Luisa Chiapparini

List of Publications by Year
in descending order

149	4,504	109137	128067
papers	citations	35 h-index	60 g-index

153	153	153	6277
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Age-related Iron Deposition in the Basal Ganglia: Quantitative Analysis in Healthy Subjects. Radiology, 2009, 252, 165-172.	3.6	266
2	Antiâ€“amyloid Î² autoantibodies in cerebral amyloid angiopathyâ€“related inflammation: Implications for amyloidâ€modifying therapies. Annals of Neurology, 2013, 73, 449-458.	2.8	179
3	Exome Sequence Reveals Mutations in CoA Synthase as a Cause of Neurodegeneration with Brain Iron Accumulation. American Journal of Human Genetics, 2014, 94, 11-22.	2.6	176
4	Stimulation of the globus pallidus internus for childhood-onset dystonia. Movement Disorders, 2005, 20, 1194-1200.	2.2	162
5	Periventricular Nodular Heterotopia: Classification, Epileptic History, and Genesis of Epileptic Discharges. Epilepsia, 2006, 47, 86-97.	2.6	150
6	Spinal radiological findings in nine patients with spontaneous intracranial hypotension. Neuroradiology, 2002, 44, 143-150.	1.1	139
7	Rasmussenâ€™s encephalitis. Neurology, 2003, 60, 422-425.	1.5	137
8	Overlapping phenotypes in complex spastic paraplegias SPG11, SPG15, SPG35 and SPG48. Brain, 2014, 137, 1907-1920.	3.7	133
9	Iron-related MRI images in patients with pantothenate kinase-associated neurodegeneration (PKAN) treated with deferiprone: Results of a phase II pilot trial. Movement Disorders, 2011, 26, 1755-1759.	2.2	125
10	Diagnostic imaging in 13Âcases of Rasmussen's encephalitis: can early MRI suggest the diagnosis?. Neuroradiology, 2003, 45, 171-183.	1.1	116
11	Adult-Onset Rasmussen's Encephalitis: Anatomical-Electrographic-Clinical Features of 7 Italian Cases. Epilepsia, 2006, 47, 41-46.	2.6	108
12	Diffusion Tensor Imaging Shows Different Topographic Involvement of the Thalamus in Progressive Supranuclear Palsy and Corticobasal Degeneration. American Journal of Neuroradiology, 2009, 30, 1482-1487.	1.2	105
13	Can MR Imaging Diagnose Adult-Onset Alexander Disease?. American Journal of Neuroradiology, 2008, 29, 1190-1196.	1.2	99
14	Spontaneous intracranial hypotension with deep brain swelling. Brain, 2007, 130, 1884-1893.	3.7	98
15	Spontaneous intracranial hypotension syndrome: a novel speculative physiopathological hypothesis and a novel patch method in a series of 28 consecutive patients. Journal of Neurosurgery, 2010, 112, 300-306.	0.9	93
16	Pain Processing in Medication Overuse Headache: A Functional Magnetic Resonance Imaging (fMRI) Study. Pain Medicine, 2012, 13, 255-262.	0.9	74
17	Chronic Migraine With Medication Overuse Preâ€“Post Withdrawal of Symptomatic Medication: Clinical Results and fMRI Correlations. Headache, 2010, 50, 998-1004.	1.8	68
18	Response assessment in diffuse intrinsic pontine glioma: recommendations from the Response Assessment in Pediatric Neuro-Oncology (RAPNO) working group. Lancet Oncology, The, 2020, 21, e330-e336.	5.1	59

