Junnichi Takanashi

List of Publications by Year
in descending order

145 papers	4,518 citations	109321 35 h-index	128289 60 g-index
146 all docs	146 docs citations	146 times ranked	4965 citing authors

#	Article	IF	Citations
1	Increased cytokines/chemokines and hyponatremia as a possible cause of clinically mild encephalitis/encephalopathy with a reversible splenial lesion associated with acute focal bacterial nephritis. Brain and Development, 2022, 44, 30-35.	1.1	13
2	Unilateral internal carotid artery absence in trisomy 18. Brain and Development, 2022, , .	1.1	0
3	Neurochemistry evaluated by MR spectroscopy in a patient with SPTAN1-related developmental and epileptic encephalopathy. Brain and Development, 2022, 44, 415-420.	1.1	3
4	Characteristic nail lesions in Kawasaki disease: Case series and literature review. Journal of Dermatology, 2022, 49, 232-238.	1.2	4
5	Author reply to â€œOnycholysis associated with Kawasaki disease: A comment on characteristic nail lesions in Kawasaki disease: Case series and literature reviewâ€. Journal of Dermatology, 2022, 49, .	1.2	0
6	A child with mitochondrial DNA deletion presenting diabetes mellitus as an initial symptom. Radiology Case Reports, 2022, 17, 2915-2918.	0.6	0
7	Prenatal clinical manifestations in individuals with COL4A1/2 variants. Journal of Medical Genetics, 2021, 58, 505-513.	3.2	22
8	Guidelines for the diagnosis and treatment of acute encephalopathy in childhood. Brain and Development, 2021, 43, 2-31.	1.1	67
9	Evaluation of the Diagnostic Criteria for Anti-NMDA Receptor Encephalitis in Japanese Children. Neurology, 2021, 96, e2070-e2077.	1.1	7
10	Spontaneous tracheal rupture caused by acute asthma exacerbation. Pediatrics International, 2021, 63, 1541-1543.	0.5	0
11	Cerebral white matter lacerations in children caused by repetitive head trauma. Brain and Development, 2020, 42, 83-87.	1.1	0
12	Altered MR imaging findings in a Japanese female child with PRUNE1-related disorder. Brain and Development, 2020, 42, 302-306.	1.1	9
13	Prognostic value of MR spectroscopy in patients with acute excitotoxic encephalopathy. Journal of the Neurological Sciences, 2020, 408, 116636.	0.6	12
14	POLR1C variants dysregulate splicing and cause hypomyelinating leukodystrophy. Neurology: Genetics, 2020, 6, e524.	1.9	4
15	Neurochemistry of hyponatremic encephalopathy evaluated by MR spectroscopy. Brain and Development, 2020, 42, 767-770.	1.1	5
16	Epidemiological changes of acute encephalopathy in Japan based on national surveillance for 2014â€“2017. Brain and Development, 2020, 42, 508-514.	1.1	41
17	Thermolabile polymorphism of carnitine palmitoyltransferase 2: A genetic risk factor of overall acute encephalopathy. Brain and Development, 2019, 41, 862-869.	1.1	10
18	Comprehensive analysis of coding variants highlights genetic complexity in developmental and epileptic encephalopathy. Nature Communications, 2019, 10, 2506.	12.8	46

