Anna CzÅ, onkowska

List of Publications by Year in descending order

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312 papers 17,706 citations

25034 57 h-index 17592 121 g-index

325 all docs 325 docs citations

times ranked

325

19892 citing authors

#	Article	lF	CITATIONS
1	Risk factors for ischaemic and intracerebral haemorrhagic stroke in 22 countries (the INTERSTROKE) Tj ETQq1 1 (0.784314 r 13.7	gBT_/Qverloc 2,565
2	Global and regional effects of potentially modifiable risk factors associated with acute stroke in 32 countries (INTERSTROKE): a case-control study. Lancet, The, 2016, 388, 761-775.	13.7	1,414
3	The benefits and harms of intravenous thrombolysis with recombinant tissue plasminogen activator within 6 h of acute ischaemic stroke (the third international stroke trial [IST-3]): a randomised controlled trial. Lancet, The, 2012, 379, 2352-2363.	13.7	1,018
4	Rivaroxaban for Stroke Prevention after Embolic Stroke of Undetermined Source. New England Journal of Medicine, 2018, 378, 2191-2201.	27.0	730
5	Siponimod versus placebo in secondary progressive multiple sclerosis (EXPAND): a double-blind, randomised, phase 3 study. Lancet, The, 2018, 391, 1263-1273.	13.7	684
6	Wilson disease. Nature Reviews Disease Primers, 2018, 4, 21.	30.5	466
7	The Inflammatory Reaction Following 1-Methyl-4-phenyl-1,2,3,6-tetrahydropyridine Intoxication in Mouse. Experimental Neurology, 1999, 156, 50-61.	4.1	338
8	Tranexamic acid for hyperacute primary IntraCerebral Haemorrhage (TICH-2): an international randomised, placebo-controlled, phase 3 superiority trial. Lancet, The, 2018, 391, 2107-2115.	13.7	309
9	Teriflunomide versus subcutaneous interferon beta-1a in patients with relapsing multiple sclerosis: a randomised, controlled phase 3 trial. Multiple Sclerosis Journal, 2014, 20, 705-716.	3.0	295
10	Microglial and astrocytic involvement in a murine model of Parkinson's disease induced by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP). Immunopharmacology, 1998, 39, 167-180.	2.0	261
11	Frequency and Prognostic Value of Cognitive Disorders in Stroke Patients. Dementia and Geriatric Cognitive Disorders, 2008, 26, 356-363.	1.5	255
12	Influence of Gender on Baseline Features and Clinical Outcomes among 17,370 Patients with Confirmed Ischaemic Stroke in the International Stroke Trial. Neuroepidemiology, 2005, 24, 123-128.	2.3	251
13	Microglial Reaction in MPTP (1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine) Induced Parkinson's Disease Mice Model. Experimental Neurology, 1996, 5, 137-143.	1.7	245
14	Late-Onset Wilson's Disease. Gastroenterology, 2007, 132, 1294-1298.	1.3	227
15	Effects of long-term treatment in Wilson's disease withd-penicillamine and zinc sulphate. Journal of Neurology, 1996, 243, 269-273.	3.6	172
16	MicroRNAs as Diagnostic and Prognostic Biomarkers in Ischemic Stroke—A Comprehensive Review and Bioinformatic Analysis. Cells, 2018, 7, 249.	4.1	131
17	Frameshift and nonsense mutations in the gene for ATPase7B are associated with severe impairment of copper metabolism and with an early clinical manifestation of Wilson's disease. Clinical Genetics, 2005, 68, 524-532.	2.0	124
18	Gender differences in Wilson's disease. Journal of the Neurological Sciences, 2012, 312, 31-35.	0.6	119

