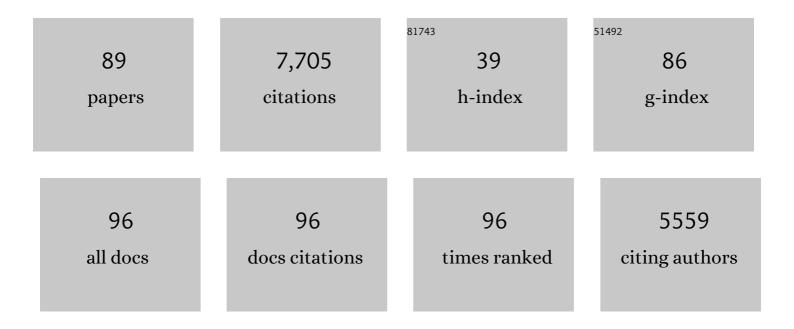
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Kleine-Levin syndrome is associated with birth difficulties and genetic variants in the <i>TRANK1</i> gene loci. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	3.3	26
2	Idiopathic hypersomnia: a homogeneous or heterogeneous disease?. Sleep Medicine, 2021, 80, 86-91.	0.8	17
3	2018 worldwide survey of health-care providers caring for patients with narcolepsy. Sleep Medicine, 2021, 82, 23-28.	0.8	4
4	Delayed sleep-wake phase disorder: Can polysomnography be useful?. Pediatric Neurology, 2021, 127, 28-31.	1.0	1
5	Developmental Language Disorder: Wake and Sleep Epileptiform Discharges and Co-morbid Neurodevelopmental Disorders. Brain Sciences, 2020, 10, 910.	1.1	3
6	Sleep-related rhythmic movements and rhythmic movement disorder beyond early childhood. Sleep Medicine, 2019, 64, 112-115.	0.8	20
7	Relapsing encephalopathy with cerebellar ataxia are caused by variants involving p.Arg756 in ATP1A3. European Journal of Paediatric Neurology, 2019, 23, 448-455.	0.7	33
8	The MSLT is Repeatable in Narcolepsy Type 1 But Not Narcolepsy Type 2: A Retrospective Patient Study. Journal of Clinical Sleep Medicine, 2018, 14, 65-74.	1.4	69
9	Childhood narcolepsy and autism spectrum disorders: four case reports. Sleep Medicine, 2018, 51, 167-170.	0.8	13
10	Higher body mass index in narcolepsy with cataplexy: lifelong experience. Sleep Medicine, 2017, 32, 277.	0.8	3
11	Cardiovascular fitness in narcolepsy is inversely related to sleepiness and the number of cataplexy episodes. Sleep Medicine, 2017, 34, 7-12.	0.8	11
12	Sleep in Neurological and Neurodevelopmental Disorders. , 2017, , 357-387.		0
13	Disorders Associated with Increased Sleepiness. , 2017, , 281-304.		2
14	Narcolepsy with cataplexy in patients aged over 60 years: a case-control study. Sleep Medicine, 2016, 26, 79-84.	0.8	20
15	Smoking Prevalence and Its Clinical Correlations in Patients with Narcolepsy-cataplexy. Prague Medical Report, 2016, 117, 81-89.	0.4	2
16	Stable or improved neurological manifestations during miglustat therapy in patients from the international disease registry for Niemann-Pick disease type C: an observational cohort study. Orphanet Journal of Rare Diseases, 2015, 10, 65.	1.2	83
17	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhood—a study of 155 patients. Orphanet Journal of Rare Diseases, 2015, 10, 123.	1.2	117
18	Cataplexy and Sleep Disorders in Niemann-Pick Type C Disease. Current Neurology and Neuroscience Reports, 2015, 15, 522.	2.0	27

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19	Antidepressants substantially affect basic REM sleep characteristics in narcolepsy-cataplexy patients. Neuroendocrinology Letters, 2015, 36, 430-3.	0.2	3
20	The Diagnosis and Treatment of Pediatric Narcolepsy. Current Neurology and Neuroscience Reports, 2014, 14, 469.	2.0	41
21	Nightmares in narcolepsy: underinvestigated symptom?. Sleep Medicine, 2014, 15, 967-972.	0.8	42
22	Childhood narcolepsy and H1N1 vaccination: stirring up a sleeping menace?. Sleep Medicine, 2014, 15, 159-160.	0.8	1
23	HLA DQB1*06:02 Negative Narcolepsy with Hypocretin/Orexin Deficiency. Sleep, 2014, 37, 1601-1608.	0.6	59
24	Childhood parasomnia – A disorder of sleep maturation?. European Journal of Paediatric Neurology, 2013, 17, 615-619.	0.7	26
25	Narcolepsy: clinical differences and association with other sleep disorders in different age groups. Journal of Neurology, 2013, 260, 767-775.	1.8	49
26	ImmunoChip Study Implicates Antigen Presentation to T Cells in Narcolepsy. PLoS Genetics, 2013, 9, e1003270.	1.5	206
27	Narcolepsy and pregnancy: a retrospective <scp>E</scp> uropean evaluation of 249 pregnancies. Journal of Sleep Research, 2013, 22, 496-512.	1.7	54
28	Alteration of the Circadian Clock in Children with Smith-Magenis Syndrome. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E312-E318.	1.8	35
29	Charcot–Marie–Tooth neuropathy due to a novel EGR2 gene mutation with mild phenotype – Usefulness of human mapping chip linkage analysis in a Czech family. Neuromuscular Disorders, 2012, 22, 742-746.	0.3	9
