Mariasavina Severino

List of Publications by Year
in descending order

192	3,917	186265	189892
papers	citations	28 h-index	50 g-index

203	203	203	6487
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Structural brain abnormalities in the common epilepsies assessed in a worldwide ENIGMA study. Brain, 2018, 141, 391-408.	7.6	352
2	ADA2 deficiency (DADA2) as an unrecognised cause of early onset polyarteritis nodosa and stroke: a multicentre national study. Annals of the Rheumatic Diseases, 2017, 76, 1648-1656.	0.9	199
3	Definitions and classification of malformations of cortical development: practical guidelines. Brain, 2020, 143, 2874-2894.	7.6	145
4	White matter abnormalities across different epilepsy syndromes in adults: an ENIGMA-Epilepsy study. Brain, 2020, 143, 2454-2473.	7.6	123
5	Network-based atrophy modeling in the common epilepsies: A worldwide ENIGMA study. Science Advances, 2020, 6, .	10.3	97
6	Diffuse Leptomeningeal Glioneuronal Tumors: A New Entity?. Brain Pathology, 2010, 20, 361-366.	4.1	95
7	Congenital tumors of the central nervous system. Neuroradiology, 2010, 52, 531-548.	2.2	87
8	The pulvinar sign: frequency and clinical correlations in Fabry disease. Journal of Neurology, 2008, 255, 738-744.	3.6	82
9	Novel DyneinDYNC1H1Neck and Motor Domain Mutations Link Distal Spinal Muscular Atrophy and Abnormal Cortical Development. Human Mutation, 2014, 35, 298-302.	2.5	77
10	Optic pathway glioma: Long-term visual outcome in children without neurofibromatosis type-1. Pediatric Blood and Cancer, 2010, 55, 1083-1088.	1.5	70
11	Low-grade intraventricular hemorrhage: is ultrasound good enough?. Journal of Maternal-Fetal and Neonatal Medicine, 2015, 28, 2261-2264.	1.5	68
12	Sinus pericranii: diagnosis and management in 21 pediatric patients. Journal of Neurosurgery: Pediatrics, 2015, 15, 60-70.	1.3	67
13	New MR sequences (diffusion, perfusion, spectroscopy) in brain tumours. Pediatric Radiology, 2010, 40, 999-1009.	2.0	53
14	Grading and outcome prediction of pediatric diffuse astrocytic tumors with diffusion and arterial spin labeling perfusion MRI in comparison with 18Fâ€“DOPA PET. European Journal of Nuclear Medicine and Molecular Imaging, 2017, 44, 2084-2093.	6.4	53
15	Accuracy of ultrasound in assessing cerebellar haemorrhages in very low birthweight babies. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2015, 100, F289-F292.	2.8	51
16	A Delayed Methadone Encephalopathy: Clinical and Neuroradiological Findings. Journal of Child Neurology, 2010, 25, 748-751.	1.4	50
17	Loss of SMPD4 Causes a Developmental Disorder Characterized by Microcephaly and Congenital Arthrogryposis. American Journal of Human Genetics, 2019, 105, 689-705.	6.2	48
18	The Shrunken, Bright Cerebellum: A Characteristic MRI Finding in Congenital Disorders of Glycosylation Type 1a. American Journal of Neuroradiology, 2012, 33, 2062-2067.	2.4	47

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19	The <scp>ENIGMAâ€Epilepsy</scp> working group: Mapping disease from large data sets. Human Brain Mapping, 2022, 43, 113-128.	3.6	47
20	Update on neuroimaging phenotypes of mid-hindbrain malformations. Neuroradiology, 2015, 57, 113-138.	2.2	45
21	Pontine tegmental cap dysplasia: developmental and cognitive outcome in three adolescent patients. Orphanet Journal of Rare Diseases, 2011, 6, 36.	2.7	44
22	Pediatric astrocytic tumor grading: comparison between arterial spin labeling and dynamic susceptibility contrast MRI perfusion. Neuroradiology, 2018, 60, 437-446.	2.2	43