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19	Early neurological worsening in patients with Wilson's disease. Journal of the Neurological Sciences, 2015, 355, 162-167.	0.6	116
20	Restenosis and risk of stroke after stenting or endarterectomy for symptomatic carotid stenosis in the International Carotid Stenting Study (ICSS): secondary analysis of a randomised trial. Lancet Neurology, The, 2018, 17, 587-596.	10.2	114
21	Dâ€penicillamine versus zinc sulfate as firstâ€line therapy for Wilson's disease. European Journal of Neurology, 2014, 21, 599-606.	3.3	113
22	Estrogen and Cytokines Production - The Possible Cause of Gender Differences in Neurological Diseases. Current Pharmaceutical Design, 2005, 11, 1017-1030.	1.9	112
23	Bis-choline tetrathiomolybdate in patients with Wilson's disease: an open-label, multicentre, phase 2 study. The Lancet Gastroenterology and Hepatology, 2017, 2, 869-876.	8.1	110
24	Age and Sex but Not ATP7B Genotype Effectively Influence the Clinical Phenotype of Wilson Disease. Hepatology, 2019, 69, 1464-1476.	7.3	110
25	Risk factors for falls in stroke patients during inpatient rehabilitation. Clinical Rehabilitation, 2009, 23, 176-188.	2.2	107
26	Dexamethasone protects against dopaminergic neurons damage in a mouse model of Parkinson's disease. International Immunopharmacology, 2004, 4, 1307-1318.	3.8	106
27	Factors Influencing In-Hospital Delay in Treatment With Intravenous Thrombolysis. Stroke, 2012, 43, 1578-1583.	2.0	104
28	Intracerebral Hematoma Morphologic Appearance on Noncontrast Computed Tomography Predicts Significant Hematoma Expansion. Stroke, 2015, 46, 3111-3116.	2.0	103
29	Gender Differences in Neurological Disease: Role of Estrogens and Cytokines. Endocrine, 2006, 29, 243-256.	2.2	98
30	Microglia and microglia-derived brain macrophages in culture: generation from axotomized rat facial nuclei, identification and characterization in vitro. Brain Research, 1989, 492, 1-14.	2.2	97
31	Practice patterns and outcomes after stroke across countries at different economic levels (INTERSTROKE): an international observational study. Lancet, The, 2018, 391, 2019-2027.	13.7	96
32	Global Impact of COVID-19 on Stroke Care and IV Thrombolysis. Neurology, 2021, 96, e2824-e2838.	1.1	95
33	Evaluation of the Unified Wilson's Disease Rating Scale (UWDRS) in German patients with treated Wilson's disease. Movement Disorders, 2008, 23, 54-62.	3.9	94
34	Transcranial Magnetic Stimulation Combined With Physiotherapy in Rehabilitation of Poststroke Hemiparesis. Neurorehabilitation and Neural Repair, 2012, 26, 1072-1079.	2.9	94
35	Wilson's diseaseâ€"cause of mortality in 164 patients during 1992â€"2003 observation period. Journal of Neurology, 2005, 252, 698-703.	3.6	89
36	The third international stroke trial (IST-3) of thrombolysis for acute ischaemic stroke. Trials, 2008, 9, 37.	1.6	86

