Nathalie Boddaert

List of Publications by Year
in descending order

132	3,799	147801	168389
papers	citations	31 h-index	53 g-index

144	144	144	6466
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Arterial Spin Labeling for the Etiological Workup of Intracerebral Hemorrhage in Children. Stroke, 2022, 53, 185-193.	2.0	6
2	A malignant choroid plexus tumour with prevailing immature blastematous elements. Neuropathology and Applied Neurobiology, 2022, 48, .	3.2	1
3	Deciphering the genetic and epigenetic landscape of pediatric bithalamic tumors. Brain Pathology, 2022, 32, e13039.	4.1	5
4	Treatment of two infants with PIK3CA-related overgrowth spectrum by alpelisib. Journal of Experimental Medicine, 2022, 219, .	8.5	27
5	A novel LARGE1-AFF2 fusion expanding the molecular alterations associated with the methylation class of neuroepithelial tumors with PATZ1 fusions. Acta Neuropathologica Communications, 2022, 10, 15.	5.2	3
6	A partial form of inherited human USP18 deficiency underlies infection and inflammation. Journal of Experimental Medicine, 2022, 219, .	8.5	28
7	Spectrum of Neuroradiologic Findings Associated with Monogenic Interferonopathies. American Journal of Neuroradiology, 2022, 43, 2-10.	2.4	6
8	Feasibility and Added Value of Fetal DTI Tractography in the Evaluation of an Isolated Short Corpus Callosum: Preliminary Results. American Journal of Neuroradiology, 2022, 43, 132-138.	2.4	7
9	Imaging Features with Histopathologic Correlation of CNS High-Grade Neuroepithelial Tumors with a BCOR Internal Tandem Duplication. American Journal of Neuroradiology, 2022, 43, 151-156.	2.4	17
10	Suleiman-El-Hattab syndrome: a histone modification disorder caused by TASP1 deficiency. Human Molecular Genetics, 2022, 31, 3083-3094.	2.9	3
11	Neuroinflammatory Disease following Severe Acute Respiratory Syndrome Coronavirus 2 Infection in Children. Journal of Pediatrics, 2022, 247, 22-28.e2.	1.8	15
12	The genomic landscape of dysembryoplastic neuroepithelial tumours and a comprehensive analysis of recurrent cases. Neuropathology and Applied Neurobiology, 2022, 48, .	3.2	4
13	Imaging features of medulloblastoma: Conventional imaging, diffusion-weighted imaging, perfusion-weighted imaging, and spectroscopy: From general features to subtypes and characteristics. Neurochirurgie, 2021, 67, 6-13.	1.2	19
14	Cerebral blood flow and acute episodes of Leigh syndrome in neurometabolic disorders. Developmental Medicine and Child Neurology, 2021, 63, 705-711.	2.1	6
15	Arterial abnormalities identified in kidneys transplanted into children during the COVID-19 pandemic. American Journal of Transplantation, 2021, 21, 1937-1943.	4.7	3
16	Hemorrhage Expansion After Pediatric Intracerebral Hemorrhage. Stroke, 2021, 52, 588-594.	2.0	4
17	Fastâ€track virtual reality for cardiac imaging in congenital heart disease. Journal of Cardiac Surgery, 2021, 36, 2598-2602.	0.7	21
18	Evolution of acute myocarditis in a pediatric population: An MRI based study. International Journal of Cardiology, 2021, 329, 226-233.	1.7	10

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19	Acute surgical management of children with ruptured brain arteriovenous malformation. Journal of Neurosurgery: Pediatrics, 2021, 27, 437-445.	1.3	2
20	Topographic variability of the normal circle of Willis anatomy on a paediatric population. Brain Communications, 2021, 3, fcab055.	3.3	0
21	Inherited deficiency of stress granule ZNFX1 in patients with monocytosis and mycobacterial disease. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	47
22	Radiogenomics of diffuse intrinsic pontine gliomas (DIPGs): correlation of histological and biological characteristics with multimodal MRI features. European Radiology, 2021, 31, 8913-8924.	4.5	11
