Salvatore Savasta

List of Publications by Year
in descending order

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87 papers	1,680 citations	304743 22 h-index	345221 36 g-index
91 all docs	91 docs citations	91 times ranked	2435 citing authors

#	Article	IF	Citations
1	Headache in progressive facial hemiatrophy (Parry-Romberg syndrome): A paradigmatic case and systematic review of the literature. Cephalalgia, 2022, 42, 409-425.	3.9	2
2	Absence of lingual frenulum in children with Ehlers-Danlos Syndrome: a retrospective study of forty cases and literature review of a twenty years long debate. Minerva Pediatrics, 2021, 73, .	0.4	3
3	Absence of lingual frenulum in children with Ehlers-Danlos Syndrome: a retrospective study of forty cases and literature review of a twenty years long debate. Minerva Pediatrics, 2021, 73, 230-235.	0.4	3
4	Further delineation and long-term evolution of electroclinical phenotype in Mowat Wilson Syndrome. A longitudinal study in 40 individuals. Epilepsy and Behavior, 2021, 124, 108315.	1.7	5
5	A Novel 4q32.3 Deletion in a Boy: Additional Signs and the Role of MARCH1. Journal of Pediatric Genetics, 2021, 10, 259-265.	0.7	1
6	Gelastic seizures not associated with hypothalamic hamartoma: A long-term follow-up study. Epilepsy and Behavior, 2020, 103, 106578.	1.7	8
7	Clinical variability in children with dolichoarteriopathies of the internal carotid artery. Child's Nervous System, 2020, 36, 621-628.	1.1	3
8	Challenges and management of neurological and psychiatric manifestations in SARS-CoV-2 (COVID-19) patients. Neurological Sciences, 2020, 41, 2353-2366.	1.9	60
9	Pai syndrome: a review. Child's Nervous System, 2020, 36, 2635-2640.	1.1	4
10	Challenges and management of neurological and psychiatric manifestations in SARS-CoV-2 (COVID-19) patients. , 2020, 41, 2353.		1
11	Postural analysis in a pediatric cohort of patients with Ehlers-Danlos Syndrome: a pilot study. Minerva Pediatrica, 2020, 72, 73-78.	2.7	2
12	Innovative therapies for malignant brain tumors: the road to a tailored cure. Acta Biomedica, 2020, 91, 5-17.	0.3	21
13	Adoptive immunotherapies in neuro-oncology: classification, recent advances, and translational challenges. Acta Biomedica, 2020, 91, 18-31.	0.3	12
14	Gene therapies for high-grade gliomas: from the bench to the bedside. Acta Biomedica, 2020, 91, 32-50.	0.3	9
15	The impact of stem cells in neuro-oncology: applications, evidence, limitations and challenges. Acta Biomedica, 2020, 91, 51-60.	0.3	10
16	Potential roads for reaching the summit: an overview on target therapies for high-grade gliomas. Acta Biomedica, 2020, 91, 61-78.	0.3	12
17	Targeting the medulloblastoma: a molecular-based approach. Acta Biomedica, 2020, 91, 79-100.	0.3	14
18	Advanced pharmacological therapies for neurofibromatosis type 1-related tumors. Acta Biomedica, 2020, 91, 101-114.	0.3	7

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19	Gradenigoâ€™s syndrome with abscess of the petrous apex in pediatric patients: what is the best treatment?. Child's Nervous System, 2019, 35, 2265-2272.	1.1	6
20	Lower Airway Microbiota. Frontiers in Pediatrics, 2019, 7, 393.	1.9	38
21	Relapse risk factors in antiâ€Nâ€methylâ€Dâ€aspartate receptor encephalitis. Developmental Medicine and Child Neurology, 2019, 61, 1101-1107.	2.1	40
22	Sleep in Mowat-Wilson Syndrome: a clinical and video-polysomnographic study. Sleep Medicine, 2019, 61, 44-51.	1.6	4
23	Melkerssonâ€“Rosenthal Syndrome in Childhood: Report of Three Paediatric Cases and a Review of the Literature. International Journal of Environmental Research and Public Health, 2019, 16, 1289.	2.6	28
24	Acute ataxia in paediatric emergency departments: a multicentre Italian study. Archives of Disease in Childhood, 2019, 104, 768-774.	1.9	27
25	Understanding Childhood Neuroimmune Diseases of the Central Nervous System. Frontiers in Pediatrics, 2019, 7, 511.	1.9	23
26	Transient symptomatic zinc deficiency in a breastâ€fed African infant: case report and literature review. International Journal of Dermatology, 2019, 58, 963-965.	1.0	2
27	Measurement of nitric oxide and assessment of airway diseases in children: an update. Minerva Pediatrica, 2019, 71, 524-532.	2.7	12
28	Phenotype and genotype of 87 patients with Mowatâ€“Wilson syndrome and recommendations for care. Genetics in Medicine, 2018, 20, 965-975.	2.4	67
29	Clinical characteristics of headache in Italian adolescents aged 11â€“16Âyears: a cross-sectional questionnaire school-based study. Italian Journal of Pediatrics, 2018, 44, 44.	2.6	24
30	Hypomelanosis of Ito. Journal of Pediatric Neurology, 2018, 16, 265-275.	0.2	2
31	Coexistence of childhood absence epilepsy and benign epilepsy with centrotemporal spikes: A case series. European Journal of Paediatric Neurology, 2017, 21, 570-575.	1.6	19
32	X-Linked Hypohidrotic Ectodermal Dysplasia: New Features and a Novel <i>EDA</i> Gene Mutation. Cytogenetic and Genome Research, 2017, 152, 111-116.	1.1	10
33	Efficacy of rituximab as third-line therapy in combined central and peripheral demyelination. Neurology: Clinical Practice, 2017, 7, 534-537.	1.6	5
34	Neuroimaging findings in Mowatâ€“Wilson syndrome: a study of 54 patients. Genetics in Medicine, 2017, 19, 691-700.	2.4	45
35	Nucleotide variation in Sabin type 3 poliovirus from an Albanian infant with agammaglobulinemia and vaccine associated poliomyelitis. BMC Infectious Diseases, 2016, 16, 277.	2.9	19
36	Total Hemi-overgrowth in Pigmentary Mosaicism of the (Hypomelanosis of) Ito Type. Medicine (United) Tj ETQq0 0 0 rgBT /Overlock 10	1.0	33

