Andrea Rossi

List of Publications by Year in descending order

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150 papers	4,466 citations	94381 37 h-index	138417 58 g-index
156 all docs	156 docs citations	156 times ranked	5962 citing authors

#	Article	IF	Citations
1	Diabetes Insipidus â€“ Diagnosis and Management. Hormone Research in Paediatrics, 2012, 77, 69-84.	0.8	222
2	Imaging of Neurologic Disease in Hospitalized Patients with COVID-19: An Italian Multicenter Retrospective Observational Study. Radiology, 2020, 297, E270-E273.	3.6	149
3	Definitions and classification of malformations of cortical development: practical guidelines. Brain, 2020, 143, 2874-2894.	3.7	145
4	Imaging of Acute Disseminated Encephalomyelitis. Neuroimaging Clinics of North America, 2008, 18, 149-161.	0.5	129
5	NANS-mediated synthesis of sialic acid is required for brain and skeletal development. Nature Genetics, 2016, 48, 777-784.	9.4	125
6	Imaging in spine and spinal cord malformations. European Journal of Radiology, 2004, 50, 177-200.	1.2	123
7	Medulloblastoma Variants: Age-Dependent Occurrence and Relation to Gorlin Syndromeâ€”A New Clinical Perspective. Clinical Cancer Research, 2009, 15, 2463-2471.	3.2	112
8	Central Diabetes Insipidus in Children and Young Adults: Etiological Diagnosis and Long-Term Outcome of Idiopathic Cases. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1264-1272.	1.8	97
9	POMT2 gene mutation in limb-girdle muscular dystrophy with inflammatory changes. Biochemical and Biophysical Research Communications, 2007, 363, 1033-1037.	1.0	91
10	Magnetic Resonance Imaging of Spinal Dysraphism. Topics in Magnetic Resonance Imaging, 2001, 12, 375-409.	0.7	89
11	Congenital tumors of the central nervous system. Neuroradiology, 2010, 52, 531-548.	1.1	87
12	Deficiency of hyccin, a newly identified membrane protein, causes hypomyelination and congenital cataract. Nature Genetics, 2006, 38, 1111-1113.	9.4	82
13	Novel DyneinDYNC1H1Neck and Motor Domain Mutations Link Distal Spinal Muscular Atrophy and Abnormal Cortical Development. Human Mutation, 2014, 35, 298-302.	1.1	77
14	Diffusion tensor imaging and fiber tractography in brain malformations. Pediatric Radiology, 2013, 43, 28-54.	1.1	71
15	The genotypic and phenotypic spectrum of PIGA deficiency. Orphanet Journal of Rare Diseases, 2015, 10, 23.	1.2	70
16	Magnetic resonance imaging spectrum of medulloblastoma. Neuroradiology, 2011, 53, 387-396.	1.1	69
17	Leigh Syndrome with COX deficiency and SURF1 gene mutations: MR imaging findings. American Journal of Neuroradiology, 2003, 24, 1188-91.	1.2	69
18	Low-grade intraventricular hemorrhage: is ultrasound good enough?. Journal of Maternal-Fetal and Neonatal Medicine, 2015, 28, 2261-2264.	0.7	68

#	Article	IF	Citations
19	Pediatric Neuroradiology. , 2005, , .		66
20	Spectrum of Nonterminal Myelocystoceles. Neurosurgery, 2006, 58, 509-515.	0.6	66
21	The Diagnosis of Children with Central Diabetes Insipidus. Journal of Pediatric Endocrinology and Metabolism, 2007, 20, 359-75.	0.4	62
22	The use of neuroimaging for assessing disorders of pituitary development. Clinical Endocrinology, 2012, 76, 161-176.	1.2	62
23	New MR sequences (diffusion, perfusion, spectroscopy) in brain tumours. Pediatric Radiology, 2010, 40, 999-1009.	1.1	53
24	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.	3.3	53
25	Tumors of the Spine in Children. Neuroimaging Clinics of North America, 2007, 17, 17-35.	0.5	51
26	No major role for the EMX2 gene in schizencephaly. American Journal of Medical Genetics, Part A, 2008, 146A, 1142-1150.	0.7	51
27	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.	1.4	51
28	Diagnostic and prognostic value of¹⁸F-DOPA PET and¹H-MR spectroscopy in pediatric supratentorial infiltrative gliomas: a comparative study. Neuro-Oncology, 2015, 17, 1637-1647.	0.6	49
29	White matter lesions in spastic paraplegia with mutations in SPG5/CYP7B1. Neuromuscular Disorders, 2009, 19, 62-65.	0.3	48
30	Pontine tegmental cap dysplasia: developmental and cognitive outcome in three adolescent patients. Orphanet Journal of Rare Diseases, 2011, 6, 36.	1.2	44
31	Deterioration of Growth Hormone (GH) Response and Anterior Pituitary Function in Young Adults with Childhood-Onset GH Deficiency and Ectopic Posterior Pituitary: A Two-Year Prospective Follow-Up Study. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 3875-3884.	1.8	43
32	Value of ¹⁸F-3,4-Dihydroxyphenylalanine PET/MR Image Fusion in Pediatric Supratentorial Infiltrative Astrocytomas: A Prospective Pilot Study. Journal of Nuclear Medicine, 2014, 55, 718-723.	2.8	43
33	Pediatric astrocytic tumor grading: comparison between arterial spin labeling and dynamic susceptibility contrast MRI perfusion. Neuroradiology, 2018, 60, 437-446.	1.1	43
34	MR imaging of brain-stem hypoplasia in horizontal gaze palsy with progressive scoliosis. American Journal of Neuroradiology, 2004, 25, 1046-8.	1.2	43
35	Cerebrovascular disease and varicella in children. Brain and Development, 2006, 28, 366-370.	0.6	42
36	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.	2.3	41