23	Neuroimaging Changes in Menkes Disease, Part 1. American Journal of Neuroradiology, 2017, 38, 1850-1857.	2.4	42
24	The effects of mild germinal matrix-intraventricular haemorrhage on the developmental white matter microstructure of preterm neonates: a DTI study. European Radiology, 2018, 28, 1157-1166.	4.5	41
25	Advanced MR imaging and 18F-DOPA PET characteristics of H3K27M-mutant and wild-type pediatric diffuse midline gliomas. European Journal of Nuclear Medicine and Molecular Imaging, 2019, 46, 1685-1694.	6.4	41
26	Three de novo DDX3X variants associated with distinctive brain developmental abnormalities and brain tumor in intellectually disabled females. European Journal of Human Genetics, 2019, 27, 1254-1259.	2.8	41
27	Bilateral lesions of the basal ganglia and thalami (central grey matter)â€"pictorial review. Neuroradiology, 2020, 62, 1565-1605.	2.2	36
28	Atlas of lesion locations and postsurgical seizure freedom in focal cortical dysplasia: A MELD study. Epilepsia, 2022, 63, 61-74.	5.1	36
29	Neuroradiologic findings and followâ€up with magnetic resonance imaging of the genetic forms of haemophagocytic lymphohistiocytosis with CNS involvement. Pediatric Blood and Cancer, 2012, 58, 810-814.	1.5	32
30	Expanded spectrum of Pelizaeusâ€"Merzbacher-like disease: literature revision and description of a novel GJC2 mutation in an unusually severe form. European Journal of Human Genetics, 2013, 21, 34-39.	2.8	30
31	Differences in subependymal vein anatomy may predispose preterm infants to GMHâ€"IVH. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2018, 103, F59-F65.	2.8	30
32	Early Pain Exposure Influences Functional Brain Connectivity in Very Preterm Neonates. Frontiers in Neuroscience, 2019, 13, 899.	2.8	30
33	De Novo Pathogenic Variants in N-cadherin Cause a Syndromic Neurodevelopmental Disorder with Corpus Callosum, Axon, Cardiac, Ocular, and Genital Defects. American Journal of Human Genetics, 2019, 105, 854-868.	6.2	29
34	Cryopyrin-associated Periodic Syndromes in Italian Patients: Evaluation of the Rate of Somatic NLRP3 Mosaicism and Phenotypic Characterization. Journal of Rheumatology, 2017, 44, 1667-1673.	2.0	28
35	Improvement in White Matter Tract Reconstruction with Constrained Spherical Deconvolution and Track Density Mapping in Low Angular Resolution Data: A Pediatric Study and Literature Review. Frontiers in Pediatrics, 2017, 5, 182.	1.9	28
36	CNNM2 homozygous mutations cause severe refractory hypomagnesemia, epileptic encephalopathy and brain malformations. European Journal of Medical Genetics, 2019, 62, 198-203.	1.3	28

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37	Biallelic variants in LIG3 cause a novel mitochondrial neurogastrointestinal encephalomyopathy. Brain, 2021, 144, 1451-1466.	7.6	28
38	Congenital aural atresia associated with agenesis of internal carotid artery in a girl with a FOXI3 deletion. American Journal of Medical Genetics, Part A, 2015, 167, 537-544.	1.2	27
39	Spinal Cord Infarction Due to Fibrocartilaginous Embolization: The Role of Diffusion Weighted Imaging and Short-Tau Inversion Recovery Sequences. Journal of Child Neurology, 2010, 25, 1024-1028.	1.4	26
40	Skull base osteomyelitis and potential cerebrovascular complications in children. Pediatric Radiology, 2012, 42, 867-874.	2.0	25
41	Ability of 18F-DOPA PET/CT and fused 18F-DOPA PET/MRI to assess striatal involvement in paediatric glioma. European Journal of Nuclear Medicine and Molecular Imaging, 2016, 43, 1664-1672.	6.4	25