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37	Cognitive functioning in neurologically symptomatic and asymptomatic forms of Wilson's disease. Movement Disorders, 2002, 17, 1077-1083.	3.9	83
38	Rivaroxaban for secondary stroke prevention in patients with embolic strokes of undetermined source: Design of the NAVIGATE ESUS randomized trial. European Stroke Journal, 2016, 1, 146-154.	5.5	83
39	Genetic control of multiple sclerosis: Increased production of lymphotoxin and tumor necrosis factor-? by HLA-DR2+ T cells. Annals of Neurology, 1995, 38, 723-730.	5.3	81
40	Transcranial Magnetic Stimulation Combined with Speech and Language Training in Early Aphasia Rehabilitation: A Randomized Double-Blind Controlled Pilot Study. Topics in Stroke Rehabilitation, 2013, 20, 250-261.	1.9	81
41	Inflammation and gliosis in neurological diseases – clinical implications. Journal of Neuroimmunology, 2011, 231, 78-85.	2.3	78
42	Hepatobiliary malignancies in Wilson disease. Liver International, 2015, 35, 1615-1622.	3.9	78
43	The Relation of the Brain-Derived Neurotrophic Factor with MicroRNAs in Neurodegenerative Diseases and Ischemic Stroke. Molecular Neurobiology, 2021, 58, 329-347.	4.0	78
44	Brain metal accumulation in Wilson's disease. Journal of the Neurological Sciences, 2013, 329, 55-58.	0.6	77
45	Wilson Disease and Other Neurodegenerations with Metal Accumulations. Neurologic Clinics, 2015, 33, 175-204.	1.8	76
46	Immunological observations on patients with acute cerebral vascular disease. Journal of the Neurological Sciences, 1979, 43, 455-464.	0.6	74
47	Alteplase for Acute Ischemic Stroke. Stroke, 2015, 46, 746-756.	2.0	74
48	Monozygotic female twins discordant for phenotype of Wilson's disease. Movement Disorders, 2009, 24, 1066-1069.	3.9	72
49	Compliant treatment with antiâ€copper agents prevents clinically overt <scp>W</scp> ilson's disease in preâ€symptomatic patients. European Journal of Neurology, 2014, 21, 332-337.	3.3	70
50	Immune processes in the pathogenesis of Parkinson's disease - a potential role for microglia and nitric oxide. Medical Science Monitor, 2002, 8, RA165-77.	1.1	69
51	p.H1069Q mutation inATP7B and biochemical parameters of copper metabolism and clinical manifestation of Wilson's disease. Movement Disorders, 2006, 21, 245-248.	3.9	68
52	Psychiatric manifestations in Wilson's disease: possibilities and difficulties for treatment. Therapeutic Advances in Psychopharmacology, 2018, 8, 199-211.	2.7	68
53	Late onset Wilson's disease: Therapeutic implications. Movement Disorders, 2008, 23, 896-898.	3.9	67
54	Effect of Low-Frequency Repetitive Transcranial Magnetic Stimulation on Naming Abilities in Early-Stroke Aphasic Patients: A Prospective, Randomized, Double-Blind Sham-Controlled Study. Scientific World Journal, The, 2012, 2012, 1-8.	2.1	66