23	Loss of function mutations in GEMIN5 cause a neurodevelopmental disorder. Nature Communications, 2021, 12, 2558.	12.8	28
24	Pineal alveolar rhabdomyosarcoma with PAX3:NCOA2 fusion inducing OLIG2 expression, a potential pitfall in the central nervous system. Histopathology, 2021, 79, 437-439.	2.9	3
25	A novel case of cribriform neuroepithelial tumor: A potential diagnostic pitfall in the ventricular system. Pediatric Blood and Cancer, 2021, 68, e29037.	1.5	3
26	A CBF decrease in the left supplementary motor areas: New insight into postoperative pediatric cerebellar mutism syndrome using arterial spin labeling perfusion MRI. Journal of Cerebral Blood Flow and Metabolism, 2021, 41, 3339-3349.	4.3	10
27	Fatal encephalitis caused by Newcastle disease virus in a child. Acta Neuropathologica, 2021, 142, 605-608.	7.7	9
28	Brain perfusion magnetic resonance imaging using pseudocontinuous arterial spin labeling in 314 dogs and cats. Journal of Veterinary Internal Medicine, 2021, 35, 2327-2341.	1.6	3
29	Alternative pathways for the development of lymphoid structures in humans. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	11
30	Pediatric brain arteriovenous malformation recurrence: a cohort study, systematic review and meta-analysis. Journal of NeuroInterventional Surgery, 2021, , neurintsurg-2021-017777.	3.3	10
31	Prenatalâ€onset of congenital neuronal ceroid lipofuscinosis with a novel CTSD mutation. Birth Defects Research, 2021, 113, 1324-1332.	1.5	1
32	Variants in the MIPEP gene presenting with complex neurological phenotype without cardiomyopathy, impair OXPHOS protein maturation and lead to a reduced OXPHOS abundance in patient cells. Molecular Genetics and Metabolism, 2021, 134, 267-273.	1.1	4
33	Rest Functional Brain Maturation during the First Year of Life. Cerebral Cortex, 2021, 31, 1776-1785.	2.9	11
34	Object Detection Improves Tumour Segmentation in MR Images of Rare Brain Tumours. Cancers, 2021, 13, 6113.	3.7	9
35	Prevalence of Venovenous Shunting and High-Output State Quantified with 4D Flow MRI in Patients with Fontan Circulation. Radiology: Cardiothoracic Imaging, 2021, 3, e210161.	2.5	5
36	Myocardial involvement in children with post-COVID multisystem inflammatory syndrome: a cardiovascular magnetic resonance based multicenter international studyâ€"the CARDOVID registry. Journal of Cardiovascular Magnetic Resonance, 2021, 23, 140.	3.3	33

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37	Arterial Spin Labeling and Central Precocious Puberty. Clinical Neuroradiology, 2020, 30, 137-144.	1.9	5
38	Biometric and morphological features on magnetic resonance imaging of fetal bladder in lower urinary tract obstruction: new perspectives for fetal cystoscopy. Ultrasound in Obstetrics and Gynecology, 2020, 56, 86-95.	1.7	6
39	Central nervous system complications in adult cystinosis patients. Journal of Inherited Metabolic Disease, 2020, 43, 348-356.	3.6	14
40	Scurvy: A New Old Cause of Skeletal Pain in Young Children. Frontiers in Pediatrics, 2020, 8, 8.	1.9	25
41	Posterior Fossa Arachnoid Cyst in a Pediatric Population is Associated with Social Perception and Rest Cerebral Blood Flow Abnormalities. Cerebellum, 2020, 19, 58-67.	2.5	2
42	The pediatric supratentorial MYCN-amplified high-grade gliomas methylation class presents the same radiological, histopathological and molecular features as their pontine counterparts. Acta Neuropathologica Communications, 2020, 8, 104.	5.2	24
43	Focal Areas of High Signal Intensity in Children with Neurofibromatosis Type 1: Expected Evolution on MRI. American Journal of Neuroradiology, 2020, 41, 1733-1739.	2.4	8
44	Complete hemispherotomy leads to lateralized functional organization and lower level of consciousness in the isolated hemisphere. Epilepsia Open, 2020, 5, 537-549.	2.4	3