42	T2*-based MR imaging (gradient echo or susceptibility-weighted imaging) in midline and off-midline intracranial germ cell tumors: a pilot study. Neuroradiology, 2018, 60, 89-99.	2.2	25
43	Damaging de novo missense variants inEEF1A2lead to a developmental and degenerative epilepticâ€dyskinetic encephalopathy. Human Mutation, 2020, 41, 1263-1279.	2.5	24
44	Expanding the spectrum of congenital anomalies of the diencephalicâ€“mesencephalic junction. Neuroradiology, 2016, 58, 33-44.	2.2	23
45	Brain-injured Survivors of Monochorionic Twin Pregnancies Complicated by Single Intrauterine Death: MR Findings in a Multicenter Study. Radiology, 2018, 288, 582-590.	7.3	23
46	Midbrain-Hindbrain Involvement in Septo-Optic Dysplasia. American Journal of Neuroradiology, 2014, 35, 1586-1592.	2.4	22
47	Novel AMPD2 mutation in pontocerebellar hypoplasia, dysmorphisms, and teeth abnormalities. Neurology: Genetics, 2017, 3, e179.	1.9	22
48	Role of MRI T2-DRIVE in the assessment of pituitary stalk abnormalities without gadolinium in pituitary diseases. European Journal of Endocrinology, 2018, 178, 613-622.	3.7	22
49	Loss of Wwox Perturbs Neuronal Migration and Impairs Early Cortical Development. Frontiers in Neuroscience, 2020, 14, 644.	2.8	22
50	A systemsâ€level analysis highlights microglial activation as a modifying factor in common epilepsies. Neuropathology and Applied Neurobiology, 2022, 48, .	3.2	22
51	Delayed rotation of the cerebellar vermis: a pitfall in early second-trimester fetal magnetic resonance imaging. Ultrasound in Obstetrics and Gynecology, 2016, 48, 121-124.	1.7	21
52	Biallelic MFSD2A variants associated with congenital microcephaly, developmental delay, and recognizable neuroimaging features. European Journal of Human Genetics, 2020, 28, 1509-1519.	2.8	21
53	Transverse dural sinuses: incidence of anatomical variants and flow artefacts with 2D time-of-flight MR venography at 1 Tesla. Radiologia Medica, 2010, 115, 326-338.	7.7	20
54	Neuroimaging of Infectious and Inflammatory Diseases of the Pediatric Cerebellum and Brainstem. Neuroimaging Clinics of North America, 2016, 26, 471-487.	1.0	20

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55	White matter involvement in a family with a novel PDGFB mutation. Neurology: Genetics, 2016, 2, e77.	1.9	19
56	Regional impairment of cortical and deep gray matter perfusion in preterm neonates with low-grade germinal matrix-intraventricular hemorrhage: an ASL study. Neuroradiology, 2020, 62, 1689-1699.	2.2	19
57	Intermittentâ€relapsing pyruvate dehydrogenase complex deficiency: a case with clinical, biochemical, and neuroradiological reversibility. Developmental Medicine and Child Neurology, 2012, 54, 472-476.	2.1	18
58	Beyond spinal muscular atrophy with lower extremity dominance: cerebellar hypoplasia associated with a novel mutation in <scp>BICD</scp>2. European Journal of Neurology, 2016, 23, e19-21.	3.3	18
59	Posterior Fossa Malformations. Neuroimaging Clinics of North America, 2019, 29, 367-383.	1.0	18
60	Substrate reduction therapy with Miglustat in pediatric patients with GM1 type 2 gangliosidosis delays neurological involvement: A multicenter experience. Molecular Genetics & Genomic Medicine, 2020, 8, e1371.	1.2	18
61	Topographic divergence of atypical cortical asymmetry and atrophy patterns in temporal lobe epilepsy. Brain, 2022, 145, 1285-1298.	7.6	18