30	De novo mutations in ATP1A3 cause alternating hemiplegia of childhood. Nature Genetics, 2012, 44, 1030-1034.	9.4	345
31	Predictors of Hypocretin (Orexin) Deficiency in Narcolepsy Without Cataplexy. Sleep, 2012, 35, 1247-1255.	0.6	182
32	Decreased serum antioxidant capacity in patients with Wilson disease is associated with neurological symptoms. Journal of Inherited Metabolic Disease, 2012, 35, 541-548.	1.7	28
33	Alternating hemiplegia of childhood: Metabolic studies in the largest European series of patients. European Journal of Paediatric Neurology, 2012, 16, 10-14.	0.7	20
34	Clinical experience suggests that modafinil is an effective and safe treatment for paediatric narcolepsy. Journal of Sleep Research, 2012, 21, 481-483.	1.7	51
35	Total Antioxidative Capacity in Serum Correlates With the Phenotypic Manifestation of Wilson Disease. Gastroenterology, 2011, 140, S-939.	0.6	0
36	Sleep disorders in Wilson's disease. European Journal of Neurology, 2011, 18, 184-190.	1.7	56

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37	Diagnosing narcolepsy with cataplexy on history alone: challenging the International Classification of Sleep Disorders (ICSD-2) criteria. European Journal of Neurology, 2011, 18, 1017-1020.	1.7	12
38	Longâ€ŧerm followâ€up of Wilson Disease: natural history, treatment, mutations analysis and phenotypic correlation. Liver International, 2011, 31, 83-91.	1.9	114
39	Common variants in P2RY11 are associated with narcolepsy. Nature Genetics, 2011, 43, 66-71.	9.4	215
40	Clinical features of childhood narcolepsy. Can cataplexy be foretold?. European Journal of Paediatric Neurology, 2011, 15, 320-325.	0.7	29
41	Genome-Wide Association Study Identifies Novel Restless Legs Syndrome Susceptibility Loci on 2p14 and 16q12.1. PLoS Genetics, 2011, 7, e1002171.	1.5	163
42	Genetic disorders and sleepiness. , 2011, , 335-350.		4
43	Gabapentin in the treatment of dementia-associated nocturnal agitation. Medical Science Monitor, 2011, 17, CS149-CS151.	0.5	6
44	Anti-Tribbles Homolog 2 (TRIB2) Autoantibodies in Narcolepsy are Associated with Recent Onset of Cataplexy. Sleep, 2010, 33, 869-874.	0.6	113
45	Interictal epileptiform discharges andÂphasic phenomena ofÂREM sleep. Epileptic Disorders, 2010, 12, 217-221.	0.7	13
46	Evidence of a non-progressive course of alternating hemiplegia of childhood: study of a large cohort of children and adults. Brain, 2010, 133, 3598-3610.	3.7	126
47	Sodium oxybate is an effective and safe treatment for narcolepsy. Sleep Medicine, 2010, 11, 105-106.	0.8	29
48	Olfactory dysfunction in narcolepsy with and without cataplexy. Sleep Medicine, 2010, 11, 558-561.	0.8	18
49	Sleep disorders and daytime sleepiness in children with attention-deficit/hyperactivity disorder: A two-night polysomnographic study with a multiple sleep latency test. Sleep Medicine, 2010, 11, 922-928.	0.8	71
50	Assessment of pregnancy outcomes in Czech and Slovak women with narcolepsy. Medical Science Monitor, 2010, 16, SR35-40.	0.5	16
51	Decreased CSF Histamine in Narcolepsy With and Without Low CSF Hypocretin-1 in Comparison to Healthy Controls. Sleep, 2009, 32, 175-180.	0.6	142
52	Elevated Anti-Streptococcal Antibodies in Patients with Recent Narcolepsy Onset. Sleep, 2009, 32, 979-983.	0.6	311
53	Replication of restless legs syndrome loci in three European populations. Journal of Medical Genetics, 2009, 46, 315-318.	1.5	78
54	Narcolepsy is strongly associated with the T-cell receptor alpha locus. Nature Genetics, 2009, 41, 708-711.	9.4	445

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55	Narcolepsy in childhood. Sleep Medicine Reviews, 2009, 13, 169-180.	3.8	102
56	Arousals in nocturnal groaning. Sleep Medicine, 2009, 10, 1051-1055.	0.8	20
57	Does age at the onset of narcolepsy influence the course and severity of the disease?. Sleep Medicine, 2009, 10, 967-972.	0.8	51
58	Suggestive evidence for linkage for restless legs syndrome on chromosome 19p13. Neurogenetics, 2008, 9, 75-82.	0.7	61
59	PTPRD (protein tyrosine phosphatase receptor type delta) is associated with restless legs syndrome. Nature Genetics, 2008, 40, 946-948.	9.4	252