45	Risk Factors for Early Brain AVM Rupture: Cohort Study of Pediatric and Adult Patients. American Journal of Neuroradiology, 2020, 41, 2358-2363.	2.4	16
46	Histone H3 wild-type DIPG/DMG overexpressing EZHIP extend the spectrum diffuse midline gliomas with PRC2 inhibition beyond H3-K27M mutation. Acta Neuropathologica, 2020, 139, 1109-1113.	7.7	104
47	Neonatal factors related to survival and intellectual and developmental outcome of patients with early-onset urea cycle disorders. Molecular Genetics and Metabolism, 2020, 130, 110-117.	1.1	4
48	Loss of Function of RIMS2 Causes a Syndromic Congenital Cone-Rod Synaptic Disease with Neurodevelopmental and Pancreatic Involvement. American Journal of Human Genetics, 2020, 106, 859-871.	6.2	22
49	High-grade gliomas in adolescents and young adults highlight histomolecular differences from their adult and pediatric counterparts. Neuro-Oncology, 2020, 22, 1190-1202.	1.2	50
50	The â€œsalt and pepperâ€•pattern on renal ultrasound in a group of children with molecular-proven diagnosis of ciliopathy-related renal diseases. Pediatric Nephrology, 2020, 35, 1033-1040.	1.7	10
51	Pediatric cardiac computed tomography angiography: Expert consensus from the Filiale de Cardiologie PÃ©diatrique et CongÃ©nitale (FCPC) and the SociÃ©tÃ© FranÃ§aise dâ€™Imagerie Cardiaque et Vasculaire diagnostique et interventionnelle (SFICV). Diagnostic and Interventional Imaging, 2020, 101, 335-345.	3.2	5
52	Imaging features of complete congenital atresia of left coronary artery. Diagnostic and Interventional Imaging, 2020, 101, 421-423.	3.2	2
53	Neural basis of interindividual variability in social perception in typically developing children and adolescents using diffusion tensor imaging. Scientific Reports, 2020, 10, 6379.	3.3	2
54	Hydrocephalus in children with ruptured cerebral arteriovenous malformation. Journal of Neurosurgery: Pediatrics, 2020, 26, 283-287.	1.3	2

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55	DIPG-61. RESCUE REGIMENS AFTER BIOMEDE: POSSIBLE INFLUENCE ON OS ASSESSMENT. Neuro-Oncology, 2020, 22, iii299-iii299.	1.2	0
56	Mutations in the MRPS28 gene encoding the small mitoribosomal subunit protein bS1m in a patient with intrauterine growth retardation, craniofacial dysmorphism and multisystemic involvement. Human Molecular Genetics, 2019, 28, 1445-1462.	2.9	19
57	Elevated thrombin generation in patients with congenital disorder of glycosylation and combined coagulation factor deficiencies. Journal of Thrombosis and Haemostasis, 2019, 17, 1798-1807.	3.8	18
58	Incidental Brain MRI Findings in Children: A Systematic Review and Meta-Analysis. American Journal of Neuroradiology, 2019, 40, 1818-1823.	2.4	25
59	Defects in t6A tRNA modification due to GON7 and YRDC mutations lead to Galloway-Mowat syndrome. Nature Communications, 2019, 10, 3967.	12.8	66
60	Neural and behavioral signature of human social perception. Scientific Reports, 2019, 9, 9252.	3.3	8
61	Aortic atresia and interrupted aortic arch communicating through external carotid anastomosis. Cardiology in the Young, 2019, 29, 699-700.	0.8	0
62	An integrative radiological, histopathological and molecular analysis of pediatric pontine histone-wildtype glioma with MYCN amplification (HGG-MYCN). Acta Neuropathologica Communications, 2019, 7, 87.	5.2	22
63	E-047â€...Vessel wall imaging and brain arteriovenous malformations: initial description of enhancement patterns. , 2019, , .		0
64	E-065â€...Ruptured brain arterio-venous malformations in children and adults: angioarchitectural variations at presentation across the lifespan. , 2019, , .		0
65	Corpus callosum metrics predict severity of visuospatial and neuromotor dysfunctions in ARID1B mutations with Coffinâ€“Siris syndrome. Psychiatric Genetics, 2019, 29, 237-242.	1.1	8