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19	In Medicationâ€Overuse Headache, <scp>fMRI</scp> Shows Longâ€Lasting Dysfunction in Midbrain Areas. Headache, 2012, 52, 1520-1534.	1.8	58
20	Iatrogenic intracranial pseudoaneurysms: neuroradiological and therapeutical considerations, including endovascular options. Neurological Sciences, 2006, 27, 317-322.	0.9	56
21	Dural sinus thrombosis in spontaneous intracranial hypotension. Journal of Neurology, 2006, 253, 1197-1202.	1.8	51
22	Substantia nigra in Parkinsonâ€™s disease: a multimodal MRI comparison between early and advanced stages of the disease. Neurological Sciences, 2014, 35, 753-758.	0.9	50
23	Headache and intracranial hypotension: neuroradiological findings. Neurological Sciences, 2004, 25, s138-s141.	0.9	48
24	Epileptic and imaging findings in perinatal hypoxic-ischemic encephalopathy with ulegyria. Epilepsy Research, 2003, 55, 235-243.	0.8	45
25	Hypomorphic Recessive Variants in SUFU Impair the Sonic Hedgehog Pathway and Cause Joubert Syndrome with Cranio-facial and Skeletal Defects. American Journal of Human Genetics, 2017, 101, 552-563.	2.6	45
26	Hyperhomocysteinemia and other thrombophilic risk factors in 26 patients with cerebral venous thrombosis. European Journal of Neurology, 2004, 11, 405-409.	1.7	44
27	Angiographically proven cervical venous engorgement: a possible concurrent cause in the pathophysiology of Hirayamaâ€™s myelopathy. Neurological Sciences, 2010, 31, 845-848.	0.9	44
28	Pontine tegmental cap dysplasia: developmental and cognitive outcome in three adolescent patients. Orphanet Journal of Rare Diseases, 2011, 6, 36.	1.2	44
29	Application of IHS Criteria to Headache Attributed to Spontaneous Intracranial Hypotension in a Large Population. Cephalalgia, 2009, 29, 418-422.	1.8	43
30	Spontaneous low cerebrospinal pressure: a mini review. Neurological Sciences, 2004, 25, s135-s137.	0.9	42
31	Electroclinical spectrum of the neuronal ceroid lipofuscinoses associated with CLN6 mutations. Neurology, 2015, 85, 316-324.	1.5	40
32	Clinical spectrum of PTEN mutation in pediatric patients. A bicenter experience. European Journal of Medical Genetics, 2019, 62, 103596.	0.7	39
33	Neuroimaging in chronic migraine. Neurological Sciences, 2010, 31, 19-22.	0.9	38
34	C19orf12 and FA2H Mutations Are Rare in Italian Patients With Neurodegeneration With Brain Iron Accumulation. Seminars in Pediatric Neurology, 2012, 19, 75-81.	1.0	38
35	Imaging and outcome in severe complications of lumbar epidural anaesthesia: report of 16 cases. Neuroradiology, 2000, 42, 564-571.	1.1	37
36	Clinical features and outcomes in spontaneous intracranial hypotension: a survey of 90 consecutive patients. Neurological Sciences, 2009, 30, 11-13.	0.9	37

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37	Brownâ€“Vialettoâ€“van Laere and Fazioâ€“Londe overlap syndromes: A clinical, biochemical and genetic study. Neuromuscular Disorders, 2012, 22, 1075-1082.	0.3	36
38	Aicardiâ€GoutiÃ¨res syndrome: description of a late onset case. Developmental Medicine and Child Neurology, 2008, 50, 631-634.	1.1	35
39	Hemophagocytic lymphohistiocytosis with neurological presentation: MRI findings and a nearly miss diagnosis. Neurological Sciences, 2011, 32, 473-477.	0.9	35
40	Neuroradiological diagnosis of Chiari malformations. Neurological Sciences, 2011, 32, 283-286.	0.9	34
41	The â€œEye-of-the-Tigerâ€•Sign may be Absent in the Early Stages of Classic Pantothenate Kinase Associated Neurodegeneration. Neuropediatrics, 2011, 42, 159-162.	0.3	34
42	Low-Functioning Autism and Nonsyndromic Intellectual Disability. Journal of Child Neurology, 2015, 30, 1658-1663.	0.7	32
43	KARS-related diseases: progressive leukoencephalopathy with brainstem and spinal cord calcifications as new phenotype and a review of literature. Orphanet Journal of Rare Diseases, 2018, 13, 45.	1.2	32
44	Treatment of spontaneous intracranial hypotension: evolution of the therapeutic and diagnostic modalities. Neurological Sciences, 2013, 34, 151-155.	0.9	29
45	Clinical, radiological and possible pathological overlap of cystic leukoencephalopathy without megalencephaly and Aicardi-GoutiÃ¨res syndrome. European Journal of Paediatric Neurology, 2016, 20, 604-610.	0.7	29
46	Molecular Genetics and Interferon Signature in the Italian Aicardi GoutiÃ¨res Syndrome Cohort: Report of 12 New Cases and Literature Review. Journal of Clinical Medicine, 2019, 8, 750.	1.0	29
47	Cranial ultrasound is a reliable first step imaging in children with suspected craniosynostosis. Child's Nervous System, 2017, 33, 1545-1552.	0.6	27
48	Iatrogenic early onset cerebral amyloid angiopathy 30â€‰years after cerebral trauma with neurosurgery: vascular amyloid deposits are made up of both Aî²40 and Aî²42. Acta Neuropathologica Communications, 2019, 7, 70.	2.4	26
49	POEMS syndrome: relapse after successful autologous peripheral blood stem cell transplantation. Neuromuscular Disorders, 2007, 17, 980-982.	0.3	25
50	Treatment for Chiari 1 malformation (CIM): analysis of a pediatric surgical series. Neurological Sciences, 2011, 32, 321-324.	0.9	25
51	Defective functional connectivity between posterior hypothalamus and regions of the diencephalic-mesencephalic junction in chronic cluster headache. Cephalalgia, 2018, 38, 1910-1918.	1.8	25
52	Electroencephalographic Recordings of Focal Seizures in Patients Affected by Periventricular Nodular Heterotopia: Role of the Heterotopic Nodules in the Genesis of Epileptic Discharges. Journal of Child Neurology, 2005, 20, 369-377.	0.7	24
53	Second series by the Italian Association of Pediatric Hematology and Oncology of children and adolescents with intracranial ependymoma: an integrated molecular and clinical characterization with a long-term follow-up. Neuro-Oncology, 2021, 23, 848-857.	0.6	24
54	Diagnosis and treatment of Chiari malformation type 1 in children: the International Consensus Document. Neurological Sciences, 2022, 43, 1311-1326.	0.9	24

