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	Diagnosis and treatment of Chiari malformation type 1 in children: the International Consensus Document. Neurological Sciences, 2022, 43, 1311-1326.	0.9	24
2	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
3	Treatment and outcome of intracranial ependymoma after first relapse in the 2nd AIEOP protocol. Neuro-Oncology, 2022, 24, 467-479.	0.6	5
4	Kearns-Sayre syndrome: expanding spectrum of a â€œnovelâ€•mitochondrial leukomyeloencephalopathy. Neurological Sciences, 2022, 43, 2081.	0.9	1
5	MRI in Late-Onset Rasmussen Encephalitis: A Long-Term Follow-Up Study. Diagnostics, 2022, 12, 502.	1.3	4
6	Superficial siderosis in long-standing pilocytic astrocytoma. Neurological Sciences, 2022, , 1.	0.9	0
7	Mesocorticolimbic system abnormalities in chronic cluster headache patients: A neural signature?. Cephalalgia, 2022, 42, 1039-1049.	1.8	4
8	In vivo brain MR spectroscopy in gliomas: clinical and pre-clinical chances. Clinical and Translational Imaging, 2022, 10, 495-515.	1.1	6
9	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
10	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.5	6
11	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
12	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
13	<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
14	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
15	<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
16	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
17	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
18	Biallelic mutations in RNF220 cause laminopathies featuring leukodystrophy, ataxia and deafness. Brain, 2021, 144, 3020-3035.	3.7	11

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19	Neuroimaging of Pediatric Cerebellum in Inherited Neurodegenerative Diseases. Applied Sciences (Switzerland), 2021, 11, 8522.	1.3	1
20	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
21	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
22	Quantitative Muscle MRI Protocol as Possible Biomarker in Becker Muscular Dystrophy. Clinical Neuroradiology, 2021, 31, 257-266.	1.0	12
23	Sleeve-Shaped Neurothekeoma of the Ulnar Nerve: A Unique Case of a Still Unclear Pathological Entity. Hand, 2020, 15, NP7-NP10.	0.7	1
24	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
25	Phenotypic spectrum of short-chain enoyl-Coa hydratase-1 (ECHS1) deficiency. European Journal of Paediatric Neurology, 2020, 28, 151-158.	0.7	18
26	Abnormal cerebellar foliation in EBF3 mutation. Neurology, 2020, 94, 933-935.	1.5	6
27	MYORG -related disease is associated with central pontine calcifications and atypical parkinsonism. Neurology: Genetics, 2020, 6, e399.	0.9	13
28	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
29	Harmful Iron-Calcium Relationship in Pantothenate kinase Associated Neurodegeneration. International Journal of Molecular Sciences, 2020, 21, 3664.	1.8	19
30	Discovering the Italian phenotype of cerebral amyloid angiopathy (CAA): the SENECA project. Neurological Sciences, 2020, 41, 2193-2200.	0.9	3
31	Understanding Cluster Headache Using Magnetic Resonance Imaging. Frontiers in Neurology, 2020, 11, 535.	1.1	8
32	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
33	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
34	Chiari 1 malformation and untreated sagittal synostosis: a new subset of complex Chiari?. Child's Nervous System, 2019, 35, 1741-1753.	0.6	11
35	Novel mutations in SLC16A2 associated with a less severe phenotype of MCT8 deficiency. Metabolic Brain Disease, 2019, 34, 1565-1575.	1.4	12
36	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