66	TP53 Pathway Alterations Drive Radioresistance in Diffuse Intrinsic Pontine Gliomas (DIPG). Clinical Cancer Research, 2019, 25, 6788-6800.	7.0	66
67	CT and Multimodal MR Imaging Features of Embryonal Tumors with Multilayered Rosettes in Children. American Journal of Neuroradiology, 2019, 40, 732-736.	2.4	9
68	High predictive value of brain MRI imaging in primary mitochondrial respiratory chain deficiency. Journal of Medical Genetics, 2018, 55, 378-383.	3.2	21
69	Historadiological correlations in high-grade glioma with the histone 3.3 G34R mutation. Journal of Neuroradiology, 2018, 45, 316-322.	1.1	26
70	Comprehensive molecular screening strategy of <scp>OCLN</scp> in bandâ€like calcification with simplified gyration and polymicrogyria. Clinical Genetics, 2018, 93, 228-234.	2.0	9
71	Predictors of Outcome in Patients with Pediatric Intracerebral Hemorrhage: Development and Validation of a Modified Score. Radiology, 2018, 286, 651-658.	7.3	31
72	Prenatally diagnosed periventricular nodular heterotopia: Further delineation of the imaging phenotype and outcome. European Journal of Medical Genetics, 2018, 61, 773-782.	1.3	8

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73	Reverse-Transcriptase Inhibitors in the Aicardiâ€“GoutiÃ¨res Syndrome. New England Journal of Medicine, 2018, 379, 2275-2277.	27.0	106
74	Inhibition of mitochondrial translation in fibroblasts from a patient expressing the KARS p.(Pro228Leu) variant and presenting with sensorineural deafness, developmental delay, and lactic acidosis. Human Mutation, 2018, 39, 2047-2059.	2.5	14
75	Further refinement of COL4A1 and COL4A2 related cortical malformations. European Journal of Medical Genetics, 2018, 61, 765-772.	1.3	29
76	Challenges in managing epilepsy associated with focal cortical dysplasia in children. Epilepsy Research, 2018, 145, 1-17.	1.6	25
77	PLA2G6-associated neurodegeneration: Lessons from neurophysiological findings. European Journal of Paediatric Neurology, 2018, 22, 854-861.	1.6	9
78	TLE1, a key player in neurogenesis, a new candidate gene for autosomal recessive postnatal microcephaly. European Journal of Medical Genetics, 2018, 61, 729-732.	1.3	8
79	NSRG-05. SAFETY OF ULTRASOUND-INDUCED BLOOD-BRAIN BARRIER OPENING IN PEDIATRIC PATIENTS WITH REFRACTORY SUS-TENTORIAL MALIGNANT BRAIN TUMORS BEFORE CHEMOTHERAPY ADMINISTRATION â€“ THE SONOKID CLINICAL TRIAL. Neuro-Oncology, 2018, 20, i146-i146.	1.2	2
80	Recurrent RTTN mutation leading to severe microcephaly, polymicrogyria and growth restriction. European Journal of Medical Genetics, 2018, 61, 755-758.	1.3	9
81	Myocardial inflammation detected by cardiac MRI in Arrhythmogenic right ventricular cardiomyopathy: A paediatric case series. International Journal of Cardiology, 2018, 271, 81-86.	1.7	52
82	Targeted therapy in patients with PIK3CA-related overgrowth syndrome. Nature, 2018, 558, 540-546.	27.8	374
83	Arterial Spin-Labeling to Discriminate Pediatric Cervicofacial Soft-Tissue Vascular Anomalies. American Journal of Neuroradiology, 2017, 38, 633-638.	2.4	20
84	Prenatal and postnatal presentations of corpus callosum agenesis with polymicrogyria caused by EGP5 mutation. American Journal of Medical Genetics, Part A, 2017, 173, 706-711.	1.2	12
85	Moyamoya syndrome in children with neurofibromatosis type 1: Italianâ€“French experience. American Journal of Medical Genetics, Part A, 2017, 173, 1521-1530.	1.2	36
86	Multimodal Magnetic Resonance Imaging of Treatment-Induced Changes to Diffuse Infiltrating Pontine Gliomas in Children and Correlation to Patient Progression-Free Survival. International Journal of Radiation Oncology Biology Physics, 2017, 99, 476-485.	0.8	18
87	Epileptic spasms in congenital disorders of glycosylation. Epileptic Disorders, 2017, 19, 15-23.	1.3	12