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55	Neurological Disorders Associated with Striatal Lesions: Classification and Diagnostic Approach. Current Neurology and Neuroscience Reports, 2016, 16, 54.	2.0	23
56	Chiari malformation-related headache: outcome after surgical treatment. Neurological Sciences, 2017, 38, 95-98.	0.9	23
57	Chiari malformation type 1-related headache: the importance of a multidisciplinary study. Neurological Sciences, 2017, 38, 91-93.	0.9	23
58	Ataxia With Oculomotor Apraxia Type 1 (AOA1): Clinical and Neuropsychological Features in 2 New Patients and Differential Diagnosis. Journal of Child Neurology, 2008, 23, 895-900.	0.7	22
59	A Case of Pediatric Tumefactive Demyelinating Lesion Misdiagnosed and Treated as Glioblastoma. Journal of Child Neurology, 2008, 23, 944-947.	0.7	22
60	Functional-MRI evaluation of pain processing in chronic migraine with medication overuse. Neurological Sciences, 2009, 30, 71-74.	0.9	22
61	The Italian Alzheimer's Disease Neuroimaging Initiative (I-ADNI): Validation of Structural MR Imaging. Journal of Alzheimer's Disease, 2014, 40, 941-952.	1.2	22
62	Late presentation of leucoencephalopathy with calcifications and cysts: report of two cases. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 1303-1304.	0.9	21
63	Xâ€linked Charcotâ€Marieâ€Tooth type 1: strokeâ€like presentation of a novel <scp>GJB1</scp> mutation. Journal of the Peripheral Nervous System, 2014, 19, 183-186.	1.4	20
64	Substantia Nigra Swelling and Dentate Nucleus T2 Hyperintensity May Be Early Magnetic Resonance Imaging Signs of Î²â€Propeller Proteinâ€Associated Neurodegeneration. Movement Disorders Clinical Practice, 2019, 6, 51-56.	0.8	20
65	Treatment of alterations in CSF dynamics. Neurological Sciences, 2011, 32, 117-120.	0.9	19
66	Chronic daily headache in the adults: differential diagnosis between symptomatic Chiari I malformation and spontaneous intracranial hypotension. Neurological Sciences, 2011, 32, 291-294.	0.9	19
67	Dandy-Walker malformation and Wisconsin syndrome: novel cases add further insight into the genotype-phenotype correlations of 3q23q25 deletions. Orphanet Journal of Rare Diseases, 2013, 8, 75.	1.2	19
68	Neuroimaging Findings in 41 Low-Functioning Children With Autism Spectrum Disorder. Journal of Child Neurology, 2014, 29, 1626-1631.	0.7	19
69	Risk of Optic Pathway Glioma in Neurofibromatosis Type 1: No Evidence of Genotypeâ€“Phenotype Correlations in a Large Independent Cohort. Cancers, 2019, 11, 1838.	1.7	19
70	Harmful Iron-Calcium Relationship in Pantothenate kinase Associated Neurodegeneration. International Journal of Molecular Sciences, 2020, 21, 3664.	1.8	19
71	Phenotypic spectrum of short-chain enoyl-Coa hydratase-1 (ECHS1) deficiency. European Journal of Paediatric Neurology, 2020, 28, 151-158.	0.7	18
72	Lumbar epidural blood patch: effectiveness on orthostatic headache and MRI predictive factors in 101 consecutive patients affected by spontaneous intracranial hypotension. Journal of Neurosurgery, 2020, 132, 809-817.	0.9	18