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55	Effects of Repeated Anodal tDCS Coupled With Cognitive Training for Patients With Severe Traumatic Brain Injury. Journal of Head Trauma Rehabilitation, 2014, 29, E20-E29.	1.7	65
56	New approach to the rehabilitation of post-stroke focal cognitive syndrome: Effect of levodopa combined with speech and language therapy on functional recovery from aphasia. Journal of the Neurological Sciences, 2009, 283, 214-218.	0.6	61
57	Spot Sign Number Is the Most Important Spot Sign Characteristic for Predicting Hematoma Expansion Using First-Pass Computed Tomography Angiography. Stroke, 2013, 44, 972-977.	2.0	61
58	Role of Preexisting Disability in Patients Treated With Intravenous Thrombolysis for Ischemic Stroke. Stroke, 2014, 45, 770-775.	2.0	60
59	Brain iron accumulation in Wilson disease: a <i>post mortem</i> 7 Tesla MRI – histopathological study. Neuropathology and Applied Neurobiology, 2017, 43, 514-532.	3.2	60
60	The International Stroke Trial database. Trials, 2011, 12, 101.	1.6	59
61	Neurologic impairment in Wilson disease. Annals of Translational Medicine, 2019, 7, S64-S64.	1.7	58
62	Persistence with treatment in patients with Wilson disease. Neurologia I Neurochirurgia Polska, 2010, 44, 260-263.	1.2	54
63	Early stroke-related deep venous thrombosis: risk factors and influence on outcome. Journal of Thrombosis and Thrombolysis, 2011, 32, 96-102.	2.1	54
64	Wilson disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2017, 142, 101-119.	1.8	52
65	Association of IL1A, IL1B, ILRN, IL6, IL10 and TNF-α polymorphisms with risk and clinical course of multiple sclerosis in a Polish population. Journal of Neuroimmunology, 2011, 236, 87-92.	2.3	51
66	Venous Phase of Computed Tomography Angiography Increases Spot Sign Detection, but Intracerebral Hemorrhage Expansion Is Greater in Spot Signs Detected in Arterial Phase. Stroke, 2014, 45, 734-739.	2.0	51
67	Intravenous tranexamic acid for hyperacute primary intracerebral hemorrhage: Protocol for a randomized, placebo-controlled trial. International Journal of Stroke, 2016, 11, 683-694.	5.9	50
68	Ultraearly hematoma growth in active intracerebral hemorrhage. Neurology, 2016, 87, 357-364.	1.1	50
69	Peripheral Blood <i>MCEMP1</i> Gene Expression as a Biomarker for Stroke Prognosis. Stroke, 2016, 47, 652-658.	2.0	48
70	Genetic variability in the methylenetetrahydrofolate reductase gene (MTHFR) affects clinical expression of Wilson's disease. Journal of Hepatology, 2011, 55, 913-919.	3.7	47
71	Wilson disease – currently used anticopper therapy. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2017, 142, 181-191.	1.8	47
72	Elevated Levels of Anti-Heat Shock Protein Antibodies in Patients with Cerebral Ischemia. Cerebrovascular Diseases, 2001, 12, 235-239.	1.7	46

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73	Characterization of Patients with Embolic Strokes of Undetermined Source in the NAVIGATE ESUS Randomized Trial. Journal of Stroke and Cerebrovascular Diseases, 2018, 27, 1673-1682.	1.6	46
74	Intravenous Alteplase in Ischemic Stroke Patients not Fully Adhering to the Current Drug License in Central and Eastern Europe. International Journal of Stroke, 2012, 7, 615-622.	5.9	44
75	Wilson's disease: update on pathogenesis, biomarkers and treatments. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 1053-1061.	1.9	44
76	Characteristics of a newly diagnosed Polish cohort of patients with neurological manifestations of Wilson disease evaluated with the Unified Wilson's Disease Rating Scale. BMC Neurology, 2018, 18, 34.	1.8	43
77	Semiquantitative Scale for Assessing Brain MRI Abnormalities in Wilson Disease: A Validation Study. Movement Disorders, 2020, 35, 994-1001.	3.9	43
78	Apolipoprotein E gene (APOE) genotype in Wilson's disease: Impact on clinical presentation. Parkinsonism and Related Disorders, 2012, 18, 367-369.	2.2	42
79	The effect of gender on brain MRI pathology in Wilson's disease. Metabolic Brain Disease, 2013, 28, 69-75.	2.9	42
80	Anodal transcranial direct current stimulation in early rehabilitation of patients with post-stroke non-fluent aphasia: A randomized, double-blind, sham-controlled pilot study. Restorative Neurology and Neuroscience, 2013, 31, 761-771.	0.7	42
81	Cyclooxygenases mRNA and protein expression in striata in the experimental mouse model of Parkinson's disease induced by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine administration to mouse. Brain Research, 2004, 1019, 144-151.	2.2	41
82	Anticardiolipin antibodies are an independent risk factor for ischemic stroke. Neurological Research, 1999, 21, 653-657.	1.3	40
83	Underfunding of Stroke Research. Stroke, 2004, 35, 2368-2371.	2.0	40
84	Symptomatic treatment of neurologic symptoms in Wilson disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2017, 142, 211-223.	1.8	39
85	Age- and sex-differences in the nitric oxide synthase expression and dopamine concentration in the murine model of Parkinson's disease induced by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine. Brain Research, 2009, 1261, 7-19.	2.2	38
86	Update on the third international stroke trial (IST-3) of thrombolysis for acute ischaemic stroke and baseline features of the 3035 patients recruited. Trials, 2011, 12, 252.	1.6	38
87	Wholeâ€exome sequencing identifies novel pathogenic variants across the <i>ATP7B </i> gene and some modifiers of Wilson's disease phenotype. Liver International, 2019, 39, 177-186.	3.9	38
88	Clinical manifestations of Wilson disease in organs other than the liver and brain. Annals of Translational Medicine, 2019, 7, S62-S62.	1.7	38
89	BDNF A196G and C270T gene polymorphisms and susceptibility to multiple sclerosis in the polish population. Gender differences. Journal of Neuroimmunology, 2008, 193, 170-172.	2.3	36
90	Hyperperfusion Syndrome after Carotid Endarterectomy and Carotid Stenting. Cerebrovascular Diseases, 2013, 35, 531-537.	1.7	36

