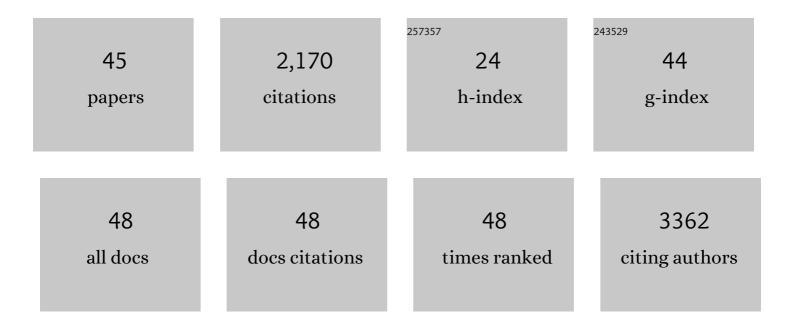
German Moris

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Frequency, symptoms, risk factors, and outcomes of autoimmune encephalitis after herpes simplex encephalitis: a prospective observational study and retrospective analysis. Lancet Neurology, The, 2018, 17, 760-772.	4.9	422
2	Autoimmune post–herpes simplex encephalitis of adults and teenagers. Neurology, 2015, 85, 1736-1743.	1.5	226
3	The Challenge of Drug-Induced Aseptic Meningitis. Archives of Internal Medicine, 1999, 159, 1185.	4.3	157
4	Profile of microRNAs in the plasma of Parkinson's disease patients and healthy controls. Journal of Neurology, 2013, 260, 1420-1422.	1.8	132
5	<i>SPG7</i> mutational screening in spastic paraplegia patients supports a dominant effect for some mutations and a pathogenic role for p.A510V. Clinical Genetics, 2013, 83, 257-262.	1.0	94
6	Clinical and therapeutic features of myasthenia gravis in adults based on age at onset. Neurology, 2020, 94, e1171-e1180.	1.5	88
7	Angiotensin converting enzyme and endothelial nitric oxide synthase DNA polymorphisms and late onset Alzheimer's disease. Journal of Neurology, Neurosurgery and Psychiatry, 1999, 67, 733-736.	0.9	83
8	Inflammatory bowel disease: An increased risk factor for neurologic complications. World Journal of Gastroenterology, 2014, 20, 1228.	1.4	80
9	LRRK2 mutations are a common cause of Parkinson's disease in Spain. European Journal of Neurology, 2006, 13, 391-394.	1.7	60
10	MiRNA Profile in the Substantia Nigra of Parkinson's Disease and Healthy Subjects. Journal of Molecular Neuroscience, 2014, 54, 830-836.	1.1	58
11	Neurologic disorders associated with inflammatory bowel disease. European Journal of Neurology, 2011, 18, 138-143.	1.7	55
12	The Challenge of Drug-Induced Aseptic Meningitis Revisited. JAMA Internal Medicine, 2014, 174, 1511.	2.6	53
13	Mutational spectrum of the SPG4 (SPAST) and SPG3A (ATL1) genes in Spanish patients with hereditary spastic paraplegia. BMC Neurology, 2010, 10, 89.	0.8	49
14	A Search for SNCA 3′ UTR Variants Identified SNP rs356165 as a Determinant of Disease Risk and Onset Age in Parkinson's Disease. Journal of Molecular Neuroscience, 2012, 47, 425-430.	1.1	49
15	Association between an α2 Macroglobulin DNA Polymorphism and Late-Onset Alzheimer's Disease. Biochemical and Biophysical Research Communications, 1999, 264, 48-50.	1.0	48
16	FGF20 rs12720208 SNP and microRNA-433 variation: No association with Parkinson's disease in Spanish patients. Neuroscience Letters, 2010, 479, 22-25.	1.0	46
17	Trunk muscle involvement in late-onset Pompe disease: Study of thirty patients. Neuromuscular Disorders, 2012, 22, S148-S154.	0.3	45
18	SUNCT Syndrome and Seborrheic Dermatitis Associated with Craneosynostosis. Cephalalgia, 2001, 21, 157-159.	1.8	42