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73	Headache attributed to spontaneous intracranial hypotension. Neurological Sciences, 2008, 29, 164-165.	0.9	17
74	Neurological pictures in Paediatric Chiari I malformation. Neurological Sciences, 2011, 32, 295-298.	0.9	17
75	Early-onset progressive spastic paraplegia caused by a novel TUBB4A mutation: brain MRI and FDG-PET findings. Journal of Neurology, 2016, 263, 591-593.	1.8	17
76	Adult leukoencephalopathies with prominent infratentorial involvement can be caused by Erdheimâ€“Chester disease. Journal of Neurology, 2018, 265, 273-284.	1.8	17
77	Encephalopathies with intracranial calcification in children: clinical and genetic characterization. Orphanet Journal of Rare Diseases, 2018, 13, 135.	1.2	17
78	Spontaneous intracranial hypotension: diagnostic and therapeutic implications in neurosurgical practice. Neurological Sciences, 2011, 32, 287-290.	0.9	16
79	Progressive myoclonus epilepsy caused by a gain-of-function KCNA2 mutation. Seizure: the Journal of the British Epilepsy Association, 2019, 65, 106-108.	0.9	16
80	<scp>YY1</scp>â€Related Dystonia: Clinical Aspects and Longâ€Term Response to Deep Brain Stimulation. Movement Disorders, 2021, 36, 1461-1462.	2.2	16
81	Rhythmic cortical myoclonus in a case of HIV-related encephalopathy. Movement Disorders, 2003, 18, 1533-1538.	2.2	14
82	Prolonged Focal Negative Motor Seizures: A Video-EEG Study. Epilepsia, 2006, 47, 1949-1952.	2.6	14
83	Chiari I malformation in a child with PTEN hamartoma tumor syndrome: Association or coincidence?. European Journal of Medical Genetics, 2017, 60, 261-264.	0.7	14
84	A PDE10A de novo mutation causes childhoodâ€onset chorea with diurnal fluctuations. Movement Disorders, 2017, 32, 1646-1647.	2.2	13
85	Thiamine-responsive disease due to mutation of tpk1 : Importance of avoiding misdiagnosis. Neurology, 2017, 89, 870-871.	1.5	13
86	MYORG -related disease is associated with central pontine calcifications and atypical parkinsonism. Neurology: Genetics, 2020, 6, e399.	0.9	13
87	Consolidating the Role of TDP2 Mutations in Recessive Spinocerebellar Ataxia Associated with Pediatric Onset Drug Resistant Epilepsy and Intellectual Disability (SCAR23). Cerebellum, 2019, 18, 972-975.	1.4	12
88	Novel mutations in SLC16A2 associated with a less severe phenotype of MCT8 deficiency. Metabolic Brain Disease, 2019, 34, 1565-1575.	1.4	12
89	Quantitative Muscle MRI Protocol as Possible Biomarker in Becker Muscular Dystrophy. Clinical Neuroradiology, 2021, 31, 257-266.	1.0	12
90	Impact of obesity and binge eating disorder on patients with idiopathic intracranial hypertension. Cephalalgia, 2017, 37, 278-283.	1.8	11