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91	Effect of medical complications on the after-stroke rehabilitation outcome. NeuroRehabilitation, 2017, 40, 223-232.	1.3	36
92	Middle-aged heterozygous carriers of Wilson's disease do not present with significant phenotypic deviations related to copper metabolism. Journal of Genetics, 2010, 89, 463-467.	0.7	35
93	The influence of AAV2-mediated gene transfer of human IL-10 on neurodegeneration and immune response in a murine model of Parkinson's disease. Pharmacological Reports, 2014, 66, 660-669.	3.3	35
94	Diverse attention deficits in patients with neurologically symptomatic and asymptomatic Wilson's disease Neuropsychology, 2015, 29, 25-30.	1.3	34
95	Wilson diseaseâ€"treatment perspectives. Annals of Translational Medicine, 2019, 7, S68-S68.	1.7	34
96	Unified Wilson's Disease Rating Scale - a proposal for the neurological scoring of Wilson's disease patients. Neurologia I Neurochirurgia Polska, 2007, 41, 1-12.	1.2	34
97	MR spectroscopy in monitoring the treatment of Wilson's disease patients. Movement Disorders, 2008, 23, 1560-1566.	3.9	33
98	Dynamics of expression of the mRNA for cytokines and inducible nitric synthase in a murine model of the Parkinson's disease. Acta Neurobiologiae Experimentalis, 2003, 63, 117-26.	0.7	33
99	No effects of anodal transcranial direct stimulation on language abilities in early rehabilitation of post-stroke aphasic patients. Neurologia I Neurochirurgia Polska, 2013, 47, 414-422.	1.2	32
100	Symptomatic copper deficiency in three Wilson's disease patients treated with zinc sulphate. Neurologia I Neurochirurgia Polska, 2014, 48, 214-218.	1.2	32
101	Glyceryl Trinitrate for Acute Intracerebral Hemorrhage. Stroke, 2016, 47, 44-52.	2.0	32
102	Epigenomic signatures in liver and blood of Wilson disease patients include hypermethylation of liver-specific enhancers. Epigenetics and Chromatin, 2019, 12, 10.	3.9	32
103	Oxfordshire Community Stroke Project Clinical Stroke Syndrome and Appearances of Tissue and Vascular Lesions on Pretreatment CT in Hyperacute Ischemic Stroke Among the First 510 Patients in the Third International Stroke Trial (IST-3). Stroke, 2009, 40, 743-748.	2.0	31
104	BDNF â^270 C>T polymorphisms might be associated with stroke type and BDNF â^2196 G>A corresponds to early neurological deficit in hemorrhagic stroke. Journal of Neuroimmunology, 2012, 249, 71-75.	2.3	31
105	Encephalopathy in Wilson Disease: Copper Toxicity or Liver Failure?. Journal of Clinical and Experimental Hepatology, 2015, 5, S88-S95.	0.9	31
106	The sunflower cataract in Wilson's disease: pathognomonic sign or rare finding?. Acta Neurologica Belgica, 2016, 116, 325-328.	1.1	31
107	Late Onset of Wilson's Disease. Archives of Neurology, 1981, 38, 729.	4.5	30
108	The Activity of Malignancy May Determine Stroke Pattern in Cancer Patients. Journal of Stroke and Cerebrovascular Diseases, 2015, 24, 778-783.	1.6	30