88	A novel recurrent LIS1 splice site mutation in classic lissencephaly. American Journal of Medical Genetics, Part A, 2017, 173, 561-564.	1.2	6
89	Clinical, laboratory and molecular findings and long-term follow-up data in 96 French patients with PMM2-CDG (phosphomannomutase 2-congenital disorder of glycosylation) and review of the literature. Journal of Medical Genetics, 2017, 54, 843-851.	3.2	88
90	WDR81 mutations cause extreme microcephaly and impair mitotic progression in human fibroblasts and Drosophila neural stem cells. Brain, 2017, 140, 2597-2609.	7.6	28

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91	Fetal MRI compared with ultrasound for the diagnosis of obstructive genital malformations. Prenatal Diagnosis, 2017, 37, 1138-1145.	2.3	14
92	Biallelic Mutations in LIPT2 Cause a Mitochondrial Lipoylation Defect Associated with Severe Neonatal Encephalopathy. American Journal of Human Genetics, 2017, 101, 283-290.	6.2	55
93	Severe neuroimaging anomalies are usually associated with random X inactivation in leucocytes circulating DNA in X-linked dominant Incontinentia Pigmenti. Molecular Genetics and Metabolism, 2017, 122, 140-144.	1.1	11
94	Usefulness of cocaine drops in investigating infant anisocoria. European Journal of Paediatric Neurology, 2017, 21, 852-857.	1.6	12
95	Neuroimaging evidence of brain abnormalities in mastocytosis. Translational Psychiatry, 2017, 7, e1197-e1197.	4.8	21
96	Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly. Nature Genetics, 2017, 49, 1529-1538.	21.4	164
97	Biallelic Mutations in MRPS34 Lead to Instability of the Small Mitoribosomal Subunit and Leigh Syndrome. American Journal of Human Genetics, 2017, 101, 239-254.	6.2	83
98	Mutations in TUBB4B Cause a Distinctive Sensorineural Disease. American Journal of Human Genetics, 2017, 101, 1006-1012.	6.2	30
99	Neuropsychological improvement after posterior fossa arachnoid cyst drainage. Child's Nervous System, 2017, 33, 135-141.	1.1	17
100	Type I interferon-mediated autoinflammation due to DNase II deficiency. Nature Communications, 2017, 8, 2176.	12.8	164
101	HG-46RECURRENT DIFFUSE INTRINSIC PONTINE GLIOMAS: CLINICAL, BIOLOGICAL, RADIOLOGICAL AND THERAPEUTIC FACTORS CORRELATING WITH THE SURVIVAL. Neuro-Oncology, 2016, 18, iii57.4-iii58.	1.2	0
102	Magnetic resonance imaging arterialâ€spinâ€labelling perfusion alterations in childhood migraine with atypical aura: a caseâ€“control study. Developmental Medicine and Child Neurology, 2016, 58, 965-969.	2.1	26
103	Cerebral Blood Flow Improvement after Indirect Revascularization for Pediatric Moyamoya Disease: A Statistical Analysis of Arterial Spin-Labeling MRI. American Journal of Neuroradiology, 2016, 37, 706-712.	2.4	41
104	Arterial spin labeling shows pre-epileptic tuber hyperperfusion in tuberous sclerosis complex. Neurology, 2016, 86, 1744-1745.	1.1	4
105	Arterial Spin Labeling to Predict Brain Tumor Grading in Children: Correlations between Histopathologic Vascular Density and Perfusion MR Imaging. Radiology, 2016, 281, 553-566.	7.3	82
106	Recurrent KIF5C mutation leading to frontal pachygyria without microcephaly. Neurogenetics, 2016, 17, 79-82.	1.4	17
107	Tuning Eye-Gaze Perception by Transitory STS Inhibition. Cerebral Cortex, 2016, 26, 2823-2831.	2.9	19
108	A nonsense variant in HERC1 is associated with intellectual disability, megalencephaly, thick corpus callosum and cerebellar atrophy. European Journal of Human Genetics, 2016, 24, 455-458.	2.8	53

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109	ISDN2014_0400: Mutations in DOCK7 in individuals with epileptic encephalopathy and cortical blindness. International Journal of Developmental Neuroscience, 2015, 47, 119-120.	1.6	0
110	Choroid Plexus Neoplasms: Toward a Distinction between Carcinoma and Papilloma Using Arterial Spin-Labeling. American Journal of Neuroradiology, 2015, 36, 1786-1790.	2.4	31