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19	Reader response: Evidence in focus: Nusinersen use in spinal muscular atrophy: Report of the Guideline Development, Dissemination, and Implementation Subcommittee of the American Academy of Neurology. Neurology, 2019, 93, 464-464.	1.1	3
20	Vital Signs as Predictor Factors of Intravenous Immunoglobulin Resistance in Patients With Kawasaki Disease. Clinical Pediatrics, 2018, 57, 1148-1153.	0.8	8
21	Infantile traumatic brain injury with a biphasic clinical course and late reduced diffusion. Journal of the Neurological Sciences, 2018, 390, 63-66.	0.6	10
22	Excitotoxicity in encephalopathy associated with STEC O-157 infection. Brain and Development, 2018, 40, 357-360.	1.1	2
23	Loss of myelinated axons and astrocytosis in an autopsy case of acute encephalopathy with biphasic seizures and late reduced diffusion. Brain and Development, 2018, 40, 947-951.	1.1	11
24	Neurochemistry evaluated by MR spectroscopy in a patient with xeroderma pigmentosum group A. Brain and Development, 2018, 40, 931-933.	1.1	1
25	Neuroimaging on Pediatric Encephalopathy in Japan. , 2018, , 53-62.		1
26	An episode of acute encephalopathy with biphasic seizures and late reduced diffusion followed by hemiplegia and intractable epilepsy observed in a patient with a novel frameshift mutation in HNRNPU. Brain and Development, 2018, 40, 813-818.	1.1	13
27	Prevalence and characteristics of human parechovirus and enterovirus infection in febrile infants. Pediatrics International, 2018, 60, 142-147.	0.5	20
28	Clinically mild infantile encephalopathy associated with excitotoxicity. Journal of the Neurological Sciences, 2017, 373, 138-141.	0.6	4
29	Identification of novel <scp>SNORD118</scp> mutations in seven patients with leukoencephalopathy with brain calcifications and cysts. Clinical Genetics, 2017, 92, 180-187.	2.0	28
30	A new infectious encephalopathy syndrome, clinically mild encephalopathy associated with excitotoxicity (MEEX). Journal of the Neurological Sciences, 2017, 380, 27-30.	0.6	5
31	A novel DARS2 mutation in a Japanese patient with leukoencephalopathy with brainstem and spinal cord involvement but no lactate elevation. Human Genome Variation, 2017, 4, 17051.	0.7	6
32	The first report of Japanese patients with asparagine synthetase deficiency. Brain and Development, 2017, 39, 236-242.	1.1	25
33	Comprehensive investigation of CASK mutations and other genetic etiologies in 41 patients with intellectual disability and microcephaly with pontine and cerebellar hypoplasia (MICPCH). PLoS ONE, 2017, 12, e0181791.	2.5	44
34	In vitro characterization of neurite extension using induced pluripotent stem cells derived from lissencephaly patients with TUBA1A missense mutations. Molecular Brain, 2016, 9, 70.	2.6	22
35	Activated microglia in acute encephalopathy with biphasic seizures and late reduced diffusion. Journal of the Neurological Sciences, 2016, 366, 91-93.	0.6	9
36	Association Between Invisible Basal Ganglia and ZNF335 Mutations: A Case Report. Pediatrics, 2016, 138, .	2.1	12

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37	Elevated taurine and glutamate in cerebral juvenile xanthogranuloma on MR spectroscopy. Brain and Development, 2016, 38, 964-967.	1.1	6
38	Drugs indicated for mitochondrial dysfunction as treatments for acute encephalopathy with onset of febrile convulsive status epileptics. Journal of the Neurological Sciences, 2016, 360, 57-60.	0.6	22
39	The magnetic resonance imaging spectrum of Pelizaeusâ€“Merzbacher disease: A multicenter study of 19 patients. Brain and Development, 2016, 38, 571-580.	1.1	18
40	Neurochemistry of Hypomyelination Investigated with MR Spectroscopy. Magnetic Resonance in Medical Sciences, 2015, 14, 85-91.	2.0	8
41	Clinical Pictures in Pelizaeus-Merzbacher Disease: A Report of a Case. Journal of Nippon Medical School, 2015, 82, 74-75.	0.9	3
42	Predictive score for early diagnosis of acute encephalopathy with biphasic seizures and late reduced diffusion (AESD). Journal of the Neurological Sciences, 2015, 358, 62-65.	0.6	37
43	Clinically mild encephalitis with a reversible splenial lesion (MERS) after mumps vaccination. Journal of the Neurological Sciences, 2015, 349, 226-228.	0.6	27
44	Progressive brain atrophy in Schinzelâ€“Giedion syndrome with a SETBP1 mutation. European Journal of Medical Genetics, 2015, 58, 369-371.	1.3	22
45	Efficacy and safety of fosphenytoin for acute encephalopathy in children. Brain and Development, 2015, 37, 418-422.	1.1	9
46	Disrupted glutamate-glutamine cycle in acute encephalopathy with biphasic seizures and late reduced diffusion. Neuroradiology, 2015, 57, 1163-1168.	2.2	39
47	Comment on â€œDelayed myelination is not a constant feature of Allanâ€“Herndonâ€“Dudley syndrome: Report of a new case and review of the literatureâ€•by Azzolini S et al. Brain & Development 2014;36:716â€“720. Brain and Development, 2015, 37, 988-989.	1.1	3
48	Diffusion-weighted MRI for early diagnosis of neonatal herpes simplex encephalitis. Brain and Development, 2015, 37, 423-431.	1.1	19
49	Expanding the phenotypic spectrum of TUBB4A -associated hypomyelinating leukoencephalopathies. Neurology, 2014, 82, 2230-2237.	1.1	45
50	Clinical and radiologic features of encephalopathy during 2011 E coli O111 outbreak in Japan. Neurology, 2014, 82, 564-572.	1.1	36
51	Neurochemistry in shiverer mouse depicted on MR spectroscopy. Journal of Magnetic Resonance Imaging, 2014, 39, 1550-1557.	3.4	10
52	Epidemiological, clinical, and genetic landscapes of hypomyelinating leukodystrophies. Journal of Neurology, 2014, 261, 752-758.	3.6	36
53	C5orf42 is the major gene responsible for OFD syndrome type VI. Human Genetics, 2014, 133, 367-377.	3.8	71
54	Serum and CSF biomarkers in acute pediatric neurological disorders. Brain and Development, 2014, 36, 489-495.	1.1	10