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37	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.	3.3	41
38	Treatment and outcome of children with cerebral cavernomas: a survey on 32 patients. Neurological Sciences, 2010, 31, 117-123.	0.9	40
39	Internal jugular vein phlebectasia and duplication: case report with magnetic resonance angiography features. Pediatric Radiology, 2001, 31, 134-134.	1.1	39
40	Phenotypic characterization of hypomyelination and congenital cataract. Annals of Neurology, 2007, 62, 121-127.	2.8	39
41	Management of diabetes insipidus and adipsia in the child. Best Practice and Research in Clinical Endocrinology and Metabolism, 2015, 29, 415-436.	2.2	39
42	POMGnT1 Mutations in Congenital Muscular Dystrophy. Archives of Neurology, 2006, 63, 1491.	4.9	38
43	Current Classification and Imaging of Congenital Spinal Abnormalities. Seminars in Roentgenology, 2006, 41, 250-273.	0.2	37
44	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. European Radiology, 2017, 27, 5080-5092.	2.3	36
45	Multimodal Magnetic Resonance Imaging and ¹⁸F-L-Dihydroxyphenylalanine Positron Emission Tomography in Early Characterization of Pseudoresponse and Nonenhancing Tumor Progression in a Pediatric Patient With Malignant Transformation of Ganglioglioma Treated With Bevacizumab. Journal of Clinical Oncology, 2013, 31, e1-e5.	0.8	35
46	Natural history of cavernous malformations in children with brain tumors treated with radiotherapy and chemotherapy. Journal of Neuro-Oncology, 2014, 117, 311-320.	1.4	35
47	Different gestational ages and changing vulnerability of the premature brain. Journal of Maternal-Fetal and Neonatal Medicine, 2015, 28, 2268-2272.	0.7	34
48	Molecular Genetic Analysis of the PLP1 Gene in 38 Families with PLP1-related disorders: Identification and Functional Characterization of 11 Novel PLP1 Mutations. Orphanet Journal of Rare Diseases, 2011, 6, 40.	1.2	32
49	A Novel Mutation in the SURF1 Gene in a Child With Leigh Disease, Peripheral Neuropathy, and Cytochrome-c Oxidase Deficiency. Journal of Child Neurology, 2002, 17, 233-236.	0.7	31
50	Genetic disorders affecting white matter in the pediatric age. American Journal of Medical Genetics Part A, 2004, 129B, 85-93.	2.4	30
51	Prematurity and brain perfusion: Arterial spin labeling MRI. NeuroImage: Clinical, 2017, 15, 401-407.	1.4	30
52	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.	1.4	30
53	Early Pain Exposure Influences Functional Brain Connectivity in Very Preterm Neonates. Frontiers in Neuroscience, 2019, 13, 899.	1.4	30
54	EXOSC3 mutations in isolated cerebellar hypoplasia and spinal anterior horn involvement. Journal of Neurology, 2013, 260, 1866-1870.	1.8	28

