, 320-324.	0.6	12
65	Neurological impairment in congenital bilateral ptosis with ophthalmoplegia. Brain and Development, 1992, 14, 107-109.	0.6	10
66	Quantification of diffuse and focal delta activity in hypsarrhythmia. Brain and Development, 1990, 12, 310-315.	0.6	8
67	Treatment response in children and adolescents with anorexia nervosa: a naturalistic, caseâ€“control study. Eating and Weight Disorders, 0, , .	1.2	1