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37	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
38	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
39	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
40	Cluster headache: insights from resting-state functional magnetic resonance imaging. Neurological Sciences, 2019, 40, 45-47.	0.9	10
41	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
42	Clinical spectrum of PTEN mutation in pediatric patients. A bicenter experience. European Journal of Medical Genetics, 2019, 62, 103596.	0.7	39
43	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
44	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
45	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
46	SLC19A3 related disorder: Treatment implication and clinical outcome of 2 new patients. European Journal of Paediatric Neurology, 2018, 22, 332-335.	0.7	6
47	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
48	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
49	Adult leukoencephalopathies with prominent infratentorial involvement can be caused by Erdheimâ€“Chester disease. Journal of Neurology, 2018, 265, 273-284.	1.8	17
50	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
51	Encephalopathies with intracranial calcification in children: clinical and genetic characterization. Orphanet Journal of Rare Diseases, 2018, 13, 135.	1.2	17
52	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
53	Conventional MRI. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 154, 219-234.	1.0	4
54	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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55	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
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	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
59	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
60	A PDE10A de novo mutation causes childhoodâ€onset chorea with diurnal fluctuations. Movement Disorders, 2017, 32, 1646-1647.	2.2	13
61	Thiamine-responsive disease due to mutation of tpk1 : Importance of avoiding misdiagnosis. Neurology, 2017, 89, 870-871.	1.5	13
62	Epileptic spikes in Rasmussenâ€™s encephalitis: Migratory pattern and short-term evolution. A MEG study. Clinical Neurophysiology, 2017, 128, 1898-1905.	0.7	2
63	Cranial ultrasound is a reliable first step imaging in children with suspected craniosynostosis. Child's Nervous System, 2017, 33, 1545-1552.	0.6	27
64	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
65	Neurological Disorders Associated with Striatal Lesions: Classification and Diagnostic Approach. Current Neurology and Neuroscience Reports, 2016, 16, 54.	2.0	23
66	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
67	A 52â€Yearâ€Old Man with Myoclonic Jerks. Brain Pathology, 2016, 26, 291-292.	2.1	2
68	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
69	Intrathecal synthesis of onconeural antibodies in patients with paraneoplastic syndromes. Journal of Neuroimmunology, 2016, 290, 119-122.	1.1	10
70	Persistent orthostatic headache without intracranial hypotension: which treatment?. Neurological Sciences, 2015, 36, 153-155.	0.9	2
71	Resting state fMRI in cluster headache: which role?. Neurological Sciences, 2015, 36, 47-50.	0.9	8
72	Low-Functioning Autism and Nonsyndromic Intellectual Disability. Journal of Child Neurology, 2015, 30, 1658-1663.	0.7	32

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73	Electroclinical spectrum of the neuronal ceroid lipofuscinoses associated with CLN6 mutations. Neurology, 2015, 85, 316-324.	1.5	40
74	A novel homozygous ISPD gene mutation causing phenotype variability in a consanguineous family. Neuromuscular Disorders, 2015, 25, 55-59.	0.3	6
75	Overlapping phenotypes in complex spastic paraplegias SPG11, SPG15, SPG35 and SPG48. Brain, 2014, 137, 1907-1920.	3.7	133
76	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
77	Exploring cerebral networks in cluster headache: Insights and perspectives. Cephalalgia, 2014, 34, 323-324.	1.8	0
78	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
79	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
80	Neuroimaging Findings in 41 Low-Functioning Children With Autism Spectrum Disorder. Journal of Child Neurology, 2014, 29, 1626-1631.	0.7	19
81	Intracranial idiopathic hypertension: 1-year follow-up study. Neurological Sciences, 2014, 35, 177-179.	0.9	8
82	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
83	Neurodegeneration with Brain Iron Accumulation. , 2014, , 171-198.		0
84	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
85	Treatment of spontaneous intracranial hypotension: evolution of the therapeutic and diagnostic modalities. Neurological Sciences, 2013, 34, 151-155.	0.9	29
86	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
87	Pearls & Oy-sters: The use of CT venography in Hirayama disease. Neurology, 2013, 80, 1539-1539.	1.5	0
88	Autosomal Dominant Spinocerebellar Ataxias and Episodic Ataxias. , 2013, , 2193-2267.		5
89	In Medicationâ€Overuse Headache, <scp>fMRI</scp> Shows Longâ€Lasting Dysfunction in Midbrain Areas. Headache, 2012, 52, 1520-1534.	1.8	58
90	Brownâ€“Vialettoâ€“van Laere and Fazioâ€“Londe overlap syndromes: A clinical, biochemical and genetic study. Neuromuscular Disorders, 2012, 22, 1075-1082.	0.3	36