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109	Persistence with treatment for Wilson disease: a retrospective study. BMC Neurology, 2019, 19, 278.	1.8	30
110	Long Non-coding RNAs as Promising Therapeutic Approach in Ischemic Stroke: a Comprehensive Review. Molecular Neurobiology, 2021, 58, 1664-1682.	4.0	30
111	Immunological changes in the MPTP-induced Parkinson's disease mouse model. Journal of Neuroimmunology, 1993, 42, 33-37.	2.3	29
112	Acute Ischemic Stroke Care and Outcome in Centers Participating in the Polish National Stroke Prevention and Treatment Registry. Stroke, 2006, 37, 1837-1843.	2.0	29
113	Neurological presentation of Wilson's disease in a patient after liver transplantation. Movement Disorders, 2008, 23, 743-746.	3.9	29
114	Cerebral vasomotor reactivity in neurodegenerative diseases. Neurologia I Neurochirurgia Polska, 2016, 50, 455-462.	1.2	29
115	Clinical features of hemolysis, elevated liver enzymes, and low platelet count syndrome in undiagnosed Wilson disease: report of two cases. Archives of Gynecology and Obstetrics, 2010, 281, 129-134.	1.7	28
116	Effect of human interleukin-10 on the expression of nitric oxide synthases in the MPTP-based model of Parkinson's disease. Pharmacological Reports, 2013, 65, 44-49.	3.3	28
117	Other organ involvement and clinical aspects of Wilson disease. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2017, 142, 157-169.	1.8	28
118	Does brain degeneration in Wilson disease involve not only copper but also iron accumulation?. Neurologia I Neurochirurgia Polska, 2013, 47, 542-546.	1.2	27
119	Association between BDNF-196 G>A and BDNF-270 C>T polymorphisms, BDNF concentration, and rTMS-supported long-term rehabilitation outcome after ischemic stroke. NeuroRehabilitation, 2013, 32, 573-582.	1.3	27
120	Routine serum C-reactive protein and stroke outcome after intravenous thrombolysis. Acta Neurologica Scandinavica, 2014, 130, 305-311.	2.1	27
121	Safety of Statin Pretreatment in Intravenous Thrombolysis for Acute Ischemic Stroke. Stroke, 2015, 46, 2681-2684.	2.0	27
122	Intravenous thrombolysis for ischemic stroke in the golden hour: propensity-matched analysis from the SITS-EAST registry. Journal of Neurology, 2017, 264, 912-920.	3.6	27
123	Fibrin clot characteristics in acute ischaemic stroke patients treated with thrombolysis: the impact on clinical outcome. Thrombosis and Haemostasis, 2017, 117, 1440-1447.	3.4	27
124	Brain volume is related to neurological impairment and to copper overload in Wilson's disease. Neurological Sciences, 2019, 40, 2089-2095.	1.9	27
125	Predictors of Recurrent Ischemic Stroke in Patients with Embolic Strokes of Undetermined Source and Effects of Rivaroxaban Versus Aspirin According to Risk Status: The NAVIGATE ESUS Trial. Journal of Stroke and Cerebrovascular Diseases, 2019, 28, 2273-2279.	1.6	27
126	Influence of Age and Gender on Cytokine Expression in a Murine Model of Parkinson's Disease. NeuroImmunoModulation, 2007, 14, 255-265.	1.8	26