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37	Epilepsy in the setting of full trisomy 18: A multicenter study on 18 affected children with and without structural brain abnormalities. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2016, 172, 288-295.	1.6	24
38	Epilepsy is a possible feature in Williamsâ€Beuren syndrome patients harboring typical deletions of the 7q11.23 critical region. American Journal of Medical Genetics, Part A, 2016, 170, 148-155.	1.2	29
39	Paediatric clinically isolated syndromes: report of seven cases, differential diagnosis and literature review. Child's Nervous System, 2016, 32, 69-77.	1.1	6
40	Long-term outcome of epilepsy in patients with Praderâ€“Willi syndrome. Journal of Neurology, 2015, 262, 116-123.	3.6	10
41	Stickler syndrome associated with epilepsy: report of three cases. European Journal of Pediatrics, 2015, 174, 697-701.	2.7	9
42	Management of Pediatric Pseudotumor Cerebri Syndrome. Journal of Pediatric Neurology, 2015, 13, 058-061.	0.2	1
43	Headache and Diplopia after Rapid Maxillary Expansion: A Clue to Underdiagnosed Pseudotumor Cerebri Syndrome?. Journal of Pediatric Neurology, 2015, 13, 031-034.	0.2	2
44	Octreotide: A Therapeutic Option for Pediatric Pseudotumor Cerebri Syndrome?. Journal of Pediatric Neurology, 2015, 13, 046-048.	0.2	0
45	Tinnitus and Pediatric Pseudotumor Cerebri Syndrome. Journal of Pediatric Neurology, 2015, 13, 038-041.	0.2	1
46	Unilateral periventricular heterotopia and epilepsy in a girl with Ehlersâ€“Danlos syndrome. Epilepsy & Behavior Case Reports, 2015, 4, 27-29.	1.5	3
47	Improving molecular diagnosis in epilepsy by a dedicated high-throughput sequencing platform. European Journal of Human Genetics, 2015, 23, 354-362.	2.8	64
48	Pediatric idiopathic intracranial hypertension and the underlying endocrine-metabolic dysfunction: a pilot study. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 107-15.	0.9	41
49	Longâ€term prognosis of patients with <scp>E</scp>hlersâ€<scp>D</scp>anlos syndrome and epilepsy. Epilepsia, 2014, 55, 1213-1219.	5.1	5
50	Clinical guidelines in pediatric headache: evaluation of quality using the AGREE II instrument. Journal of Headache and Pain, 2014, 15, 57.	6.0	30
51	Long-term follow-up in children with benign convulsions associated with gastroenteritis. European Journal of Paediatric Neurology, 2014, 18, 572-577.	1.6	30
52	Seizures in fetal alcohol spectrum disorders: Evaluation of clinical, electroencephalographic, and neuroradiologic features in a pediatric case series. Epilepsia, 2014, 55, e60-6.	5.1	24
53	Gelastic epilepsy without hypothalamic hamartoma: Three additional cases. Epilepsy and Behavior, 2014, 37, 87-90.	1.7	5
54	CHARGEâ€like presentation, craniosynostosis and mild Mowatâ€“Wilson Syndrome diagnosed by recognition of the distinctive facial gestalt in a cohort of 28 new cases. American Journal of Medical Genetics, Part A, 2014, 164, 2557-2566.	1.2	20