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55	Intragenic mutations in SMN1 may contribute more significantly to clinical severity than SMN2 copy numbers in some spinal muscular atrophy (SMA) patients. Brain and Development, 2014, 36, 914-920.	1.1	39
56	Different patterns of cerebellar abnormality and hypomyelination between POLR3A and POLR3B mutations. Brain and Development, 2014, 36, 259-263.	1.1	21
57	MR spectroscopy in 18q- syndrome suggesting other than hypomyelination. Brain and Development, 2014, 36, 57-60.	1.1	5
58	Phenotypic Spectrum of COL4A1 Mutations: Porencephaly to Schizencephaly. Annals of Neurology, 2013, 73, 48-57.	5.3	143
59	A Child with Three Episodes of Reversible Splenial Lesion. Neuropediatrics, 2013, 44, 199-202.	0.6	2
60	ADORA2A polymorphism predisposes children to encephalopathy with febrile status epilepticus. Neurology, 2013, 80, 1571-1576.	1.1	71
61	Wholeâ€exome sequencing of a unique brain malformation with periventricular heterotopia, cingulate polymicrogyria and midbrain tectal hyperplasia. Neuropathology, 2013, 33, 553-560.	1.2	7
62	MLL2 and KDM6A mutations in patients with Kabuki syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 2234-2243.	1.2	148
63	Clinical and radiological features of Japanese patients with a severe phenotype due to CASK mutations. American Journal of Medical Genetics, Part A, 2012, 158A, 3112-3118.	1.2	34
64	Hypoyelination in I-cell disease; MRI, MR spectroscopy and neuropathological correlation. Brain and Development, 2012, 34, 780-783.	1.1	7
65	Late Delirious Behavior With 2009 H1N1 Influenza: Mild Autoimmune-Mediated Encephalitis?. Pediatrics, 2012, 129, e1068-e1071.	2.1	11
66	Reversible splenial lesion in the corpus callosum following rapid withdrawal of carbamazepine after neurosurgical decompression for trigeminal neuralgia. Journal of Clinical Neuroscience, 2012, 19, 1182-1184.	1.5	18
67	Kawasaki disease complicated by mild encephalopathy with a reversible splenial lesion (MERS). Journal of the Neurological Sciences, 2012, 315, 167-169.	0.6	50
68	A severe form of epidermal nevus syndrome associated with brainstem and cerebellar malformations and neonatal medulloblastoma. Brain and Development, 2012, 34, 881-885.	1.1	8
69	Oxidative stress in patients with clinically mild encephalitis/encephalopathy with a reversible splenial lesion (MERS). Brain and Development, 2012, 34, 124-127.	1.1	60
70	Epidemiology of acute encephalopathy in Japan, with emphasis on the association of viruses and syndromes. Brain and Development, 2012, 34, 337-343.	1.1	328
71	Metabolic changes in early childhood using LCModel with corrected water scaling method. Journal of Magnetic Resonance Imaging, 2012, 35, 174-180.	3.4	11
72	Increased Nâ€acetylaspartate in model mouse of pelizaeusâ€merzbacher disease. Journal of Magnetic Resonance Imaging, 2012, 35, 418-425.	3.4	12

