Alice Bonuccelli

List of Publications by Year
in descending order

40	527	687363 13	713466 21
papers	citations	h-index	g-index

41	41	41	821
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Epilepsy, electroclinical features, and longâ€term outcomes in Pittâ€“Hopkins syndrome due to pathogenic variants in the TCF4 gene. European Journal of Neurology, 2022, 29, 19-25.	3.3	4
2	A nationwide study on Sydenham's chorea: Clinical features, treatment and prognostic factors. European Journal of Paediatric Neurology, 2022, 36, 1-6.	1.6	9
3	Targeting Inflammatory Mediators in Epilepsy: A Systematic Review of Its Molecular Basis and Clinical Applications. Frontiers in Neurology, 2022, 13, 741244.	2.4	12
4	Therapeutic aspects of Sydenham's Chorea: an update.. Acta Biomedica, 2022, 92, e2021414.	0.3	2
5	PRES-like leukoencephalopathy presenting with status epilepticus associated with Brentuximab Vedotin treatment.. Acta Biomedica, 2022, 92, e2021416.	0.3	0
6	Pediatric traumatic brain injury: a new relation between outcome and neutrophil-to-lymphocite ratio.. Acta Biomedica, 2022, 92, e2021417.	0.3	2
7	Peripheral Arterial Tonometry (EndoPAT)-measured Endothelial Dysfunction in Migraine with Aura children.. Acta Biomedica, 2022, 92, e2021345.	0.3	0
8	A prenatal case with multiple supernumerary markers identified as derivatives of chromosomes 13, 15, and 20: molecular cytogenetic characterization and review of the literature. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 1-5.	1.5	1
9	Post-traumatic stress, anxiety, and depressive symptoms in caregivers of children tested for COVID-19 in the acute phase of the Italian outbreak. Journal of Psychiatric Research, 2021, 135, 256-263.	3.1	27
10	Electroclinical features of MEF2C haploinsufficiency-related epilepsy: A multicenter European study. Seizure: the Journal of the British Epilepsy Association, 2021, 88, 60-72.	2.0	6
11	Toward the Knowledge of the Epidemiological Impact of Acute Rheumatic Fever in Italy. Frontiers in Pediatrics, 2021, 9, 746505.	1.9	4
12	Efficacy and tolerability of mycophenolate mofetil in a pediatric Rasmussen syndrome. Epilepsy and Behavior Reports, 2020, 13, 100334.	1.0	3
13	Post-traumatic stress spectrum symptoms in parents of children affected by epilepsy: Gender differences. Seizure: the Journal of the British Epilepsy Association, 2020, 80, 169-174.	2.0	10
14	Rasmussen's encephalitis: From immune pathogenesis towards targeted-therapy. Seizure: the Journal of the British Epilepsy Association, 2020, 81, 76-83.	2.0	23
15	Genomic sequencing in severe epilepsy: a step closer to precision medicine. Expert Review of Precision Medicine and Drug Development, 2020, 5, 101-108.	0.7	1
16	Targeted re-sequencing for early diagnosis of genetic causes of childhood epilepsy: the Italian experience from the â€˜beyond epilepsyâ€™ project. Italian Journal of Pediatrics, 2020, 46, 92.	2.6	17
17	Clinical spectrum and genotype-phenotype correlations in PRRT2 Italian patients. European Journal of Paediatric Neurology, 2020, 28, 193-197.	1.6	14
18	Unilateral Lisch nodules in a pediatric patient: a sign for genetic mosaicism?. Minerva Pediatrics, 2020, , .	0.4	0