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55	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.	0.9	28
56	Spontaneous rupture of middle fossa arachnoid cysts: surgical series from a single center pediatric hospital and literature review. Child's Nervous System, 2020, 36, 2789-2799.	0.6	28
57	Magnetic resonance imaging in childhood leukemia survivors treated with cranial radiotherapy: A cross sectional, single center study. Pediatric Blood and Cancer, 2011, 57, 240-246.	0.8	25
58	Skull base osteomyelitis and potential cerebrovascular complications in children. Pediatric Radiology, 2012, 42, 867-874.	1.1	25
59	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.	3.3	25
60	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.	1.1	25
61	Posterior pituitary (PP) evaluation in patients with anterior pituitary defect associated with ectopic PP and septo-optic dysplasia. European Journal of Endocrinology, 2011, 165, 411-420.	1.9	24
62	Immunomodulatory therapy in recurrent acute necrotizing encephalopathy ANE1: Is it useful?. Brain and Development, 2012, 34, 384-391.	0.6	24
63	Magnetic Resonance Spectroscopy in Metabolic Disorders. Neuroimaging Clinics of North America, 2013, 23, 425-448.	0.5	24
64	Neonatal Hypoglycemia and Brain Vulnerability. Frontiers in Endocrinology, 2021, 12, 634305.	1.5	24
65	Cervico-medullary desmoplastic infantile ganglioglioma: An unusual case with diffuse leptomeningeal dissemination at diagnosis. Pediatric Blood and Cancer, 2005, 45, 986-990.	0.8	23
66	Developmental Abnormalities of the Posterior Pituitary Gland. Endocrine Development, 2009, 14, 83-94.	1.3	23
67	Expanding the spectrum of congenital anomalies of the diencephalicâ€“mesencephalic junction. Neuroradiology, 2016, 58, 33-44.	1.1	23
68	Brain-injured Survivors of Monochorionic Twin Pregnancies Complicated by Single Intrauterine Death: MR Findings in a Multicenter Study. Radiology, 2018, 288, 582-590.	3.6	23
69	Agenesis of bilateral internal carotid arteries in the PHACE syndrome. American Journal of Neuroradiology, 2006, 27, 1602.	1.2	23
70	Hypomyelination and Congenital Cataract. Archives of Neurology, 2011, 68, 1191.	4.9	22
71	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.	1.9	22
72	Pediatric Spinal Infection and Inflammation. Neuroimaging Clinics of North America, 2015, 25, 173-191.	0.5	21

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73	Anti-N-methyl-D-Aspartate-Receptor Encephalitis in a Four-Year-Old Girl. Journal of Pediatrics, 2010, 156, 332-334.	0.9	20
74	Neuroimaging of Infectious and Inflammatory Diseases of the Pediatric Cerebellum and Brainstem. Neuroimaging Clinics of North America, 2016, 26, 471-487.	0.5	20
75	Congenital Malformations of the Spine and Spinal Cord. , 2005, , 1551-1608.		19
76	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.	1.1	19
77	MR Imaging of Neonatal Spinal Dysraphia: What to Consider?. Magnetic Resonance Imaging Clinics of North America, 2012, 20, 45-61.	0.6	18
78	Intermittentâ€relapsing pyruvate dehydrogenase complex deficiency: a case with clinical, biochemical, and neuroradiological reversibility. Developmental Medicine and Child Neurology, 2012, 54, 472-476.	1.1	18
79	Moyamoya Vasculopathy in PHACE Syndrome: Six New Cases and Review of the Literature. World Neurosurgery, 2017, 108, 291-302.	0.7	17
80	Bilateral germinoma of the basal ganglia. Pediatric Blood and Cancer, 2008, 50, 177-179.	0.8	16
81	Punctate white matter lesions of preterm infants: Risk factor analysis. European Journal of Paediatric Neurology, 2019, 23, 733-739.	0.7	16
82	Added value of diffusion weighted imaging in pediatric central nervous system embryonal tumors surveillance. Oncotarget, 2017, 8, 60401-60413.	0.8	16
83	Incidental findings on routine brain MRI scans in preterm infants. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2017, 102, F73-F78.	1.4	15
84	ABCC6 mutations and early onset stroke: Two cases of a typical Pseudoxanthoma Elasticum. European Journal of Paediatric Neurology, 2018, 22, 725-728.	0.7	15
85	Quantitative susceptibility map analysis in preterm neonates with germinal matrixâ€intraventricular hemorrhage. Journal of Magnetic Resonance Imaging, 2018, 48, 1199-1207.	1.9	15
86	Congenital spine anomalies: the closed spinal dysraphisms. Pediatric Radiology, 2015, 45, 413-419.	1.1	14
87	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.	4.6	14
88	Cystic angiomatosis of the craniocervical junction associated with Chiari I malformation. Child's Nervous System, 2007, 23, 697-700.	0.6	13
89	Diagnostic Approach to Pediatric Spine Disorders. Magnetic Resonance Imaging Clinics of North America, 2016, 24, 621-644.	0.6	13
90	Widening the Heterogeneity of Leigh Syndrome: Clinical, Biochemical, and Neuroradiologic Features in a Patient Harboring a NDUFA10 Mutation. JIMD Reports, 2017, 37, 37-43.	0.7	13