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127	Association of MMP1, MMP3, MMP9, and MMP12 polymorphisms with risk and clinical course of multiple sclerosis in a Polish population. Journal of Neuroimmunology, 2009, 214, 113-117.	2.3	26
128	Optical coherence tomography and electrophysiology of retinal and visual pathways in Wilson's disease. Metabolic Brain Disease, 2016, 31, 405-415.	2.9	26
129	Accuracy of the radioactive copper incorporation test in the diagnosis of Wilson disease. Liver International, 2018, 38, 1860-1866.	3.9	26
130	Metabolomics profiles of patients with Wilson disease reveal a distinct metabolic signature. Metabolomics, 2019, 15, 43.	3.0	26
131	Infections Up to 76ÂDays After Stroke Increase Disability and Death. Translational Stroke Research, 2017, 8, 541-548.	4.2	25
132	Variations in knowledge, awareness and treatment of hypertension and stroke risk by country income level. Heart, 2021, 107, 282-289.	2.9	25
133	Lenticular nucleus hyperechogenicity in Wilson's disease reflects local copper, but not iron accumulation. Journal of Neural Transmission, 2014, 121, 1273-1279.	2.8	24
134	Severe disease exacerbations in patients with multiple sclerosis after discontinuing fingolimod. Neurologia I Neurochirurgia Polska, 2017, 51, 156-162.	1.2	24
135	Pulse Pressure – Independent Predictor of Poor Early Outcome and Mortality following Ischemic Stroke. Cerebrovascular Diseases, 2009, 27, 187-192.	1.7	23
136	Abnormal antisaccades and smooth pursuit eye movements in patients with Wilson's disease. Movement Disorders, 2008, 23, 2067-2073.	3.9	22
137	Incidence and case fatality rates of first-ever stroke – comparison of data from two prospective population-based studies conducted in Warsaw. Neurologia I Neurochirurgia Polska, 2011, 45, 207-212.	1.2	22
138	Infections and Ischemic Stroke Outcome. Neurology Research International, 2011, 2011, 1-8.	1.3	22
139	Influence of <i>IL-1RN </i> Intron 2 Variable Number of Tandem Repeats (VNTR) Polymorphism on the Age at Onset of Neuropsychiatric Symptoms in Wilson's Disease. International Journal of Neuroscience, 2011, 121, 8-15.	1.6	22
140	Concordance rates of Wilson's disease phenotype among siblings. Journal of Inherited Metabolic Disease, 2014, 37, 131-135.	3.6	22
141	Polymorphisms of metal transporter genes DMT1 and ATP7A in Wilson's disease. Journal of Trace Elements in Medicine and Biology, 2014, 28, 8-12.	3.0	22
142	Impact of <i>BDNF </i> -196 G> A and <i>BDNF </i> -270 C> T Polymorphisms on Stroke Rehabilitation Outcome: Sex and Age Differences. Topics in Stroke Rehabilitation, 2014, 21, S33-S41.	1.9	22
143	Evolution and novel radiological changes of neurodegeneration associated with mutations in C19orf12. Parkinsonism and Related Disorders, 2017, 39, 71-76.	2.2	22
144	Optical coherence tomography as a marker of neurodegeneration in patients with Wilson's disease. Acta Neurologica Belgica, 2017, 117, 867-871.	1.1	22