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91	Pain Processing in Medication Overuse Headache: A Functional Magnetic Resonance Imaging (fMRI) Study. Pain Medicine, 2012, 13, 255-262.	0.9	74
92	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
93	Therapeutic Advances in Neurodegeneration With Brain Iron Accumulation. Seminars in Pediatric Neurology, 2012, 19, 82-86.	1.0	9
94	Paediatric Stroke: Review of the Literature and Possible Treatment Options, including Endovascular Approach. Stroke Research and Treatment, 2011, 2011, 1-11.	0.5	7
95	Hemophagocytic lymphohistiocytosis with neurological presentation: MRI findings and a nearly miss diagnosis. Neurological Sciences, 2011, 32, 473-477.	0.9	35
96	Treatment of alterations in CSF dynamics. Neurological Sciences, 2011, 32, 117-120.	0.9	19
97	Neuroradiological diagnosis of Chiari malformations. Neurological Sciences, 2011, 32, 283-286.	0.9	34
98	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
99	Spontaneous intracranial hypotension: diagnostic and therapeutic implications in neurosurgical practice. Neurological Sciences, 2011, 32, 287-290.	0.9	16
100	Treatment for Chiari 1 malformation (CIM): analysis of a pediatric surgical series. Neurological Sciences, 2011, 32, 321-324.	0.9	25
101	Neurological pictures in Paediatric Chiari I malformation. Neurological Sciences, 2011, 32, 295-298.	0.9	17
102	Pontine tegmental cap dysplasia: developmental and cognitive outcome in three adolescent patients. Orphanet Journal of Rare Diseases, 2011, 6, 36.	1.2	44
103	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
104	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
105	Neuroimaging in chronic migraine. Neurological Sciences, 2010, 31, 19-22.	0.9	38
106	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
107	Chronic Migraine With Medication Overuse Preâ€“Post Withdrawal of Symptomatic Medication: Clinical Results and fMRI Correlations. Headache, 2010, 50, 998-1004.	1.8	68
108	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

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109	Sagging and swelling of the midbrain suggest spontaneous intracranial hypotension rather than a malformation. Brain, 2010, 133, e148-e148.	3.7	4
110	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
111	Age-related Iron Deposition in the Basal Ganglia: Quantitative Analysis in Healthy Subjects. Radiology, 2009, 252, 165-172.	3.6	266
112	Clinical features and outcomes in spontaneous intracranial hypotension: a survey of 90 consecutive patients. Neurological Sciences, 2009, 30, 11-13.	0.9	37
113	Functional-MRI evaluation of pain processing in chronic migraine with medication overuse. Neurological Sciences, 2009, 30, 71-74.	0.9	22
114	Application of IHS Criteria to Headache Attributed to Spontaneous Intracranial Hypotension in a Large Population. Cephalalgia, 2009, 29, 418-422.	1.8	43
115	Sindrome da ipotensione liquorale spontanea. , 2009, , 343-345.		0
116	Headache attributed to spontaneous intracranial hypotension. Neurological Sciences, 2008, 29, 164-165.	0.9	17
117	Aicardiâ€GoutiÃ¨res syndrome: description of a late onset case. Developmental Medicine and Child Neurology, 2008, 50, 631-634.	1.1	35
118	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
119	Can MR Imaging Diagnose Adult-Onset Alexander Disease?. American Journal of Neuroradiology, 2008, 29, 1190-1196.	1.2	99
120	Adult-onset leukoencephalopathy with calcifications associated with primary antiphospholipid syndrome. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 1215-1217.	0.9	0
121	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
122	A Case of Pediatric Tumefactive Demyelinating Lesion Misdiagnosed and Treated as Glioblastoma. Journal of Child Neurology, 2008, 23, 944-947.	0.7	22
123	POEMS syndrome: relapse after successful autologous peripheral blood stem cell transplantation. Neuromuscular Disorders, 2007, 17, 980-982.	0.3	25
124	Spontaneous intracranial hypotension with deep brain swelling. Brain, 2007, 130, 1884-1893.	3.7	98
125	Headache and spontaneous low cerebrospinal fluid pressure syndrome. Neurological Sciences, 2007, 28, S232-S234.	0.9	10
126	Periventricular Nodular Heterotopia: Classification, Epileptic History, and Genesis of Epileptic Discharges. Epilepsia, 2006, 47, 86-97.	2.6	150