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19	The best evidence for progressive myoclonic epilepsy: A pathway to precision therapy. Seizure: the Journal of the British Epilepsy Association, 2019, 71, 247-257.	2.0	57
20	Relapse risk factors in antiâ€Nâ€methylâ€Dâ€aspartate receptor encephalitis. Developmental Medicine and Child Neurology, 2019, 61, 1101-1107.	2.1	40
21	Post-traumatic stress and major depressive disorders in parent caregivers of children with a chronic disorder. Psychiatry Research, 2019, 279, 195-200.	3.3	22
22	The spectrum of intermediate <scp>SCN</scp>8Aâ€related epilepsy. Epilepsia, 2019, 60, 830-844.	5.1	70
23	Advantages of Array Comparative Genomic Hybridization Using Buccal Swab DNA for Detecting Pallister-Killian Syndrome. Annals of Laboratory Medicine, 2019, 39, 232-234.	2.5	2
24	Adult Autism Subthreshold Spectrum (AdAS Spectrum) in parents of pediatric patients with epilepsy: Correlations with post-traumatic stress symptoms. Comprehensive Psychiatry, 2018, 83, 25-30.	3.1	18
25	Mothers and fathers of children with epilepsy: gender differences in post-traumatic stress symptoms and correlations with mood spectrum symptoms. Neuropsychiatric Disease and Treatment, 2018, Volume 14, 1371-1379.	2.2	45
26	Methylenetetrahydrofolate reductase polymorphism (MTHFR C677T) and headache in children: a retrospective study from a tertiary level outpatient service. Italian Journal of Pediatrics, 2018, 44, 106.	2.6	3
27	Generalized epilepsy and mild intellectual disability associated with 13q34 deletion: A potential role for SOX1 and ARHGEF7. Seizure: the Journal of the British Epilepsy Association, 2018, 59, 38-40.	2.0	5
28	DSM-5 criteria for PTSD in parents of pediatric patients with epilepsy: What are the changes with respect to DSM-IV-TR?. Epilepsy and Behavior, 2017, 70, 97-103.	1.7	18
29	A 6.5â€‰mb deletion at 3q24q25.2 narrows Wisconsin syndrome critical region to a 750â€‰kb interval: A potential role for MBNLI. American Journal of Medical Genetics, Part A, 2017, 173, 280-284.	1.2	2
30	Maternally derived 15q11.2â€q13.1 duplication in a child with Lennoxâ€“Gastautâ€type epilepsy and dysmorphic features: Clinicalâ€genetic characterization of the family and review of the literature. American Journal of Medical Genetics, Part A, 2017, 173, 556-560.	1.2	1
31	A Case of 22q11 Deletion Syndrome (22q11DS) with a Panayiotopoulos Epileptic Pattern: Are Additional Copy-Number Variations a Possible Second Hit in Modulating the 22q11DS Phenotype?. Frontiers in Pediatrics, 2017, 5, 48.	1.9	4
32	A rare case of hypomelanosis of Ito presenting with generalized alopecia. Minerva Pediatrica, 2016, 68, 382-3.	2.7	0
33	17q12 Microduplications: A challenge for clinicians. American Journal of Medical Genetics, Part A, 2015, 167, 674-676.	1.2	6
34	Molecular cytogenetic characterization of an interstitial deletion of chromosome 21 (21q22.13q22.3) in a patient with dysmorphic features, intellectual disability and severe generalized epilepsy. European Journal of Medical Genetics, 2012, 55, 362-366.	1.3	36
35	Pitt-Hopkins syndrome: report of a case with a TCF4 gene mutation. Italian Journal of Pediatrics, 2010, 36, 12.	2.6	24
36	Molecular cytogenetic characterization of a translocation t(13;22)(q22.3;q11.23) in a patient with idiopathic partial epilepsy. Epilepsy Research, 2009, 86, 89-93.	1.6	1

#	Article	IF	Citations
37	Molecular cytogenetic characterization of a de novo mosaic supernumerary ring chromosome 7: Report of a new patient. American Journal of Medical Genetics, Part A, 2008, 146A, 2955-2959.	1.2	3
38	Acute Myelitis in a Child: Current Hypotheses. Pediatric Neurology, 2006, 35, 430-432.	2.1	0
39	Migraine-like attacks in child with Sturge-Weber syndrome without facial nevus. Pediatric Neurology, 2005, 32, 131-133.	2.1	23
40	Personalized medicine in epilepsy patients. , 0, , .		8