Doris G Leung

List of Publications by Year
in descending order

31 papers	3,560 citations	361045 20 h-index	454577 30 g-index
34 all docs	34 docs citations	34 times ranked	6449 citing authors

#	Article	IF	Citations
1	Longâ€term extension study of tofacitinib in refractory dermatomyositis. Arthritis and Rheumatology, 2022, 74, 371-372.	2.9	14
2	Study of Tofacitinib in Refractory Dermatomyositis: An Openâ€Label Pilot Study of Ten Patients. Arthritis and Rheumatology, 2021, 73, 858-865.	2.9	93
3	A phase Ib/IIa, openâ€label, multiple ascendingâ€dose trial of domagrozumab in fukutinâ€related protein limbâ€girdle muscular dystrophy. Muscle and Nerve, 2021, 64, 172-179.	1.0	5
4	Broadening learning communities during COVID-19: developing a curricular framework for telemedicine education in neurology. BMC Medical Education, 2021, 21, 549.	1.0	7
5	Longitudinal functional and imaging outcome measures in FKRP limb-girdle muscular dystrophy. BMC Neurology, 2020, 20, 196.	0.8	13
6	Advancements in magnetic resonance imagingâ€based biomarkers for muscular dystrophy. Muscle and Nerve, 2019, 60, 347-360.	1.0	17
7	Mitochondrial DNA Deletions With Low-Level Heteroplasmy in Adult-Onset Myopathy. Journal of Clinical Neuromuscular Disease, 2018, 19, 117-123.	0.3	6
8	Multivoxel proton magnetic resonance spectroscopy in facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2018, 57, 958-963.	1.0	7
9	Magnetic resonance imaging in facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2018, 57, 872-874.	1.0	5
10	Myopathy: Recent Progress, Current Therapies, and Future Directions. Neurotherapeutics, 2018, 15, 837-839.	2.1	0
11	Advanced MRI Techniques for Muscle Imaging. Seminars in Musculoskeletal Radiology, 2017, 21, 459-469.	0.4	25
12	Magnetic resonance imaging patterns of muscle involvement in genetic muscle diseases: a systematic review. Journal of Neurology, 2017, 264, 1320-1333.	1.8	56
13	Association of severity of illness and intensive care unit readmission: A systematic review. Heart and Lung: Journal of Acute and Critical Care, 2016, 45, 3-9.e2.	0.8	58
14	Wholeâ€body magnetic resonance imaging evaluation of facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2015, 52, 512-520.	1.0	63
15	A Case of Progressive Quadriceps Weakness and Elevated Creatine Kinase Level Mimicking Inclusion Body Myositis. Arthritis Care and Research, 2014, 66, 328-333.	1.5	5
16	Sildenafil does not improve cardiomyopathy in <scp>D</scp>uchenne/<scp>B</scp>ecker muscular dystrophy. Annals of Neurology, 2014, 76, 541-549.	2.8	73
17	Magnetic resonance imaging phenotyping of Becker muscular dystrophy. Muscle and Nerve, 2014, 50, 962-967.	1.0	28
18	Therapeutic advances in muscular dystrophy. Annals of Neurology, 2013, 74, 404-411.	2.8	70

#	Article	IF	Citations
19	Transcriptional profiling in facioscapulohumeral muscular dystrophy to identify candidate biomarkers. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 16234-16239.	3.3	81
20	Whole genome association analysis shows that ACE is a risk factor for Alzheimer's disease and fails to replicate most candidates from Meta-analysis. International Journal of Molecular Epidemiology and Genetics, 2010, 1, 19-30.	0.4	4
21	Genetic Control of Human Brain Transcript Expression in Alzheimer Disease. American Journal of Human Genetics, 2009, 84, 445-458.	2.6	290
22	<i>Sorl1</i> as an Alzheimerâ€™s Disease Predisposition Gene?. Neurodegenerative Diseases, 2008, 5, 60-64.	0.8	73
23	GAB2 Alleles Modify Alzheimer's Risk in APOE É›4 Carriers. Neuron, 2007, 54, 713-720.	3.8	451
24	A survey of genetic human cortical gene expression. Nature Genetics, 2007, 39, 1494-1499.	9.4	488
25	The MAPT H1c risk haplotype is associated with increased expression of tau and especially of 4 repeat containing transcripts. Neurobiology of Disease, 2007, 25, 561-570.	2.1	231
26	A High-Density Whole-Genome Association Study Reveals That APOE Is the Major Susceptibility Gene for Sporadic Late-Onset Alzheimer's Disease. Journal of Clinical Psychiatry, 2007, 68, 613-618.	1.1	484
27	Linkage disequilibrium fine mapping and haplotype association analysis of the tau gene in progressive supranuclear palsy and corticobasal degeneration. Journal of Medical Genetics, 2005, 42, 837-846.	1.5	225
28	The H1c haplotype at the MAPT locus is associated with Alzheimer's disease. Human Molecular Genetics, 2005, 14, 2399-2404.	1.4	205
29	P4-165 Microarray analysis of gene expression in the frontal cortex of patients with frontotemporal dementia. Neurobiology of Aging, 2004, 25, S522.	1.5	0
30	Sulfated Tyrosines Contribute to the Formation of the C5a Docking Site of the Human C5a Anaphylatoxin Receptor. Journal of Experimental Medicine, 2001, 193, 1059-1066.	4.2	83
31	BACE2, a beta -secretase homolog, cleaves at the beta site and within the amyloid-beta region of the amyloid-beta precursor protein. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 9712-9717.	3.3	379