German Moris

List of Publications by Year
in descending order

45 papers	2,170 citations	257357 24 h-index	243529 44 g-index
48 all docs	48 docs citations	48 times ranked	3362 citing authors

#	Article	IF	Citations
1	Hydrocephalus and myasthenia gravis: neurological manifestations in autoimmune polyendocrine syndrome type II. Acta Neurologica Belgica, 2022, , 1.	0.5	0
2	Clinical and therapeutic features of myasthenia gravis in adults based on age at onset. Neurology, 2020, 94, e1171-e1180.	1.5	88
3	Drug-Induced Aseptic Meningitis and Other Mimics. , 2018, , 275-300.		0
4	Chronic pain has a strong impact on quality of life in facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2018, 57, 380-387.	1.0	33
5	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
6	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
7	Netrin-1 receptor antibodies in thymoma-associated neuromyotonia with myasthenia gravis. Neurology, 2017, 88, 1235-1242.	1.5	28
8	Diabetes mellitus mal controlada como desencadenante de un caso de enfermedad de Marchiafava-Bignami. NeurologÃa, 2016, 31, 498-500.	0.3	8
9	Peripheral neuropathy: An underreported neurologic manifestation of inflammatory bowel disease. European Journal of Internal Medicine, 2015, 26, 468-475.	1.0	19
10	Autoimmune postâ€“herpes simplex encephalitis of adults and teenagers. Neurology, 2015, 85, 1736-1743.	1.5	226
11	The screening of the 3â€²UTR sequence of LRRK2 identified an association between the rs66737902 polymorphism and Parkinsonâ€™s disease. Journal of Human Genetics, 2014, 59, 346-348.	1.1	14
12	A Painless Burn Due to Lack of Painkillers. JAMA Neurology, 2014, 71, 240.	4.5	1
13	The Challenge of Drug-Induced Aseptic Meningitis Revisited. JAMA Internal Medicine, 2014, 174, 1511.	2.6	53
14	Chronic migraine does not increase posterior circulation territory (PCT) infarct-like lesions. Journal of the Neurological Sciences, 2014, 336, 180-183.	0.3	10
15	MiRNA Profile in the Substantia Nigra of Parkinsonâ€™s Disease and Healthy Subjects. Journal of Molecular Neuroscience, 2014, 54, 830-836.	1.1	58
16	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
17	Inflammatory bowel disease: An increased risk factor for neurologic complications. World Journal of Gastroenterology, 2014, 20, 1228.	1.4	80
18	Abnormal electrocardiogram in a patient with amyotrophic lateral sclerosis mimicking myocardial ischaemia. World Journal of Clinical Cases, 2014, 2, 211.	0.3	1

#	Article	IF	Citations
19	Profile of microRNAs in the plasma of Parkinsonâ€™s disease patients and healthy controls. Journal of Neurology, 2013, 260, 1420-1422.	1.8	132
20	Mutational Screening of PARKIN Identified a 3â€² UTR Variant (rs62637702) Associated with Parkinsonâ€™s Disease. Journal of Molecular Neuroscience, 2013, 50, 264-269.	1.1	11
21	SPG7 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
22	Ipsilateral reversible diaphragmatic paralysis after pons stroke. Journal of Neurology, 2012, 259, 966-968.	1.8	7
23	Trunk muscle involvement in late-onset Pompe disease: Study of thirty patients. Neuromuscular Disorders, 2012, 22, S148-S154.	0.3	45
24	A Search for SNCA 3â€² UTR Variants Identified SNP rs356165 as a Determinant of Disease Risk and Onset Age in Parkinsonâ€™s Disease. Journal of Molecular Neuroscience, 2012, 47, 425-430.	1.1	49
25	Mitochondrial DNA polymorphisms/haplogroups in hereditary spastic paraplegia. Journal of Neurology, 2012, 259, 246-250.	1.8	9
26	Neurologic disorders associated with inflammatory bowel disease. European Journal of Neurology, 2011, 18, 138-143.	1.7	55
27	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
28	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
29	Amyloid Precursor Protein Gene (APP) Variation in Late-Onset Alzheimerâ€™s Disease. Journal of Molecular Neuroscience, 2011, 45, 5-9.	1.1	4
30	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
31	Analysis of the MicroâRNAâ€133 and PITX3 genes in Parkinson's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 1234-1239.	1.1	33
32	Hyperintensity in the basis pontis: Atypical neuroradiological findings in a woman with FXTAS. Movement Disorders, 2010, 25, 649-650.	2.2	8
33	Guillain-BarrÃ© syndrome, tuberculosis and inflammatory bowel disease: a multiple association. International Archive of Medicine, 2010, 3, 15.	1.2	16
34	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
35	Cryoglobulinaemic neuropathy: a further cause of bilateral sciatic neuropathy. International Archive of Medicine, 2008, 1, 18.	1.2	3
36	Cluster-Like Headache Heralding Cerebral Venous Thrombosis. Cephalalgia, 2008, 28, 906-907.	1.8	13

#	Article	IF	Citations
37	Delayed posterior encephalopathy syndrome following chemotherapy with oxaliplatin and gemcitabine. Journal of Neurology, 2007, 254, 534-535.	1.8	16
38	LRRK2 mutations are a common cause of Parkinson's disease in Spain. European Journal of Neurology, 2006, 13, 391-394.	1.7	60
39	Trigeminal Mononeuropathy: First Clinical Manifestation of Breast Cancer. European Neurology, 2005, 54, 212-213.	0.6	7
40	Acute deafness as an extraintestinal manifestation of ulcerative colitis. European Journal of Internal Medicine, 2005, 16, 440-442.	1.0	20
41	SUNCT Syndrome and Seborrheic Dermatitis Associated with Craneosynostosis. Cephalalgia, 2001, 21, 157-159.	1.8	42
42	The Challenge of Drug-Induced Aseptic Meningitis. Archives of Internal Medicine, 1999, 159, 1185.	4.3	157
43	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
44	Association between an Î±2 Macroglobulin DNA Polymorphism and Late-Onset Alzheimer's Disease. Biochemical and Biophysical Research Communications, 1999, 264, 48-50.	1.0	48
45	The Distinctive Headache of the Occipital Condyle Syndrome: A Report of Four Cases. Headache, 1998, 38, 308-311.	1.8	28