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73	Novel intragenic duplications and mutations of CASK in patients with mental retardation and microcephaly with pontine and cerebellar hypoplasia (MICPCH). Human Genetics, 2012, 131, 99-110.	3.8	40
74	Genetic diagnosis and acetazolamide treatment of familial hemiplegic migraine. Brain and Development, 2011, 33, 332-334.	1.1	15
75	Carnitine palmitoyl transferase II polymorphism is associated with multiple syndromes of acute encephalopathy with various infectious diseases. Brain and Development, 2011, 33, 512-517.	1.1	67
76	Moyamoya disease in children. Brain and Development, 2011, 33, 229-234.	1.1	49
77	Lesions in the central tegmental tract in autopsy cases of developmental brain disorders. Brain and Development, 2011, 33, 541-547.	1.1	17
78	Wide range of CNS manifestations of rotavirus infection. Brain and Development, 2011, 33, 9.	1.1	12
79	Mutations in POLR3A and POLR3B Encoding RNA Polymerase III Subunits Cause an Autosomal-Recessive Hypomyelinating Leukoencephalopathy. American Journal of Human Genetics, 2011, 89, 644-651.	6.2	137
80	Evaluation of ferritin-overexpressing brain in newly developed transgenic mice. Magnetic Resonance Imaging, 2011, 29, 179-184.	1.8	8
81	Pontine hypoplasia in 5p-syndrome: A key MRI finding for a diagnosis. Brain and Development, 2010, 32, 571-573.	1.1	15
82	The axonal damage marker tau protein in the cerebrospinal fluid is increased in patients with acute encephalopathy with biphasic seizures and late reduced diffusion. Brain and Development, 2010, 32, 435-439.	1.1	40
83	Sisters with clinically mild encephalopathy with a reversible splenial lesion (MERS)-like features; Familial MERS?. Journal of the Neurological Sciences, 2010, 290, 153-156.	0.6	35
84	Differences in the time course of splenial and white matter lesions in clinically mild encephalitis/encephalopathy with a reversible splenial lesion (MERS). Journal of the Neurological Sciences, 2010, 292, 24-27.	0.6	91
85	Encephalopathy with a reversible splenial lesion is associated with hyponatremia. Brain and Development, 2009, 31, 217-220.	1.1	121
86	Delirious behavior in influenza is associated with a reversible splenial lesion. Brain and Development, 2009, 31, 423-426.	1.1	32
87	Diffuse cerebral hypomyelination with cerebellar atrophy and hypoplasia of the corpus callosum. Brain and Development, 2009, 31, 582-587.	1.1	27
88	A splenial lesion with transiently reduced diffusion in clinically mild encephalitis is not always reversible: A case report. Brain and Development, 2009, 31, 710-712.	1.1	23
89	Two newly proposed infectious encephalitis/encephalopathy syndromes. Brain and Development, 2009, 31, 521-528.	1.1	244
90	Homozygous female Becker muscular dystrophy. American Journal of Medical Genetics, Part A, 2009, 149A, 1052-1055.	1.2	31

