## P Anthony Akkari

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/12046565/publications.pdf

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		1040056	1199594
12	728	9	12
papers	citations	h-index	g-index
12	12	12	1263
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Short structural variants as informative genetic markers for ALS disease risk and progression. BMC Medicine, 2022, 20, 11.	<b>5.</b> 5	4
2	Novel STMN2 Variant Linked to Amyotrophic Lateral Sclerosis Risk and Clinical Phenotype. Frontiers in Aging Neuroscience, 2021, 13, 658226.	3.4	38
3	TOMM40 â€~523' poly-T repeat length is a determinant of longitudinal cognitive decline in Parkinson's disease. Npj Parkinson's Disease, 2021, 7, 56.	5.3	2
4	Disease-modifying effects of an <i>SCAF4</i> structural variant in a predominantly <i>SOD1</i> ALS cohort. Neurology: Genetics, 2020, 6, e470.	1.9	9
5	Single Nucleotide Polymorphisms Associated With Gut Homeostasis Influence Risk and Age-at-Onset of Parkinson's Disease. Frontiers in Aging Neuroscience, 2020, 12, 603849.	3.4	16
6	Association of a structural variant within the <i>SQSTM1</i> gene with amyotrophic lateral sclerosis. Neurology: Genetics, 2020, 6, e406.	1.9	9
7	ALS Genetics, Mechanisms, and Therapeutics: Where Are We Now?. Frontiers in Neuroscience, 2019, 13, 1310.	2.8	487
8	Structural variants can be more informative for disease diagnostics, prognostics and translation than current SNP mapping and exon sequencing. Expert Opinion on Drug Metabolism and Toxicology, 2016, 12, 135-147.	3.3	23
9	Africanâ€American TOMM40'523â€ <i>APOE</i> haplotypes are admixture of West African and Caucasian alleles. Alzheimer's and Dementia, 2014, 10, 592.	0.8	32
10	An ?tropomyosin mutation alters dimer preference in nemaline myopathy. Annals of Neurology, 2005, 57, 42-49.	5.3	62
11	Production of human skeletal $\hat{l}$ ±-actin proteins by the baculovirus expression system. Biochemical and Biophysical Research Communications, 2003, 307, 74-79.	2.1	16
12	Expression and biological activity of Baculovirus generated wild-type human slow α tropomyosin and the Met9Arg mutant responsible for a dominant form of nemaline myopathy. Biochemical and Biophysical Research Communications, 2002, 296, 300-304.	2.1	30