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19	Analysis of the <i>Microâ€RNAâ€133</i> and <i>PITX3</i> genes in Parkinson's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 1234-1239.	1.1	33
20	Late-onset Alzheimer's disease is associated with mitochondrial DNA 7028C/haplogroup H and D310 poly-C tract heteroplasmy. Neurogenetics, 2011, 12, 345-346.	0.7	33
21	Chronic pain has a strong impact on quality of life in facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2018, 57, 380-387.	1.0	33
22	Alpha-synuclein transcript isoforms in three different brain regions from Parkinson's disease and healthy subjects in relation to the SNCA rs356165/rs11931074 polymorphisms. Neuroscience Letters, 2014, 562, 45-49.	1.0	30
23	The Distinctive Headache of the Occipital Condyle Syndrome: A Report of Four Cases. Headache, 1998, 38, 308-311.	1.8	28
24	Netrin-1 receptor antibodies in thymoma-associated neuromyotonia with myasthenia gravis. Neurology, 2017, 88, 1235-1242.	1.5	28
25	Acute deafness as an extraintestinal manifestation of ulcerative colitis. European Journal of Internal Medicine, 2005, 16, 440-442.	1.0	20
26	Peripheral neuropathy: An underreported neurologic manifestation of inflammatory bowel disease. European Journal of Internal Medicine, 2015, 26, 468-475.	1.0	19
27	Delayed posterior encephalopathy syndrome following chemotherapy with oxaliplatin and gemcitabine. Journal of Neurology, 2007, 254, 534-535.	1.8	16
28	Guillain-Barré syndrome, tuberculosis and inflammatory bowel disease: a multiple association. International Archive of Medicine, 2010, 3, 15.	1.2	16
29	Lack of association between protocadherin 11-X/Y (PCDH11X and PCDH11Y) polymorphisms and late onset Alzheimer's disease. Brain Research, 2011, 1383, 252-256.	1.1	16
30	The screening of the 3′UTR sequence of LRRK2 identified an association between the rs66737902 polymorphism and Parkinson's disease. Journal of Human Genetics, 2014, 59, 346-348.	1.1	14
31	Cluster-Like Headache Heralding Cerebral Venous Thrombosis. Cephalalgia, 2008, 28, 906-907.	1.8	13
32	Mutational Screening of PARKIN Identified a 3′ UTR Variant (rs62637702) Associated with Parkinson's Disease. Journal of Molecular Neuroscience, 2013, 50, 264-269.	1.1	11
33	Chronic migraine does not increase posterior circulation territory (PCT) infarct-like lesions. Journal of the Neurological Sciences, 2014, 336, 180-183.	0.3	10
34	Mitochondrial DNA polymorphisms/haplogroups in hereditary spastic paraplegia. Journal of Neurology, 2012, 259, 246-250.	1.8	9
35	Hyperintensity in the basis pontis: Atypical neuroradiological findings in a woman with FXTAS. Movement Disorders, 2010, 25, 649-650.	2.2	8
36	Diabetes mellitus mal controlada como desencadenante de un caso de enfermedad de Marchiafava-Bignami. NeurologÃa, 2016, 31, 498-500.	0.3	8

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37	Trigeminal Mononeuropathy: First Clinical Manifestation of Breast Cancer. European Neurology, 2005, 54, 212-213.	0.6	7
38	Ipsilateral reversible diaphragmatic paralysis after pons stroke. Journal of Neurology, 2012, 259, 966-968.	1.8	7
39	Sudden paraplegia after lumbar puncture as a clue in the diagnosis of a patient with spinal dural arteriovenous fistula. European Spine Journal, 2017, 26, 151-153.	1.0	7
40	Amyloid Precursor Protein Gene (APP) Variation in Late-Onset Alzheimer's Disease. Journal of Molecular Neuroscience, 2011, 45, 5-9.	1.1	4
41	Cryoglobulinaemic neuropathy: a further cause of bilateral sciatic neuropathy. International Archive of Medicine, 2008, 1, 18.	1.2	3
42	A Painless Burn Due to Lack of Painkillers. JAMA Neurology, 2014, 71, 240.	4.5	1
43	Abnormal electrocardiogram in a patient with amyotrophic lateral sclerosis mimicking myocardial ischaemia. World Journal of Clinical Cases, 2014, 2, 211.	0.3	1
44	Drug-Induced Aseptic Meningitis and Other Mimics. , 2018, , 275-300.		0
45	Hydrocephalus and myasthenia gravis: neurological manifestations in autoimmune polyendocrine syndrome type II. Acta Neurologica Belgica, 2022, , 1.	0.5	0