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91	Role of diffusion weighted imaging for differentiating cerebral pilocytic astrocytoma and ganglioglioma BRAF V600E-mutant from wild type. Neuroradiology, 2020, 62, 71-80.	1.1	13
92	Tonsillar herniation spectrum: more than just Chiari I. Update and controversies on classification and management. Neurosurgical Review, 2020, 43, 1473-1492.	1.2	13
93	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.	0.7	12
94	Novel asymptomatic CNS findings in patients withACVR1/ALK2mutations causing fibrodysplasia ossificans progressiva. Journal of Medical Genetics, 2016, 53, 859-864.	1.5	12
95	Prenatal MR imaging features of isolated cerebellar haemorrhagic lesions. European Radiology, 2016, 26, 2685-2696.	2.3	12
96	Agenesis of the putamen and globus pallidus caused by recessive mutations in the homeobox gene GSX2. Brain, 2019, 142, 2965-2978.	3.7	12
97	Dissecting the neurological phenotype in children with callosal agenesis, interhemispheric cysts and malformations of cortical development. Journal of Neurology, 2019, 266, 1167-1181.	1.8	12
98	Arterial spin labeling perfusion in neonates. Seminars in Fetal and Neonatal Medicine, 2020, 25, 101130.	1.1	12
99	Guidelines for magnetic resonance imaging in pediatric head and neck pathologies: a multicentre international consensus paper. Neuroradiology, 2022, 64, 1081-1100.	1.1	12
100	18F-DOPA Uptake of Developmental Venous Anomalies in Children With Brain Tumors. Clinical Nuclear Medicine, 2016, 41, e351-e352.	0.7	11
101	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.	1.1	11
102	Pediatric Diffuse Midline Gliomas H3 K27M-Mutant and Non-Histone Mutant Midline High-Grade Gliomas in Neurofibromatosis Type 1 in Comparison With Non-Syndromic Children: A Single-Center Pilot Study. Frontiers in Oncology, 2020, 10, 795.	1.3	11
103	White matter and cerebellar involvement in alternating hemiplegia of childhood. Journal of Neurology, 2020, 267, 1300-1311.	1.8	10
104	Cognitive and White Matter Microstructure Development in Congenital Hypothyroidism and Familial Thyroid Disorders. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3990-e4006.	1.8	10
105	Post-chemotherapy maturation of a pineoblastoma. Acta Neuropathologica, 2010, 119, 651-653.	3.9	9
106	Targeted re-sequencing in pediatric and perinatal stroke. European Journal of Medical Genetics, 2020, 63, 104030.	0.7	9
107	Sellar and Suprasellar Disorders. , 2005, , 855-891.		8
108	Hypothalamic-pituitary magnetic resonance imaging in growth hormone deficiency. Expert Review of Endocrinology and Metabolism, 2006, 1, 413-423.	1.2	8