60	Circadian rhythm in salivary melatonin in narcoleptic patients. Neuroscience Letters, 2008, 437, 162-164.	1.0	12
61	A comparison of polysomnographic and actigraphic evaluation of periodic limb movements in sleep. Neurological Research, 2008, 30, 234-238.	0.6	23
62	REM behavior disorder (RBD) can be one of the first symptoms of childhood narcolepsy. Sleep Medicine, 2007, 8, 784-786.	0.8	115
63	Family-based association study of the restless legs syndrome loci 2 and 3 in a European population. Movement Disorders, 2007, 22, 207-212.	2.2	31
64	Genetics of restless legs syndrome (RLS): State-of-the-art and future directions. Movement Disorders, 2007, 22, S449-S458.	2.2	73
65	Rhythmic Movement Disorder in Sleep Persisting into Childhood and Adulthood. Sleep, 2005, 28, 851-857.	0.6	94
66	Hypocretin deficiency in Prader-Willi syndrome. European Journal of Neurology, 2005, 12, 70-72.	1.7	88
67	Deficient Orexinergic Modulation in Narcolepsy May Influence Heart Rate Variability in Sleep. Die defekte orexinergische Modulation in Narkolepsie kann die Herzschlagvariabilitat im Schlaf beeinflussen. Somnologie, 2005, 9, 15-20.	0.9	2
68	NREM sleep alterations in narcolepsy/cataplexy. Clinical Neurophysiology, 2005, 116, 2675-2684.	0.7	68
69	Sleep and Fasciculations in Amyotrophic Lateral Sclerosis. Schlaf und Faszikulationen bei amyotropher Lateralsklerose. Somnologie, 2004, 8, 25-30.	0.9	5
70	559 CNS impairment in the children from patients with Wilson disease. Journal of Hepatology, 2004, 40, 164.	1.8	0
71	Sleep Disturbances and Hypocretin Deficiency in Niemann-Pick Disease Type C. Sleep, 2003, 26, 427-430.	0.6	104
72	Chapter 26 Excessive daytime sleepiness. Supplements To Clinical Neurophysiology, 2002, 54, 175-179.	2.1	1

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73	Increased REM Density in Narcolepsy-Cataplexy and the Polysymptomatic Form of Idiopathic Hypersomnia. Sleep, 2001, 24, 707-711.	0.6	30
74	Electromagnetic field of mobile phones affects visual event related potential in patients with narcolepsy. Bioelectromagnetics, 2001, 22, 519-528.	0.9	30
75	Low cerebrospinal fluid hypocretin (orexin) and altered energy homeostasis in human narcolepsy. Annals of Neurology, 2001, 50, 381-388.	2.8	451
76	Electromagnetic field of mobile phones affects visual event related potential in patients with narcolepsy. Bioelectromagnetics, 2001, 22, 519-528.	0.9	1
77	Chapter 39 EEG changes and epilepsy in developmental dysphasia. Supplements To Clinical Neurophysiology, 2000, 53, 271-274.	2.1	5
78	Chapter 51 Excessive sleepiness: clinical aspects. Supplements To Clinical Neurophysiology, 2000, 53, 362-365.	2.1	1
79	A mutation in a case of early onset narcolepsy and a generalized absence of hypocretin peptides in human narcoleptic brains. Nature Medicine, 2000, 6, 991-997.	15.2	1,945
80	Narcolepsy in Children. Sleep, 1994, 17, S17-S20.	0.6	140
81	Alternating hemiplegia in childhood: a cross-sectional study. Brain and Development, 1994, 16, 189-194.	0.6	15
82	A role of autoimmunity in the etiopathogenesis of Landau-Kleffner syndrome?. Brain and Development, 1992, 14, 342-345.	0.6	47
83	An EEG study of Wilson's disease. Findings in patients and heterozygous relatives. Electroencephalography and Clinical Neurophysiology, 1986, 64, 191-198.	0.3	16
84	An Alternative to the Multiple Sleep Latency Test for Determining Sleepiness in Narcolepsy and Hypersomnia: Polygraphic Score of Sleepiness. Sleep, 1986, 9, 243-245.	0.6	41
85	Multiple sulphatase deficiency in homozygotic twins. Journal of Inherited Metabolic Disease, 1984, 7, 38-40.	1.7	17
86	Biochemical and clinical changes in Wilson's disease heterozygotes. Journal of Inherited Metabolic Disease, 1984, 7, 41-45.	1.7	13
87	COMPUTED TOMOGRAPHY IN WILSON'S DISEASE. Journal of Computer Assisted Tomography, 1983, 7, 187.	0.5	0
88	Life Effects of Narcolepsy in 180 Patients from North America, Asia and Europe Compared to Matched Controls. Canadian Journal of Neurological Sciences, 1981, 8, 299-304.	0.3	211
89	REM sleep and NREM sleep in narcolepsy and hypersomnia. Electroencephalography and Clinical Neurophysiology, 1969, 26, 176-182.	0.3	51