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91	Headache frequency and symptoms of depression as predictors of disability in patients with idiopathic intracranial hypertension. Neurological Sciences, 2018, 39, 139-140.	0.9	11
92	Chiari 1 malformation and untreated sagittal synostosis: a new subset of complex Chiari?. Child's Nervous System, 2019, 35, 1741-1753.	0.6	11
93	Biallelic mutations in RNF220 cause laminopathies featuring leukodystrophy, ataxia and deafness. Brain, 2021, 144, 3020-3035.	3.7	11
94	Type II neurofibromatosis presenting as quadriceps atrophy. Italian Journal of Neurological Sciences, 1998, 19, 94-96.	0.1	10
95	Headache and spontaneous low cerebrospinal fluid pressure syndrome. Neurological Sciences, 2007, 28, S232-S234.	0.9	10
96	Intrathecal synthesis of onconeural antibodies in patients with paraneoplastic syndromes. Journal of Neuroimmunology, 2016, 290, 119-122.	1.1	10
97	Spontaneous MRI improvement and absence of cerebral calcification in Aicardi-GoutiÃ¨res syndrome: Diagnostic and disease-monitoring implications. Molecular Genetics and Metabolism, 2019, 126, 489-494.	0.5	10
98	Cluster headache: insights from resting-state functional magnetic resonance imaging. Neurological Sciences, 2019, 40, 45-47.	0.9	10
99	Therapeutic Advances in Neurodegeneration With Brain Iron Accumulation. Seminars in Pediatric Neurology, 2012, 19, 82-86.	1.0	9
100	Persistent Fixed Torticollis due to Atlanto-Axial Rotatory Fixation: Report of 4 Pediatric Cases. Neuropediatrics, 2005, 36, 45-49.	0.3	8
101	Spontaneous CSF leak treated with percutaneous CT-guided fibrin glue. Neurology, 2006, 66, 782-782.	1.5	8
102	Intracranial idiopathic hypertension: 1-year follow-up study. Neurological Sciences, 2014, 35, 177-179.	0.9	8
103	Resting state fMRI in cluster headache: which role?. Neurological Sciences, 2015, 36, 47-50.	0.9	8
104	Headache in spontaneous intracranial hypotension: an overview with indications for differential diagnosis in the clinical practice. Neurological Sciences, 2020, 41, 423-427.	0.9	8
105	Understanding Cluster Headache Using Magnetic Resonance Imaging. Frontiers in Neurology, 2020, 11, 535.	1.1	8
106	Artificial lamina-assisted laminoplasty performed in seven cases. Journal of Neurosurgery: Spine, 1999, 91, 43-49.	0.9	7
107	Paediatric Stroke: Review of the Literature and Possible Treatment Options, including Endovascular Approach. Stroke Research and Treatment, 2011, 2011, 1-11.	0.5	7
108	Pediatric intracranial ependymoma: correlating signs and symptoms at recurrence with outcome in the second prospective AIEOP protocol follow-up. Journal of Neuro-Oncology, 2018, 140, 457-465.	1.4	7

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109	A novel homozygous ISPD gene mutation causing phenotype variability in a consanguineous family. Neuromuscular Disorders, 2015, 25, 55-59.	0.3	6
110	SLC19A3 related disorder: Treatment implication and clinical outcome of 2 new patients. European Journal of Paediatric Neurology, 2018, 22, 332-335.	0.7	6
111	Abnormal cerebellar foliation in EBF3 mutation. Neurology, 2020, 94, 933-935.	1.5	6
112	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.5	6
113	In vivo brain MR spectroscopy in gliomas: clinical and pre-clinical chances. Clinical and Translational Imaging, 2022, 10, 495-515.	1.1	6
114	Autosomal Dominant Spinocerebellar Ataxias and Episodic Ataxias. , 2013, , 2193-2267.		5
115	Milder presentation of TELO2-related syndrome in two sisters homozygous for the p.Arg609His pathogenic variant. European Journal of Medical Genetics, 2021, 64, 104116.	0.7	5
116	Surgical treatment and clinical outcome of large pediatric interhemispheric cysts with callosal agenesis: A systematic literature review with four additional patients. Clinical Neurology and Neurosurgery, 2021, 204, 106600.	0.6	5
117	Medulloblastoma and familial adenomatous polyposis: Good prognosis and good quality of life in the longâ€term?. Pediatric Blood and Cancer, 2021, 68, e28912.	0.8	5
118	Treatment and outcome of intracranial ependymoma after first relapse in the 2nd AIEOP protocol. Neuro-Oncology, 2022, 24, 467-479.	0.6	5
119	Sagging and swelling of the midbrain suggest spontaneous intracranial hypotension rather than a malformation. Brain, 2010, 133, e148-e148.	3.7	4
120	Conventional MRI. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 154, 219-234.	1.0	4
121	MRI in Late-Onset Rasmussen Encephalitis: A Long-Term Follow-Up Study. Diagnostics, 2022, 12, 502.	1.3	4
122	Mesocorticolimbic system abnormalities in chronic cluster headache patients: A neural signature?. Cephalalgia, 2022, 42, 1039-1049.	1.8	4
123	Severe Pain and Edema due to a Widespread Lymphangioma: Disappearance of Symptoms and Reduction of Lesion with Spinal Cord Stimulation. World Neurosurgery, 2016, 93, 487.e1-487.e3.	0.7	3
124	Discovering the Italian phenotype of cerebral amyloid angiopathy (CAA): the SENECA project. Neurological Sciences, 2020, 41, 2193-2200.	0.9	3
125	<scp>THAP1</scp> Dystonia with Globus Pallidus <scp>T2</scp> Hypointensity: A Report of Two Cases. Movement Disorders, 2021, 36, 1463-1464.	2.2	3
126	A Missense De Novo Variant in the CASK-interactor KIRREL3 Gene Leading to Neurodevelopmental Disorder with Mild Cerebellar Hypoplasia. Neuropediatrics, 2021, 52, 484-488.	0.3	3

