

Yanchen Xie

List of Publications by Year in descending order

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papers

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citations

759055

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24
all docs

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docs citations

24
times ranked

539
citing authors

#	ARTICLE	IF	CITATIONS
1	Complement C3 polymorphism is associated with the susceptibility of myasthenia gravis in Chinese adult patients. <i>Journal of Neuroimmunology</i> , 2021, 353, 577487.	1.1	3
2	IL-4R1± Polymorphism Is Associated With Myasthenia Gravis in Chinese Han Population. <i>Frontiers in Neurology</i> , 2018, 9, 529.	1.1	6
3	The Electrophysiological Features in X-Linked Charcot-Marie-Tooth Disease With Transient Central Nervous System Deficits. <i>Frontiers in Neurology</i> , 2018, 9, 461.	1.1	7
4	Letter re: International consensus guidance for management of myasthenia gravis: Executive summary. <i>Neurology</i> , 2017, 88, 505-505.	1.5	0
5	Rs3761389 polymorphism in autoimmune regulator (AIRE) gene is associated with susceptibility of myasthenia gravis in Chinese patients. <i>Journal of Clinical Neuroscience</i> , 2017, 40, 180-184.	0.8	7
6	Correlations of TNF-1± gene promoter polymorphisms with the risk of thymoma-associated myasthenia gravis in a northern Chinese Han population. <i>Cancer Gene Therapy</i> , 2017, 24, 259-266.	2.2	4
7	TNFAIP3 gene rs7749323 polymorphism is associated with late onset myasthenia gravis. <i>Medicine (United Tj ETQq1 1 0.784314 rgB</i>	0.4	3
8	Ablation of IL-17 expression moderates experimental autoimmune myasthenia gravis disease severity. <i>Cytokine</i> , 2017, 96, 279-285.	1.4	22
9	Gene Polymorphisms for Both Auto-antigen and Immune-Modulating Proteins Are Associated with the Susceptibility of Autoimmune Myasthenia Gravis. <i>Molecular Neurobiology</i> , 2017, 54, 4771-4780.	1.9	16
10	The Role of Osteopontin and Its Gene on Glucocorticoid Response in Myasthenia Gravis. <i>Frontiers in Neurology</i> , 2017, 8, 230.	1.1	16
11	Elevated plasma interleukin-17A in a subgroup of Myasthenia Gravis patients. <i>Cytokine</i> , 2016, 78, 44-46.	1.4	40
12	Myasthenia gravis: subgroup classifications. <i>Lancet Neurology</i> , The, 2016, 15, 355-356.	4.9	8
13	Association study between IL-17A and IL-17F gene polymorphism and myasthenia gravis in Chinese patients. <i>Neurological Sciences</i> , 2016, 37, 123-130.	0.9	10
14	Precision medicine in myasthenia graves: begin from the data precision. <i>Annals of Translational Medicine</i> , 2016, 4, 106-106.	0.7	3
15	Transient, recurrent, white matter lesions in x-linked Charcot-Marie-tooth disease with novel mutation of gap junction protein beta 1 gene in China: a case report. <i>BMC Neurology</i> , 2014, 14, 156.	0.8	13
16	CTLA4 Variants and Haplotype Contribute Genetic Susceptibility to Myasthenia Gravis in Northern Chinese Population. <i>PLoS ONE</i> , 2014, 9, e101986.	1.1	20
17	The association of HLA-DQA1*0401 and DQB1*0604 with thymomatous myasthenia gravis in northern Chinese patients. <i>Journal of the Neurological Sciences</i> , 2012, 312, 57-61.	0.3	21
18	Association between HLA-DRB1 and myasthenia gravis in a northern Han Chinese population. <i>Journal of Clinical Neuroscience</i> , 2011, 18, 1524-1527.	0.8	28

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19	Association Between Alzheimer's Disease and the NOS3 gene Glu298Asp Polymorphism in Chinese. <i>Journal of Molecular Neuroscience</i> , 2008, 34, 173-176.	1.1	13
20	The association of mitochondrial aldehyde dehydrogenase gene (ALDH2) polymorphism with susceptibility to late-onset Alzheimer's disease in Chinese. <i>Journal of the Neurological Sciences</i> , 2008, 268, 172-175.	0.3	81
21	Genetic analysis of tumor necrosis factor- α (TNF- α) G-308A and Saitohin Q7R polymorphisms with Alzheimer's disease. <i>Journal of the Neurological Sciences</i> , 2008, 270, 148-151.	0.3	28
22	Trinucleotide expansions in the SCA7 gene in a large family with spinocerebellar ataxia and craniocervical dystonia. <i>Neuroscience Letters</i> , 2008, 434, 230-233.	1.0	13
23	Lack of an Association between Alzheimer's Disease and the Cystatin C (CST3) Gene G73A Polymorphism in Mainland Chinese. <i>Dementia and Geriatric Cognitive Disorders</i> , 2008, 25, 461-464.	0.7	14
24	Association analysis of NAD(P)H:quinone oxidoreductase gene 609 C/T polymorphism with Alzheimer's disease. <i>Neuroscience Letters</i> , 2006, 409, 179-181.	1.0	13