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55	Intramedullary spinal cord cavernous malformations presenting with unexplained chest pain: case report and review of the literature. Child's Nervous System, 2013, 29, 323-328.	1.1	3
56	Epilepsy in Mowatâ€“Wilson syndrome: Delineation of the electroclinical phenotype. American Journal of Medical Genetics, Part A, 2013, 161, 273-284.	1.2	42
57	Epilepsy in patients with Cornelia de Lange syndrome: A clinical series. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 356-359.	2.0	23
58	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	1.8	25
59	Lacosamide in pediatric and adult patients: Comparison of efficacy and safety. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 210-216.	2.0	60
60	Agenesis of Internal Carotid Artery and Hypopituitarism: Case Report and Review of Literature. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 3414-3420.	3.6	9
61	Microcephaly associated with Legg-CalvÃ¨-Perthes disease in two siblings. Neurological Sciences, 2012, 33, 1401-1405.	1.9	2
62	Kikuchi-Fujimoto Disease Complicated by Peripheral Neuropathy. Pediatric Neurology, 2012, 46, 319-321.	2.1	10
63	Seizures and EEG patterns in Pallisterâ€“Killian syndrome: 13ÂNew Italian patients. European Journal of Paediatric Neurology, 2012, 16, 636-641.	1.6	13
64	Seizures and epilepsy in Sotos syndrome: Analysis of 19 Caucasian patients with longâ€term followâ€up. Epilepsia, 2012, 53, e102-5.	5.1	30
65	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	2.0	10
66	CDKL5 geneâ€related epileptic encephalopathy: electroclinical findings in the first year of life. Developmental Medicine and Child Neurology, 2011, 53, 354-360.	2.1	64
67	Ehlersâ€“Danlos syndrome and neurological features: a review. Child's Nervous System, 2011, 27, 365-371.	1.1	25
68	Benign convulsions associated with mild gastroenteritis: A multicenter clinical study. Epilepsy Research, 2011, 93, 107-114.	1.6	57
69	A Rare Cause of Recurrent Spontaneous Pneumothorax. Clinical Pediatrics, 2011, 50, 456-458.	0.8	4
70	Wolf-Hirschhorn syndrome: diagnosis using hand radiograph performed for bone age. Pediatric Radiology, 2010, 40, 1580-1580.	2.0	3
71	Italian Consensus Conference on Epilepsy and Pregnancy, Labor and Puerperium. Epilepsia, 2009, 50, 7-23.	5.1	31
72	Pai syndrome: a further report of a case with bifid nose, lipoma, and agenesis of the corpus callosum. Child's Nervous System, 2008, 24, 773-776.	1.1	20

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73	Neurofibromatosis type 1 with external genitalia involvement. Journal of Pediatric Surgery, 2008, 43, 1998-2003.	1.6	44
74	Ehlers-Danlos Syndromes. , 2008, , 887-906.		6
75	Infantile Encephalopathy and Defective Mitochondrial DNA Translation in Patients with Mutations of Mitochondrial Elongation Factors EFG1 and EFTu. American Journal of Human Genetics, 2007, 80, 44-58.	6.2	172
76	Subependymal Periventricular Heterotopias in a Patient With Ehlers-Danlos Syndrome: A New Case. Journal of Child Neurology, 2007, 22, 317-320.	1.4	8
77	Poly-epiphyseal overgrowth: description of a previously unreported skeletal dysplasia. Pediatric Radiology, 2007, 37, 1025-1030.	2.0	5
78	Wolfâ€“Hirschhorn syndrome-associated chromosome changes are not mediated by olfactory receptor gene clusters nor by inversion polymorphism on 4p16. Human Genetics, 2007, 122, 423-430.	3.8	12
79	Holoprosencephaly with neurogenic hypernatremia: a new case. Child's Nervous System, 2007, 24, 139-142.	1.1	6
80	Gastrointestinal phenotype of ATR-X syndrome. American Journal of Medical Genetics, Part A, 2006, 140A, 1172-1176.	1.2	24
81	Mitochondrial DNA Deletion in a Child With Mitochondrial Encephalomyopathy, Growth Hormone Deficiency, and Hypoparathyroidism. Journal of Child Neurology, 2006, 21, 983-985.	1.4	20
82	FISH characterization of a supernumerary r(1)(::cen?q22::q22?sq21::) chromosome associated with multiple anomalies and bilateral cataracts. American Journal of Medical Genetics Part A, 2001, 104, 157-164.	2.4	13
83	The motilin gene: subregional localisation, tissue expression, DNA polymorphisms and exclusion as a candidate gene for the HLA-associated immotile cilia syndrome. Human Genetics, 1994, 94, 671-4.	3.8	15
84	Infantile Mitochondria Encephalomyopathies: Report on 4 Cases. European Neurology, 1993, 33, 54-61.	1.4	5
85	Hemorrhagic Shock and Encephalopathy: Diagnostic Criteria. JAMA Pediatrics, 1992, 146, 279.	3.0	0
86	Muscle phosphofructokinase deficiency in a myopathic child with severe mental retardation and aplasia of cerebellar vermis. Child's Nervous System, 1992, 8, 237-241.	1.1	10
87	HLA haplotype segregation and ultrastructural study in familial immotile-cilia syndrome. Human Genetics, 1992, 89, 270-4.	3.8	19