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127	The Pre-Lumbar puncture Intracranial Hypertension Scale (PLIHS): A practical scale to identify subjects with normal cerebrospinal fluid pressure in the management of idiopathic intracranial hypertension. Journal of the Neurological Sciences, 2021, 429, 118058.	0.3	3
128	Trattamento combinato con stent e polimero liquido (ONYXÂ®) in un caso di aneurisma gigante intracranico. The Neuroradiology Journal, 2001, 14, 231-234.	0.1	2
129	Persistent orthostatic headache without intracranial hypotension: which treatment?. Neurological Sciences, 2015, 36, 153-155.	0.9	2
130	A 52â€Yearâ€Old Man with Myoclonic Jerks. Brain Pathology, 2016, 26, 291-292.	2.1	2
131	Epileptic spikes in Rasmussenâ€™s encephalitis: Migratory pattern and short-term evolution. A MEG study. Clinical Neurophysiology, 2017, 128, 1898-1905.	0.7	2
132	The epileptology of Aicardi-GoutiÃ¨res syndrome: electro-clinical-radiological findings. Seizure: the Journal of the British Epilepsy Association, 2021, 86, 197-209.	0.9	2
133	Spinal cord lesion due to epidural anesthesia. Neurological Sciences, 2000, 21, 411-412.	0.9	1
134	Low-signal intensity on MRI in the globus pallidus in children with long-term oral manganese exposure. NeuroToxicology, 2017, 58, 92-93.	1.4	1
135	Sleeve-Shaped Neurothekeoma of the Ulnar Nerve: A Unique Case of a Still Unclear Pathological Entity. Hand, 2020, 15, NP7-NP10.	0.7	1
136	How to look for intracranial calcification in children with neurological disorders: CT, MRI, or both of them?. Neurological Sciences, 2022, 43, 2043-2050.	0.9	1
137	Neuroimaging of Pediatric Cerebellum in Inherited Neurodegenerative Diseases. Applied Sciences (Switzerland), 2021, 11, 8522.	1.3	1
138	Boomerang-shaped hemorrhage of the splenium of corpus callosum after mild brain injury: an unusual case of delayed hematoma. Journal of Neurosurgical Sciences, 2019, 63, 611-612.	0.3	1
139	Kearns-Sayre syndrome: expanding spectrum of a â€œnovelâ€•mitochondrial leukomyeloencephalopathy. Neurological Sciences, 2022, 43, 2081.	0.9	1
140	Familiar protein C deficiency and cerebral vein thrombosis in a young adult patient. Italian Journal of Neurological Sciences, 1999, 20, 333-334.	0.1	0
141	Pseudotumor Cerebri dell'etÃ infantile e adolescenziale. The Neuroradiology Journal, 2003, 16, 497-501.	0.1	0
142	Adult-onset leukoencephalopathy with calcifications associated with primary antiphospholipid syndrome. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 1215-1217.	0.9	0
143	Pearls & Oy-sters: The use of CT venography in Hirayama disease. Neurology, 2013, 80, 1539-1539.	1.5	0
144	Exploring cerebral networks in cluster headache: Insights and perspectives. Cephalalgia, 2014, 34, 323-324.	1.8	0

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145	The importance of specific rehabilitation for an obese patient with idiopathic intracranial hypertension: a case report. International Journal of Rehabilitation Research, 2018, 41, 183-185.	0.7	0
146	Relevance of neurophysiological assessment in a case of epilepsia partialis continua caused by anaplastic large cell lymphoma. Clinical Neurophysiology, 2021, 132, 165-166.	0.7	0
147	Sindrome da ipotensione liquorale spontanea. , 2009, , 343-345.		0
148	Neurodegeneration with Brain Iron Accumulation. , 2014, , 171-198.		0
149	Superficial siderosis in long-standing pilocytic astrocytoma. Neurological Sciences, 2022, , 1.	0.9	0