62	Variability of Cerebral Deep Venous System in Preterm and Term Neonates Evaluated on MR SWI Venography. American Journal of Neuroradiology, 2016, 37, 2144-2149.	2.4	17
63	Moyamoya Vasculopathy in PHACE Syndrome: Six New Cases and Review of the Literature. World Neurosurgery, 2017, 108, 291-302.	1.3	17
64	Diagnostic Approach to Macrocephaly in Children. Frontiers in Pediatrics, 2021, 9, 794069.	1.9	17
65	Pituitary deficiency and congenital infiltrating lipomatosis of the face in a girl with deletion of chromosome 1q24.3q31.1. American Journal of Medical Genetics, Part A, 2014, 164, 495-499.	1.2	16
66	MR Imaging Diagnosis of Diencephalic-Mesencephalic Junction Dysplasia in Fetuses with Developmental Ventriculomegaly. American Journal of Neuroradiology, 2017, 38, 1643-1646.	2.4	16
67	Punctate white matter lesions of preterm infants: Risk factor analysis. European Journal of Paediatric Neurology, 2019, 23, 733-739.	1.6	16
68	Increased Childhood Peripheral Facial Palsy in the Emergency Department During COVID-19 Pandemic. Pediatric Emergency Care, 2020, 36, e595-e596.	0.9	16
69	Added value of diffusion weighted imaging in pediatric central nervous system embryonal tumors surveillance. Oncotarget, 2017, 8, 60401-60413.	1.8	16
70	Association of achondroplasia with sagittal synostosis and scaphocephaly in two patients, an underestimated condition?. American Journal of Medical Genetics, Part A, 2015, 167, 646-652.	1.2	15
71	Expanding the Clinical and Magnetic Resonance Spectrum of Leukoencephalopathy with Thalamus and Brainstem Involvement and High Lactate (LTBL) in a Patient Harboring a Novel EARS2 Mutation. JIMD Reports, 2015, 23, 85-89.	1.5	15
72	Incidental findings on routine brain MRI scans in preterm infants. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2017, 102, F73-F78.	2.8	15

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73	ABCC6 mutations and early onset stroke: Two cases of a typical Pseudoxanthoma Elasticum. European Journal of Paediatric Neurology, 2018, 22, 725-728.	1.6	15
74	Quantitative susceptibility map analysis in preterm neonates with germinal matrixâ€intraventricular hemorrhage. Journal of Magnetic Resonance Imaging, 2018, 48, 1199-1207.	3.4	15
75	A further contribution to the delineation of epileptic phenotype in PACS2-related syndrome. Seizure: the Journal of the British Epilepsy Association, 2020, 79, 53-55.	2.0	15
76	Impact on rehabilitation programs during COVID-19 containment for children with pediatric and perinatal stroke. European Journal of Physical and Rehabilitation Medicine, 2020, 56, 692-694.	2.2	15
77	Intradural Extramedullary Ependymoma with Leptomeningeal Dissemination: The First Case Report in a Child and Literature Review. World Neurosurgery, 2015, 84, 865.e13-865.e19.	1.3	14
78	Correlation of multimodal ¹⁸F-DOPA PET and conventional MRI with treatment response and survival in children with diffuse intrinsic pontine gliomas. Theranostics, 2020, 10, 11881-11891.	10.0	14
79	Diagnostic Approach to Pediatric Spine Disorders. Magnetic Resonance Imaging Clinics of North America, 2016, 24, 621-644.	1.1	13
80	Structural Connectivity Analysis in Children with Segmental Callosal Agenesis. American Journal of Neuroradiology, 2017, 38, 639-647.	2.4	13
81	Widening the Heterogeneity of Leigh Syndrome: Clinical, Biochemical, and Neuroradiologic Features in a Patient Harboring a NDUFA10 Mutation. JIMD Reports, 2017, 37, 37-43.	1.5	13
82	3q29 microduplication syndrome: Description of two new cases and delineation of the minimal critical region. European Journal of Medical Genetics, 2018, 61, 428-433.	1.3	13