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127	Prolonged Focal Negative Motor Seizures: A Video-EEG Study. Epilepsia, 2006, 47, 1949-1952.	2.6	14
128	Adult-Onset Rasmussen's Encephalitis: Anatomical-Electrographic-Clinical Features of 7 Italian Cases. Epilepsia, 2006, 47, 41-46.	2.6	108
129	Dural sinus thrombosis in spontaneous intracranial hypotension. Journal of Neurology, 2006, 253, 1197-1202.	1.8	51
130	Iatrogenic intracranial pseudoaneurysms: neuroradiological and therapeutical considerations, including endovascular options. Neurological Sciences, 2006, 27, 317-322.	0.9	56
131	Spontaneous CSF leak treated with percutaneous CT-guided fibrin glue. Neurology, 2006, 66, 782-782.	1.5	8
132	Stimulation of the globus pallidus internus for childhood-onset dystonia. Movement Disorders, 2005, 20, 1194-1200.	2.2	162
133	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
134	Persistent Fixed Torticollis due to Atlanto-Axial Rotatory Fixation: Report of 4 Pediatric Cases. Neuropediatrics, 2005, 36, 45-49.	0.3	8
135	Hyperhomocysteinemia and other thrombophilic risk factors in 26 patients with cerebral venous thrombosis. European Journal of Neurology, 2004, 11, 405-409.	1.7	44
136	Spontaneous low cerebrospinal pressure: a mini review. Neurological Sciences, 2004, 25, s135-s137.	0.9	42
137	Headache and intracranial hypotension: neuroradiological findings. Neurological Sciences, 2004, 25, s138-s141.	0.9	48
138	Diagnostic imaging in 13Âcases of Rasmussen's encephalitis: can early MRI suggest the diagnosis?. Neuroradiology, 2003, 45, 171-183.	1.1	116
139	Rhythmic cortical myoclonus in a case of HIV-related encephalopathy. Movement Disorders, 2003, 18, 1533-1538.	2.2	14
140	Epileptic and imaging findings in perinatal hypoxic-ischemic encephalopathy with ulegyria. Epilepsy Research, 2003, 55, 235-243.	0.8	45
141	Rasmussenâ€™s encephalitis. Neurology, 2003, 60, 422-425.	1.5	137
142	Pseudotumor Cerebri dell'etÃ infantile e adolescenziale. The Neuroradiology Journal, 2003, 16, 497-501.	0.1	0
143	Spinal radiological findings in nine patients with spontaneous intracranial hypotension. Neuroradiology, 2002, 44, 143-150.	1.1	139
144	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

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145	Imaging and outcome in severe complications of lumbar epidural anaesthesia: report of 16 cases. Neuroradiology, 2000, 42, 564-571.	1.1	37
146	Spinal cord lesion due to epidural anesthesia. Neurological Sciences, 2000, 21, 411-412.	0.9	1
147	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
148	Artificial lamina-assisted laminoplasty performed in seven cases. Journal of Neurosurgery: Spine, 1999, 91, 43-49.	0.9	7
149	Type II neurofibromatosis presenting as quadriceps atrophy. Italian Journal of Neurological Sciences, 1998, 19, 94-96.	0.1	10