

Ailian Du

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/2734727/ailian-du-publications-by-year.pdf>

Version: 2024-04-29

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

15
papers

114
citations

7
h-index

10
g-index

15
ext. papers

184
ext. citations

4.7
avg, IF

2.31
L-index

#	Paper	IF	Citations
15	Retinitis pigmentosa and molar tooth sign caused by novel AHI1 compound heterozygote pathogenic variants. <i>BMC Medical Genomics</i> , 2021 , 14, 242	3.7	
14	The Association between Neprilysin gene polymorphisms and Alzheimer's disease in Tibetan population. <i>Brain and Behavior</i> , 2021 , 11, e02002	3.4	2
13	5WTR CGG repeat expansion in GIPC1 is associated with oculopharyngodistal myopathy. <i>Brain</i> , 2021 , 144, 601-614	11.2	17
12	The non-syndromic clinical spectrums of mtDNA 3243A>G mutation. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2021 , 26, 128-133	1.1	0
11	Anti-inflammatory effects of α -nicotinic ACh receptors are exerted through interactions with adenylyl cyclase-6. <i>British Journal of Pharmacology</i> , 2021 , 178, 2324-2338	8.6	4
10	Characteristics of acute ischemic stroke in hospitalized patients in Tibet: a retrospective comparative study. <i>BMC Neurology</i> , 2020 , 20, 380	3.1	5
9	Design of stroke imaging package study of intracranial atherosclerosis: a multicenter, prospective, cohort study. <i>Annals of Translational Medicine</i> , 2020 , 8, 13	3.2	2
8	Prevalence and Risk Factors of White Matter Lesions in Tibetan Patients Without Acute Stroke. <i>Stroke</i> , 2020 , 51, 149-153	6.7	8
7	Response to "low heteroplasmy rates in clinically affected m.3243A>G carriers not necessarily explain the phenotype". <i>Journal of the Neurological Sciences</i> , 2020 , 409, 116631	3.2	
6	Heteroplasmy and phenotype spectrum of the mitochondrial tRNA gene m.3243A>G mutation in seven Han Chinese families. <i>Journal of the Neurological Sciences</i> , 2020 , 408, 116562	3.2	7
5	MT-ND5 Mutation Exhibits Highly Variable Neurological Manifestations at Low Mutant Load. <i>EBioMedicine</i> , 2018 , 30, 86-93	8.8	33
4	Acute mitochondrial myopathy with respiratory insufficiency and motor axonal polyneuropathy. <i>International Journal of Neuroscience</i> , 2018 , 128, 231-236	2	7
3	Deep Brain Stimulation: A Potential Treatment for Dementia in Alzheimer's Disease (AD) and Parkinson's Disease Dementia (PDD). <i>Frontiers in Neuroscience</i> , 2018 , 12, 360	5.1	14
2	Suppression of CHRN endocytosis by carbonic anhydrase CAR3 in the pathogenesis of myasthenia gravis. <i>Autophagy</i> , 2017 , 13, 1981-1994	10.2	9
1	Endoplasmic reticulum stress contributes to acetylcholine receptor degradation by promoting endocytosis in skeletal muscle cells. <i>Journal of Neuroimmunology</i> , 2016 , 290, 109-14	3.5	6