83	Role of diffusion weighted imaging for differentiating cerebral pilocytic astrocytoma and ganglioglioma BRAF V600E-mutant from wild type. Neuroradiology, 2020, 62, 71-80.	2.2	13
84	Targeted re-sequencing in malformations of cortical development: genotype-phenotype correlations. Seizure: the Journal of the British Epilepsy Association, 2020, 80, 145-152.	2.0	13
85	Anorectal malformation and spinal dysraphism: the value of diffusion-weighted imaging in detecting associated intradural (epi)dermoid cyst. Journal of Pediatric Surgery, 2008, 43, 1935-1938.	1.6	12
86	New insights into central nervous system involvement in FOP: Case report and review of the literature. American Journal of Medical Genetics, Part A, 2015, 167, 2817-2821.	1.2	12
87	Pretransplant management of basilar artery aneurysm and moyamoya disease in a child with Alagille syndrome. Liver Transplantation, 2015, 21, 1227-1230.	2.4	12
88	Novel asymptomatic CNS findings in patients withACVR1/ALK2mutations causing fibrodysplasia ossificans progressiva. Journal of Medical Genetics, 2016, 53, 859-864.	3.2	12
89	Agenesis of the putamen and globus pallidus caused by recessive mutations in the homeobox gene GSX2. Brain, 2019, 142, 2965-2978.	7.6	12
90	Dissecting the neurological phenotype in children with callosal agenesis, interhemispheric cysts and malformations of cortical development. Journal of Neurology, 2019, 266, 1167-1181.	3.6	12

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91	Adult-onset glutaric aciduria type I: rare presentation of a treatable disorder. Neurogenetics, 2020, 21, 179-186.	1.4	12
92	Arterial spin labeling perfusion in neonates. Seminars in Fetal and Neonatal Medicine, 2020, 25, 101130.	2.3	12
93	Guidelines for magnetic resonance imaging in pediatric head and neck pathologies: a multicentre international consensus paper. Neuroradiology, 2022, 64, 1081-1100.	2.2	12
94	Chronic cystic lesion of the sacrum: characterisation with diffusion-weighted MR imaging. Radiologia Medica, 2008, 113, 739-746.	7.7	11
95	Analysis of NADP+-dependent isocitrate dehydrogenase-1/2 gene mutations in pediatric brain tumors: report of a secondary anaplastic astrocytoma carrying the IDH1 mutation. Journal of Neuro-Oncology, 2012, 109, 477-484.	2.9	11
96	Severe growth hormone deficiency and pituitary malformation in a patient with chromosome 2p25 duplication and 2q37 deletion. Molecular Cytogenetics, 2014, 7, 41.	0.9	11
97	Early impairment of somatosensory evoked potentials in very young children with achondroplasia with foramen magnum stenosis. Developmental Medicine and Child Neurology, 2017, 59, 192-198.	2.1	11
98	Familial ROBO1 deletion associated with ectopic posterior pituitary, duplication of the pituitary stalk and anterior pituitary hypoplasia. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 95-99.	0.9	11
99	Listeria meningitis complicated by hydrocephalus in an immunocompetent child: case report and review of the literature. Italian Journal of Pediatrics, 2020, 46, 111.	2.6	11
100	Spatial coefficient of variation applied to arterial spin labeling MRI may contribute to predict surgical revascularization outcomes in pediatric moyamoya vasculopathy. Neuroradiology, 2020, 62, 1003-1015.	2.2	11
101	Biallelic variants in KARS1 are associated with neurodevelopmental disorders and hearing loss recapitulated by the knockout zebrafish. Genetics in Medicine, 2021, 23, 1933-1943.	2.4	11