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91	Brainstem disconnection associated with nodular heterotopia and proatlantal arteries. American Journal of Medical Genetics, Part A, 2009, 149A, 2479-2483.	1.2	15
92	Periventricular Haemosiderin Deposition in Patients with Congenital Hemiplegia. Developmental Medicine and Child Neurology, 2008, 37, 1016-1019.	2.1	0
93	Relationship between Dry Weight at Heading and the Number ofSpikelets on Individual Rice Tillers. Plant Production Science, 2007, 10, 430-441.	2.0	16
94	Reversible restricted diffusion of entire corpus callosum. Journal of the Neurological Sciences, 2007, 254, 106.	0.6	0
95	Mild influenza encephalopathy with biphasic seizures and late reduced diffusion. Journal of the Neurological Sciences, 2007, 256, 86-89.	0.6	30
96	The evolving MR imaging appearance of lissencephaly: A case report. Brain and Development, 2007, 29, 522-524.	1.1	3
97	Reversible restricted diffusion of entire corpus callosum. Journal of the Neurological Sciences, 2006, 247, 101-104.	0.6	33
98	14-3-3 Protein Detection in the Cerebrospinal Fluid of Patients With Influenza-Associated Encephalopathy. Journal of Child Neurology, 2006, 21, 562-565.	1.4	8
99	Recurrent Meningitis Associated With a Petrous Apex Cephalocele. Journal of Child Neurology, 2005, 20, 168-170.	1.4	21
100	Finger Cold-Induced Vasodilatation, Sympathetic Skin Response, and Râ€"R Interval Variation in Patients With Progressive Spinal Muscular Atrophy. Journal of Child Neurology, 2005, 20, 871-875.	1.4	26
101	Neonate Showing Reversible Splenial Lesion. Archives of Neurology, 2005, 62, 1481.	4.5	38
102	Grain Growth and Endosperm Cell Size Under High Night Temperatures in Rice (Oryza sativa L.). Annals of Botany, 2005, 95, 695-701.	2.9	218
103	Intracranial dural venous anomalies in familial cervical cystic hygroma. Pediatric Neurology, 2005, 32, 50-52.	2.1	3
104	Magnetic resonance imaging confirms periventricular venous infarction in a term-born child with congenital hemiplegia. Developmental Medicine and Child Neurology, 2005, 47, 706.	2.1	25
105	Pituitary cysts in childhood evaluated by MR imaging. American Journal of Neuroradiology, 2005, 26, 2144-7.	2.4	45
106	Influenza-associated encephalitis/encephalopathy with a reversible lesion in the splenium of the corpus callosum: a case report and literature review. American Journal of Neuroradiology, 2004, 25, 798-802.	2.4	93
107	Reversible white matter lesion in methionine adenosyltransferase I/III deficiency. American Journal of Neuroradiology, 2004, 25, 1843-5.	2.4	37
108	Magnetic resonance imaging in late-onset ornithine transcarbamylase deficiency. Brain and Development, 2003, 25, 40-44.	1.1	25

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109	Gorlin syndrome with ulcerative colitis in a Japanese girl. American Journal of Medical Genetics Part A, 2003, 121A, 65-68.	2.4	11
110	Posterior Pituitary Bright Spot in Large Adenomas: MR Assessment of Its Disappearance or Relocation along the Stalk. Radiology, 2003, 226, 359-365.	7.3	21
111	Decreased Cerebrospinal Fluid Hypocretin-1 Levels Near the Onset of Narcolepsy in 2 Prepubertal Children. Sleep, 2003, 26, 555-557.	1.1	44
112	Brain MR imaging in acute hyperammonemic encephalopathy arising from late-onset ornithine transcarbamylase deficiency. American Journal of Neuroradiology, 2003, 24, 390-3.	2.4	69
113	Middle interhemispheric variant of holoprosencephaly associated with diffuse polymicrogyria. American Journal of Neuroradiology, 2003, 24, 394-7.	2.4	22
114	The changing MR imaging appearance of polymicrogyria: a consequence of myelination. American Journal of Neuroradiology, 2003, 24, 788-93.	2.4	50
115	Brain MR imaging in neonatal hyperammonemic encephalopathy resulting from proximal urea cycle disorders. American Journal of Neuroradiology, 2003, 24, 1184-7.	2.4	72
116	A Case of Acute Disseminated Encephalomyelitis Presenting Hypersomnia With Decreased Hypocretin Level in Cerebrospinal Fluid. Journal of Child Neurology, 2002, 17, 537-539.	1.4	44
117	Effect of High Temperature on Ripening in Rice Plants.. Japanese Journal of Crop Science, 2002, 71, 102-109.	0.2	44
118	Retrocerebellar arachnoid cysts in siblings with mental retardation and undescended testis. Brain and Development, 2002, 24, 310-313.	1.1	10
119	Slit ventricle syndrome after cyst-peritoneal shunting for temporal arachnoid cyst in children â€“ a clinical entity difficult to detect on neuroimaging study. Brain and Development, 2002, 24, 776-779.	1.1	21
120	Nursery Method for Mechanical Transplanting of Giant-embryo Rice Cultivar 'Haiminori'.. Japanese Journal of Crop Science, 2002, 71, 76-83.	0.2	1
121	Empty sella in children as a key for diagnosis. Brain and Development, 2001, 23, 422-423.	1.1	19
122	Differences in the Rate of Seedling Emergence among Rice Cultivars under Low Soil-Moisture Conditions. Plant Production Science, 2001, 4, 94-102.	2.0	5
123	Effects of Dry Matter Production, Translocation of Nonstructural Carbohydrates and Nitrogen Application on Grain Filling in Rice Cultivar Takanari, a Cultivar Bearing a Large Number of Spikelets. Plant Production Science, 2001, 4, 173-183.	2.0	77
124	Q FEVER ENCEPHALITIS WITH CYTOKINE PROFILES IN SERUM AND CEREBROSPINAL FLUID. Pediatric Infectious Disease Journal, 2001, 20, 318-319.	2.0	15
125	Estimation of photoparoxysmal response elicited by half-field visual stimulation. NeuroReport, 2000, 11, 203-206.	1.2	3
126	Dipole Tracing Examination for the Electric Source of Photoparoxysmal Response Provoked by Half-Field Stimulation. Epilepsia, 2000, 41, 60-60.	5.1	1

