

Alice Bonuccelli

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

527
citations

687363

13
h-index

713466

21
g-index

41
all docs

41
docs citations

41
times ranked

821
citing authors

#	ARTICLE	IF	CITATIONS
1	The spectrum of intermediate <i><sc>SCN</sc>8A</i>-related epilepsy. <i>Epilepsia</i> , 2019, 60, 830-844.	5.1	70
2	The best evidence for progressive myoclonic epilepsy: A pathway to precision therapy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 71, 247-257.	2.0	57
3	Mothers and fathers of children with epilepsy: gender differences in post-traumatic stress symptoms and correlations with mood spectrum symptoms. <i>Neuropsychiatric Disease and Treatment</i> , 2018, Volume 14, 1371-1379.	2.2	45
4	Relapse risk factors in anti- <i>CD&Casp</i> caspartate receptor encephalitis. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 1101-1107.	2.1	40
5	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. <i>European Journal of Medical Genetics</i> , 2012, 55, 362-366.	1.3	36
6	Post-traumatic stress, anxiety, and depressive symptoms in caregivers of children tested for COVID-19 in the acute phase of the Italian outbreak. <i>Journal of Psychiatric Research</i> , 2021, 135, 256-263.	3.1	27
7	Pitt-Hopkins syndrome: report of a case with a TCF4 gene mutation. <i>Italian Journal of Pediatrics</i> , 2010, 36, 12.	2.6	24
8	Migraine-like attacks in child with Sturge-Weber syndrome without facial nevus. <i>Pediatric Neurology</i> , 2005, 32, 131-133.	2.1	23
9	Rasmussen's encephalitis: From immune pathogenesis towards targeted-therapy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020, 81, 76-83.	2.0	23
10	Post-traumatic stress and major depressive disorders in parent caregivers of children with a chronic disorder. <i>Psychiatry Research</i> , 2019, 279, 195-200.	3.3	22
11	DSM-5 criteria for PTSD in parents of pediatric patients with epilepsy: What are the changes with respect to DSM-IV-TR?. <i>Epilepsy and Behavior</i> , 2017, 70, 97-103.	1.7	18
12	Adult Autism Subthreshold Spectrum (AdAS Spectrum) in parents of pediatric patients with epilepsy: Correlations with post-traumatic stress symptoms. <i>Comprehensive Psychiatry</i> , 2018, 83, 25-30.	3.1	18
13	Targeted re-sequencing for early diagnosis of genetic causes of childhood epilepsy: the Italian experience from the "beyond epilepsy" project. <i>Italian Journal of Pediatrics</i> , 2020, 46, 92.	2.6	17
14	Clinical spectrum and genotype-phenotype correlations in PRRT2 Italian patients. <i>European Journal of Paediatric Neurology</i> , 2020, 28, 193-197.	1.6	14
15	Targeting Inflammatory Mediators in Epilepsy: A Systematic Review of Its Molecular Basis and Clinical Applications. <i>Frontiers in Neurology</i> , 2022, 13, 741244.	2.4	12
16	Post-traumatic stress spectrum symptoms in parents of children affected by epilepsy: Gender differences. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020, 80, 169-174.	2.0	10
17	A nationwide study on Sydenham's chorea: Clinical features, treatment and prognostic factors. <i>European Journal of Paediatric Neurology</i> , 2022, 36, 1-6.	1.6	9
18	Personalized medicine in epilepsy patients. , 0, , .		8

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19	17q12 Microduplications: A challenge for clinicians. American Journal of Medical Genetics, Part A, 2015, 167, 674-676.	1.2	6
20	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
21	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
22	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
23	Epilepsy, electroclinical features, and long-term outcomes in Pitt-Hopkins syndrome due to pathogenic variants in the <i>TCF4</i> gene. European Journal of Neurology, 2022, 29, 19-25.	3.3	4
24	Toward the Knowledge of the Epidemiological Impact of Acute Rheumatic Fever in Italy. Frontiers in Pediatrics, 2021, 9, 746505.	1.9	4
25	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
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	Efficacy and tolerability of mycophenolate mofetil in a pediatric Rasmussen syndrome. Epilepsy and Behavior Reports, 2020, 13, 100334.	1.0	3
28	A 6.5 μ m deletion at 3q24q25.2 narrows Wisconsin syndrome critical region to a 750 μ kb interval: A potential role for <i>MBNL1</i> . American Journal of Medical Genetics, Part A, 2017, 173, 280-284.	1.2	2
29	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
30	Therapeutic aspects of Sydenham's Chorea: an update.. Acta Biomedica, 2022, 92, e2021414.	0.3	2
31	Pediatric traumatic brain injury: a new relation between outcome and neutrophil-to-lymphocyte ratio.. Acta Biomedica, 2022, 92, e2021417.	0.3	2
32	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
33	Maternally derived 15q11.2-q13.1 duplication in a child with Lennox-Gastaut type epilepsy and dysmorphic features: Clinical genetic characterization of the family and review of the literature. American Journal of Medical Genetics, Part A, 2017, 173, 556-560.	1.2	1
34	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
35	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
36	Acute Myelitis in a Child: Current Hypotheses. Pediatric Neurology, 2006, 35, 430-432.	2.1	0

#	ARTICLE	IF	CITATIONS
37	Unilateral Lisch nodules in a pediatric patient: a sign for genetic mosaicism?. <i>Minerva Pediatrics</i> , 2020, , .	0.4	0
38	A rare case of hypomelanosis of Ito presenting with generalized alopecia. <i>Minerva Pediatrica</i> , 2016, 68, 382-3.	2.7	0
39	PRES-like leukoencephalopathy presenting with status epilepticus associated with Brentuximab Vedotin treatment.. <i>Acta Biomedica</i> , 2022, 92, e2021416.	0.3	0
40	Peripheral Arterial Tonometry (EndoPAT)-measured Endothelial Dysfunction in Migraine with Aura children.. <i>Acta Biomedica</i> , 2022, 92, e2021345.	0.3	0