102	Spinal motor neuron involvement in a patient with homozygous PRUNE mutation. European Journal of Paediatric Neurology, 2018, 22, 541-543.	1.6	10
103	White matter and cerebellar involvement in alternating hemiplegia of childhood. Journal of Neurology, 2020, 267, 1300-1311.	3.6	10
104	Homozygous missense WIPI2 variants cause a congenital disorder of autophagy with neurodevelopmental impairments of variable clinical severity and disease course. Brain Communications, 2021, 3, fcab183.	3.3	10
105	Widening the Neuroimaging Features of Adenosine Deaminase 2 Deficiency. American Journal of Neuroradiology, 2021, 42, 975-979.	2.4	10
106	Cognitive and White Matter Microstructure Development in Congenital Hypothyroidism and Familial Thyroid Disorders. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3990-e4006.	3.6	10
107	Idiopathic intervertebral disc calcification in childhood. Archives of Disease in Childhood, 2009, 94, 233-234.	1.9	9
108	Novel CNS malformations and skeletal anomalies in a patient with Beaulieuâ€boycottâ€Innes syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 2835-2840.	1.2	9

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109	Intragenic Microdeletion of <i>ULK4</i> and Partial Microduplication of <i>BRWD3</i> in Siblings with Neuropsychiatric Features and Obesity. Cytogenetic and Genome Research, 2018, 156, 14-21.	1.1	9
110	Novel compound heterozygous pathogenic variants in nucleotide-binding protein like protein (NUBPL) cause leukoencephalopathy with multi-systemic involvement. Molecular Genetics and Metabolism, 2020, 129, 26-34.	1.1	9
111	Targeted re-sequencing in pediatric and perinatal stroke. European Journal of Medical Genetics, 2020, 63, 104030.	1.3	9
112	Characteristic Cochlear Hypoplasia in Patients with Walker-Warburg Syndrome: A Radiologic Study of the Inner Ear in Î±-Dystroglycanâ€“Related Muscular Disorders. American Journal of Neuroradiology, 2021, 42, 167-172.	2.4	9
113	A novel homozygous MCOLN1 double mutant allele leading to TRP channel domain ablation underlies Mucolipidosis IV in an Italian Child. Metabolic Brain Disease, 2015, 30, 681-686.	2.9	8
114	Interstitial de novo 18q22.3q23 deletion: clinical, neuroradiological and molecular characterization of a new case and review of the literature. Molecular Cytogenetics, 2016, 9, 78.	0.9	8
115	Unusual white matter involvement in EAST syndrome associated with novel KCNJ10 mutations. Journal of Neurology, 2018, 265, 1419-1425.	3.6	8
116	Novel homozygous TSFM pathogenic variant associated with encephalocardiomyopathy with sensorineural hearing loss and peculiar neuroradiologic findings. Neurogenetics, 2019, 20, 165-172.	1.4	8
117	Clinical and neuroimaging features of the m.10197G>A mtDNA mutation: New case reports and expansion of the phenotype variability. Journal of the Neurological Sciences, 2019, 399, 69-75.	0.6	8
118	L1CAM variants cause two distinct imaging phenotypes on fetal MRI. Annals of Clinical and Translational Neurology, 2021, 8, 2004-2012.	3.7	8
119	Biallelic ADAM22 pathogenic variants cause progressive encephalopathy and infantile-onset refractory epilepsy. Brain, 2022, 145, 2301-2312.	7.6	8
120	Further genotypeâ€“phenotype correlation emerging from two families with PLP1 exon 4 skipping. Clinical Genetics, 2014, 85, 267-272.	2.0	7
121	Clinico-radiological and molecular characterization of a child with ring chromosome 2 presenting growth failure, microcephaly, kidney and brain malformations. Molecular Cytogenetics, 2015, 8, 17.	0.9	7