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127	Empty sella syndrome in nevoid basal cell carcinoma syndrome. Brain and Development, 2000, 22, 272-274.	1.1	13
128	Comparative performance of magnetic resonance angiography and conventional angiography in moyamoya disease. Journal of Clinical Neuroscience, 2000, 7, 112-115.	1.5	20
129	Î³-Irradiation Deregulates Cell Cycle Control and Apoptosis in Nevoid Basal Cell Carcinoma Syndrome-derived Cells. Japanese Journal of Cancer Research, 1999, 90, 1351-1357.	1.7	14
130	Proteolipid protein gene duplications causing Pelizaeus-Merzbacher disease: Molecular mechanism and phenotypic manifestations. Annals of Neurology, 1999, 45, 624-632.	5.3	126
131	Neuroradiologic findings in glutaric aciduria type II. Pediatric Neurology, 1999, 20, 142-145.	2.1	24
132	Vacuolating leukoencephalopathy with subcortical cysts with late onset athetotic movements. Journal of the Neurological Sciences, 1999, 165, 90-93.	0.6	11
133	Proteolipid protein gene duplications causing Pelizaeusâ€Merzbacher disease: Molecular mechanism and phenotypic manifestations. Annals of Neurology, 1999, 45, 624-632.	5.3	2
134	Investigation of Electrical Focus and Generator Mechanism for Photoparoxysmal Discharges with a Dipole Tracing Method. Epilepsia, 1998, 39, 79-79.	5.1	0
135	Evaluation of magnetic resonance angiography with selective maximum intensity projection in patients with childhood moyamoya disease. European Journal of Paediatric Neurology, 1998, 2, 83-89.	1.6	24
136	Adolescent case of Alexander disease: MR imaging and MR spectroscopy. Pediatric Neurology, 1998, 18, 67-70.	2.1	27
137	Dichloroacetate treatment in Leigh syndrome caused by mitochondrial DNA mutation. Journal of the Neurological Sciences, 1997, 145, 83-86.	0.6	26
138	Longitudinal MR imaging and proton MR spectroscopy in herpes simplex encephalitis. Journal of the Neurological Sciences, 1997, 149, 99-102.	0.6	39
139	Availability of frequency-selective fat-Saturation pulse (fat-Sat) MRI in childhood optic neuritis. Pediatric Neurology, 1996, 14, 64-65.	2.1	2
140	Optic neuritis with silent cerebral lesions: Availability of FLAIR sequences. Pediatric Neurology, 1995, 12, 152-154.	2.1	7
141	Antiphospholipid antibody syndrome in childhood strokes. Pediatric Neurology, 1995, 13, 323-326.	2.1	42
142	Moyamoya syndrome in a patient with Down syndrome presenting with chorea. Pediatric Neurology, 1993, 9, 396-398.	2.1	51
143	Comparison of MRI white matter changes with neuropsychologic impairment in Cockayne syndrome. Pediatric Neurology, 1992, 8, 295-298.	2.1	18
144	Pelizaeus-Merzbacher disease: Cellular hypersensitivity to ultraviolet light. Brain and Development, 1992, 14, 44-47.	1.1	5

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145	Comparison of cellular sensitivity to UV killing with neuropsychological impairment in cockayne syndrome patients. Brain and Development, 1991, 13, 163-166.	1.1	10