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145	Dysregulated Choline, Methionine, and Aromatic Amino Acid Metabolism in Patients with Wilson Disease: Exploratory Metabolomic Profiling and Implications for Hepatic and Neurologic Phenotypes. International Journal of Molecular Sciences, 2019, 20, 5937.	4.1	22
146	Leukoaraiosis and stroke outcome. Journal of Stroke and Cerebrovascular Diseases, 2002, 11, 336-340.	1.6	21
147	Agreement between the clinical Oxfordshire Community Stroke Project classification and CT findings in Poland. European Journal of Neurology, 2004, 11, 91-96.	3.3	21
148	Polymorphisms in the factor VII gene and ischemic stroke in young adults. Blood Coagulation and Fibrinolysis, 2010, 21, 442-447.	1.0	21
149	WTX101 – an investigational drug for the treatment of Wilson disease. Expert Opinion on Investigational Drugs, 2018, 27, 561-567.	4.1	21
150	Deep Venous Thrombosis in Acute Stroke Patients. Clinical and Applied Thrombosis/Hemostasis, 2012, 18, 258-264.	1.7	20
151	Self-Rated Emotional Functioning of Patients With Neurological or Asymptomatic Form of Wilson's Disease. Clinical Neuropsychologist, 2003, 17, 367-373.	2.3	19
152	Lack of experience of intravenous thrombolysis for acute ischaemic stroke does not influence the proportion of patients treated. Emergency Medicine Journal, 2007, 24, 96-99.	1.0	19
153	Management of ischemic stroke in Central and Eastern Europe. International Journal of Stroke, 2015, 10, 125-127.	5.9	19
154	The accuracy of prehospital diagnosis of acute cerebrovascular accidents: an observational study. Archives of Medical Science, 2015, 3, 530-535.	0.9	19
155	Intravenous Thrombolysis for Stroke Recurring Within 3 Months From the Previous Event. Stroke, 2015, 46, 3184-3189.	2.0	19
156	Immunological observations on patients with Wilson's disease. Journal of the Neurological Sciences, 1976, 29, 411-421.	0.6	18
157	Knowledge of Risk Factors and Stroke Symptoms among Nonstroke Patients. European Neurology, 2012, 67, 220-225.	1.4	18
158	Families with Wilson's disease in subsequent generations: Clinical and genetic analysis. Movement Disorders, 2014, 29, 1828-1832.	3.9	18
159	Small intracerebral hemorrhages have a low spot sign prevalence and are less likely to expand. International Journal of Stroke, 2016, 11, 191-197.	5.9	18
160	A heterozygous mutation in GOT1 is associated with familial macro-aspartate aminotransferase. Journal of Hepatology, 2017, 67, 1026-1030.	3.7	18
161	Epigenetic changes of the thioredoxin system in the tx-j mouse model and in patients with Wilson disease. Human Molecular Genetics, 2018, 27, 3854-3869.	2.9	18
162	High-Sensitivity Cardiac Troponin T for Risk Stratification in Patients With Embolic Stroke of Undetermined Source. Stroke, 2020, 51, 2386-2394.	2.0	18

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163	High dose of intravenously given glucocorticosteroids decrease IL-8 production by monocytes in multiple sclerosis patients treated during relapse. Journal of Neuroimmunology, 2006, 176, 134-140.	2.3	17
164	Informed consent for clinical trials in acute coronary syndromes and stroke following the European Clinical Trials Directive: investigators' experiences and attitudes. Trials, 2008, 9, 45.	1.6	17
165	Heterozygous carriers for Wilson's disease—magnetic spectroscopy changes in the brain. Metabolic Brain Disease, 2009, 24, 463-468.	2.9	17
166	Acute focal dystonia induced by a tricyclic antidepressant in a patient with Wilson disease: a case report. Neurologia I Neurochirurgia Polska, 2013, 47, 502-506.	1.2	17
167	Gene variants encoding proteins involved in antioxidant defense system and the clinical expression of Wilson disease. Liver International, 2015, 35, 215-222.	3.9	17
168	Evolution of diagnostic criteria for multiple sclerosis. Neurologia I Neurochirurgia Polska, 2015, 49, 313-321.	1.2	17
169	Cardiac assessment in Wilson's disease patients based on electrocardiography and echocardiography examination. Archives of Medical Science, 2019, 15, 857-864.	0.9	17
170	Intravenous thrombolysis for acute ischaemic stroke in patients not fully adhering to the European licence in Poland. Neurologia I Neurochirurgia Polska, 2012, 46, 3-14.	1.2	16
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