122	Torcular pseudomass: a potential diagnostic pitfall in infants and young children. Pediatric Radiology, 2017, 47, 227-234.	2.0	7
123	CASK related disorder: Epilepsy and developmental outcome. European Journal of Paediatric Neurology, 2021, 31, 61-69.	1.6	7
124	Ganglionic Eminence Anomalies and Coexisting Cerebral Developmental Anomalies on Fetal MR Imaging: Multicenter-Based Review of 60 Cases. American Journal of Neuroradiology, 2021, 42, 1151-1156.	2.4	7
125	Nosological Differences in the Nature of Punctate White Matter Lesions in Preterm Infants. Frontiers in Neurology, 2021, 12, 657461.	2.4	7
126	Successful urgent neurosugery management with rFVIIa mega doses in a child with haemophilia A and high titre inhibitor. Blood Coagulation and Fibrinolysis, 2014, 25, 518-521.	1.0	6

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127	Clinical and Molecular Characterization of Two Patients with CNTN6 Copy Number Variations. Cytogenetic and Genome Research, 2018, 156, 144-149.	1.1	6
128	Asymmetric cavernous sinus enlargement: a novel finding in Sturgeâ€“Weber syndrome. Neuroradiology, 2019, 61, 595-602.	2.2	6
129	Perinatal Arterial Ischemic Stroke in Fetal Vascular Malperfusion: A Case Series and Literature Review. American Journal of Neuroradiology, 2020, 41, 2377-2383.	2.4	6
130	Cortical formation abnormalities on foetal MR imaging: a proposed classification system trialled on 356 cases from Italian and UK centres. European Radiology, 2020, 30, 5250-5260.	4.5	6
131	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.1	6
132	An atypical case of post-varicella stroke in a child presenting with hemichorea followed by late-onset inflammatory focal cerebral arteriopathy. Quantitative Imaging in Medicine and Surgery, 2021, 11, 463-471.	2.0	6
133	A rare triad of morning glory disc anomaly, moyamoya vasculopathy, and transsphenoidal cephalocele: pathophysiological considerations and surgical management. Neurological Sciences, 2021, 42, 5433-5439.	1.9	6
134	Expanding the clinical and neuroimaging features of post-varicella arteriopathy of childhood. Journal of Neurology, 2021, 268, 4846-4865.	3.6	6
135	Epileptic encephalopathy caused by <scp>ARV1</scp> deficiency: Refinement of the genotypeâ€“phenotype spectrum and functional impact on <scp>GPI</scp>â€anchored proteins. Clinical Genetics, 2021, 100, 607-614.	2.0	6
136	Radiological-Pathological Comparison in a Case of Conjoined Gnatho-Thoracopagus Twins. Fetal Diagnosis and Therapy, 2009, 26, 223-226.	1.4	5
137	MR Imaging of Neonatal Brain Infections. Magnetic Resonance Imaging Clinics of North America, 2011, 19, 761-775.	1.1	5
138	Spontaneously Regressing Leukoencephalopathy With Bilateral Temporal Cysts in Congenital Rubella Infection. Pediatric Infectious Disease Journal, 2014, 33, 422-424.	2.0	5
139	Teaching Neuro Images : Figure of 8. Neurology, 2017, 89, e172-e173.	1.1	5
140	Reversible cerebral vasoconstriction complicating cerebral atherosclerotic vascular disease in Schimke immuno-osseous dysplasia. Neuroradiology, 2018, 60, 885-888.	2.2	5
141	Prenatal magnetic resonance imaging within the 26th week of gestation may predict the fate of isolated upward rotation of the cerebellar vermis: insights from a multicentre study. European Radiology, 2020, 30, 2161-2170.	4.5	5
142	Bivalirudin anticoagulation to overcome heparin resistance in a neonate with cerebral sinovenus thrombosis. Blood Coagulation and Fibrinolysis, 2020, 31, 97-100.	1.0	5