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109	Unusual white matter involvement in EAST syndrome associated with novel KCNJ10 mutations. Journal of Neurology, 2018, 265, 1419-1425.	1.8	8
110	A consensus response on the complete picture: reply to LynÃ¸e and Eriksson. Pediatric Radiology, 2019, 49, 424-428.	1.1	8
111	L1CAM variants cause two distinct imaging phenotypes on fetal MRI. Annals of Clinical and Translational Neurology, 2021, 8, 2004-2012.	1.7	8
112	Hypomyelination and congenital cataract: Identification of novel mutations in two unrelated families. European Journal of Paediatric Neurology, 2013, 17, 108-111.	0.7	7
113	Further genotypeâ€“phenotype correlation emerging from two families with PLP1 exon 4 skipping. Clinical Genetics, 2014, 85, 267-272.	1.0	7
114	Urological outcome in patients with Currarino syndrome. Journal of Pediatric Surgery, 2014, 49, 1643-1646.	0.8	7
115	Torcular pseudomass: a potential diagnostic pitfall in infants and young children. Pediatric Radiology, 2017, 47, 227-234.	1.1	7
116	Nosological Differences in the Nature of Punctate White Matter Lesions in Preterm Infants. Frontiers in Neurology, 2021, 12, 657461.	1.1	7
117	Management: opinions from different centersâ€”the Istituto Giannina Gaslini experience. Child's Nervous System, 2019, 35, 1905-1909.	0.6	6
118	Late Persistent Increased Putaminal 18F-DOPA Uptake Following Ipsilateral Frontal Resection. Clinical Nuclear Medicine, 2015, 40, e451-e452.	0.7	5
119	Teaching Neuro Images : Figure of 8. Neurology, 2017, 89, e172-e173.	1.5	5
120	Placental Pathology Findings and the Risk of Intraventricular and Cerebellar Hemorrhage in Preterm Neonates. Frontiers in Neurology, 2020, 11, 761.	1.1	5
121	Focal status and acute encephalopathy in a 13-year-old boy with de novo DNM1L mutation: Video-polygraphic pattern and clues for differential diagnosis. Brain and Development, 2021, 43, 644-651.	0.6	5
122	External ventricular drainage for posthemorrhagic ventricular dilatation in preterm infants: insights on efficacy and failure. Journal of Neurosurgery: Pediatrics, 2021, 28, 563-571.	0.8	5
123	Role of Dynamic Parameters of 18F-DOPA PET/CT in Pediatric Gliomas. Clinical Nuclear Medicine, 2022, 47, 517-524.	0.7	5
124	Cognitive Profiles and Brain Volume Are Affected in Patients with Silverâ€“Russell Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e1478-e1488.	1.8	4
125	Coexisting Retrocerebellar Arachnoid Cyst and Chiari Type 1 Malformation: 3 Pediatric Cases of Surgical Management Tailored to the Pathogenic Mechanism and Systematic Review of the Literature. World Neurosurgery, 2021, 148, 44-53.	0.7	4
126	Recurrent obstructive hydrocephalus in a 4-month-old infant. Child's Nervous System, 2010, 26, 133-136.	0.6	3

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127	Cerebellar hypoplasia and brainstem thinning associated with severe white matter and basal ganglia abnormalities in a child with an mtDNA deletion. Journal of Inherited Metabolic Disease, 2011, 34, 1225-1227.	1.7	3
128	Endocrine Outcomes In Central Diabetes Insipidus: the Predictive Value of Neuroimaging â€œMismatch Patternâ€. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 3562-3574.	1.8	3
129	Imaging of Pediatric Brain and Spinal Cord Tumors. , 2012, , 203-214.		2
130	Congenital Malformations of the Spine, Spinal Cord, and Cranio-Cervical Junction. Medical Radiology, 2007, , 3-40.	0.0	2
131	SPG5-related spastic paraplegia and white matter abnormalities. Neuromuscular Disorders, 2009, 19, 507-508.	0.3	1
132	Pediatric Sellar and Suprasellar Disorders. , 2015, , 1-66.		1
133	Pediatric Pineal Region Tumors. , 2015, , 1-31.		1
134	Malformations of Cortical Development. , 2021, , 1-237.		1
135	Imaging characteristics and neurosurgical outcome in subjects with agenesis of the corpus callosum and interhemispheric cysts. Neuroradiology, 2022, 64, 2163-2177.	1.1	1
136	Pediatric Spine and Spinal Cord. The Neuroradiology Journal, 2003, 16, 188-191.	0.1	0
137	Diagnosi radiologica dei difetti ipofisari in etÃ pediatrica. L Endocrinologo, 2008, 9, 14-20.	0.0	0
138	Ectopic Posterior Pituitary Lobe. , 0, , 87-88.		0
139	Acute Disseminated Encephalomyelitis (ADEM). , 0, , 231-232.		0
140	Choroid Plexus Papilloma. , 0, , 305-306.		0
141	Hybrid Imaging in Pediatric Central Nervous System Disorders. , 2016, , 195-217.		0
142	The Acute Pediatric Spine and Spinal Cord. , 2016, , 317-336.		0
143	Imaging in Spine and Spinal Cord Developmental Malformations. , 2019, , 1609-1640.		0
144	Imaging in Spine and Spinal Cord Developmental Malformations. , 2019, , 1-32.		0

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145	Acquired Infectious and Autoimmune Diseases of the Pediatric Central Nervous System. , 2012, , 280-295.		0
146	Infectious and Inflammatory Diseases of the Spine in Children. , 2015, , 1-55.		0
147	Hydrosyringomyelia, Cysts, and Other Disorders of the Cerebrospinal Fluid Spaces in the Spine. , 2015, , 1-18.		0
148	Pituitary Gland Imaging. , 2016, , 123-146.		0
149	Inflammatory, Demyelinating, and Autoimmune Diseases in Infants and Children. , 2019, , 1-48.		0
150	Stepwise approach for vertebral hemangioma in children: case-reports and treatment algorithm proposal. European Spine Journal, 0, , .	1.0	0