111	Contiguous mutation syndrome in the era of highâ€throughput sequencing. Molecular Genetics & Genomic Medicine, 2015, 3, 215-220.	1.2	9
112	Mutations in NONO lead to syndromic intellectual disability and inhibitory synaptic defects. Nature Neuroscience, 2015, 18, 1731-1736.	14.8	65
113	Arterial spin labeling magnetic resonance imaging: toward noninvasive diagnosis and follow-up of pediatric brain arteriovenous malformations. Journal of Neurosurgery: Pediatrics, 2015, 15, 451-458.	1.3	35
114	From splitting GLUT1 deficiency syndromes to overlapping phenotypes. European Journal of Medical Genetics, 2015, 58, 443-454.	1.3	52
115	Myocardial inflammation on cardiovascular magnetic resonance predicts left ventricular function recovery in children with recent dilated cardiomyopathy. European Heart Journal Cardiovascular Imaging, 2015, 16, 756-762.	1.2	15
116	IFT81, encoding an IFT-B core protein, as a very rare cause of a ciliopathy phenotype. Journal of Medical Genetics, 2015, 52, 657-665.	3.2	32
117	A neuropathological study of cerebrovascular abnormalities in a signal transducer and activator of transcription 3â€“deficient patient. Journal of Allergy and Clinical Immunology, 2015, 136, 1418-1421.e5.	2.9	5
118	Arterial Spin Labeling MRI: A step forward in non-invasive delineation of focal cortical dysplasia in children. Epilepsy Research, 2014, 108, 1932-1939.	1.6	46
119	Loss-of-Function Mutations in WDR73 Are Responsible for Microcephaly and Steroid-Resistant Nephrotic Syndrome: Galloway-Mowat Syndrome. American Journal of Human Genetics, 2014, 95, 637-648.	6.2	108
120	Long-Term Outcome of 106 Consecutive Pediatric Ruptured Brain Arteriovenous Malformations After Combined Treatment. Stroke, 2014, 45, 1664-1671.	2.0	86
121	Recurrent Streptococcus pyogenes genital infection in a woman: test and treat the partner!. International Journal of Infectious Diseases, 2014, 29, 37-39.	3.3	8
122	Mutations in QARS, Encoding Glutaminyl-tRNA Synthetase, Cause Progressive Microcephaly, Cerebral-Cerebellar Atrophy, and Intractable Seizures. American Journal of Human Genetics, 2014, 94, 547-558.	6.2	106
123	Early epileptic encephalopathies associated with STXBP1 mutations: Could we better delineate the phenotype?. European Journal of Medical Genetics, 2014, 57, 15-20.	1.3	50
124	Mutations in DOCK7 in Individuals with Epileptic Encephalopathy and Cortical Blindness. American Journal of Human Genetics, 2014, 94, 891-897.	6.2	44
125	Brain imaging in mitochondrial respiratory chain deficiency: combination of brain MRI features as a useful tool for genotype/phenotype correlations. Journal of Medical Genetics, 2014, 51, 429-435.	3.2	40
126	Clinical and imaging diagnosis for heredodegenerative diseases. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2013, 111, 63-78.	1.8	124

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127	Massive and exclusive pontocerebellar damage in mitochondrial disease and NUBPL mutations. Neurology, 2012, 79, 391-391.	1.1	27
128	Mutation in PNPT1 , which Encodes a Polyribonucleotide Nucleotidyltransferase, Impairs RNA Import into Mitochondria and Causes Respiratory-Chain Deficiency. American Journal of Human Genetics, 2012, 91, 912-918.	6.2	81
129	Childhood gangliogliomas with ependymal differentiation. Neuropathology and Applied Neurobiology, 2009, 35, 437-441.	3.2	3
130	Multiple bur hole surgery for the treatment of moyamoya disease in children. Journal of Neurosurgery: Pediatrics, 2006, 105, 437-443.	1.3	37
131	Atypical case of hemiconvulsionsâ€hemiplegiaâ€epilepsy syndrome revealing contralateral focal cortical dysplasia. Developmental Medicine and Child Neurology, 2005, 47, 830-834.	2.1	1
132	Functional neuroimaging and childhood autism. Pediatric Radiology, 2002, 32, 1-7.	2.0	107