143	Placental Pathology Findings and the Risk of Intraventricular and Cerebellar Hemorrhage in Preterm Neonates. Frontiers in Neurology, 2020, 11, 761.	2.4	5
144	Sinus pericranii, skull defects, and structural brain anomalies in TRAF7 â€related disorder. Birth Defects Research, 2020, 112, 1085-1092.	1.5	5

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145	Role of visual evoked potentials and optical coherence tomography in the screening for optic pathway gliomas in patients with neurofibromatosis type I. European Journal of Ophthalmology, 2021, 31, 698-703.	1.3	5
146	Photoparoxysmal response in ADCK3 autosomal recessive ataxia: a case report and literature review. Epileptic Disorders, 2021, 23, 153-160.	1.3	5
147	External ventricular drainage for posthemorrhagic ventricular dilatation in preterm infants: insights on efficacy and failure. Journal of Neurosurgery: Pediatrics, 2021, 28, 563-571.	1.3	5
148	Epilepsy in KAT6A syndrome: Description of two individuals and revision of the literature. European Journal of Medical Genetics, 2022, 65, 104380.	1.3	5
149	Loss of Neuron Navigator 2 Impairs Brain and Cerebellar Development. Cerebellum, 2022, , 1.	2.5	5
150	Noninvasive Assessment of Hemodynamic Stress Distribution after Indirect Revascularization for Pediatric Moyamoya Vasculopathy. American Journal of Neuroradiology, 2018, 39, 1157-1163.	2.4	4
151	Severe early-onset developmental and epileptic encephalopathy (DEE) associated with novel compound heterozygous pathogenic variants in SLC25A22: Case report and literature review. Seizure: the Journal of the British Epilepsy Association, 2019, 70, 56-58.	2.0	4
152	Cognitive Profiles and Brain Volume Are Affected in Patients with Silverâ€“Russell Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1478-e1488.	3.6	4
153	Congenital multifocal rhabdoid tumor: a case with peculiar biological behavior and different response to treatment according to location (central nervous system and kidney). Cancer Genetics, 2014, 207, 441-444.	0.4	3
154	Reversible cerebral vasoconstriction mimicking posterior reversible encephalopathy syndrome in an infant with end-stage renal disease. Cephalalgia, 2015, 35, 1031-1033.	3.9	3
155	Crossed Pontine Hemiatrophy Associated with Unilateral Cerebellar Hemorrhage in Premature Infants. Neuropediatrics, 2016, 47, 404-407.	0.6	3
156	Neonatal Developmental Venous Anomalies: Clinicoradiologic Characterization and Follow-Up. American Journal of Neuroradiology, 2020, 41, 2370-2376.	2.4	3
157	Endocrine Outcomes In Central Diabetes Insipidus: the Predictive Value of Neuroimaging â€œMismatch Patternâ€. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 3562-3574.	3.6	3
158	Balamuthia mandrillaris infection: report of 1st autochthonous, fatal case in Italy. European Journal of Clinical Microbiology and Infectious Diseases, 2022, 41, 685-687.	2.9	3
159	Calcifications in diffuse leptomeningeal glioneuronal tumors: a case series. Quantitative Imaging in Medicine and Surgery, 2022, 12, 2985-2994.	2.0	3
160	A Phenotypic-Driven Approach for the Diagnosis of WOREE Syndrome. Frontiers in Pediatrics, 2022, 10, 847549.	1.9	3
161	A 3â€YEARâ€OLD BOY WITH DRUGâ€RESISTANT COMPLEX PARTIAL SEIZURES. Brain Pathology, 2012, 22, 725-728.	4.1	2
162	Constitutional chromosomal events at 22q11 and 15q26 in a child with a pilocytic astrocytoma of the spinal cord. Molecular Cytogenetics, 2014, 7, 31.	0.9	2

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