

Ahmad Al-Khleifat

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

1,849
citations

516681

16
h-index

477281

29
g-index

36
all docs

36
docs citations

36
times ranked

3465
citing authors

#	ARTICLE	IF	CITATIONS
1	Detection of long repeat expansions from PCR-free whole-genome sequence data. <i>Genome Research</i> , 2017, 27, 1895-1903.	5.5	277
2	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. <i>Nature Genetics</i> , 2021, 53, 1636-1648.	21.4	223
3	A multicentre validation study of the diagnostic value of plasma neurofilament light. <i>Nature Communications</i> , 2021, 12, 3400.	12.8	219
4	Genomic and phenotypic insights from an atlas of genetic effects on DNA methylation. <i>Nature Genetics</i> , 2021, 53, 1311-1321.	21.4	218
5	Stage at which riluzole treatment prolongs survival in patients with amyotrophic lateral sclerosis: a retrospective analysis of data from a dose-ranging study. <i>Lancet Neurology</i> , The, 2018, 17, 416-422.	10.2	175
6	Exome sequencing in amyotrophic lateral sclerosis implicates a novel gene, DNAJC7, encoding a heat-shock protein. <i>Nature Neuroscience</i> , 2019, 22, 1966-1974.	14.8	101
7	What causes amyotrophic lateral sclerosis?. <i>F1000Research</i> , 2017, 6, 371.	1.6	94
8	C9orf72 intermediate expansions of 24-30 repeats are associated with ALS. <i>Acta Neuropathologica Communications</i> , 2019, 7, 115.	5.2	75
9	A comprehensive analysis of rare genetic variation in amyotrophic lateral sclerosis in the UK. <i>Brain</i> , 2017, 140, 1611-1618.	7.6	71
10	Comparison of the King's and MiToS staging systems for ALS. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2017, 18, 227-232.	1.7	58
11	Association of Variants in the <i>SPTLC1</i> Gene With Juvenile Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2021, 78, 1236.	9.0	46
12	Genome-wide Meta-analysis Finds the ACSL5-ZDHHC6 Locus Is Associated with ALS and Links Weight Loss to the Disease Genetics. <i>Cell Reports</i> , 2020, 33, 108323.	6.4	41
13	Younger age of onset in familial amyotrophic lateral sclerosis is a result of pathogenic gene variants, rather than ascertainment bias. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 268-271.	1.9	38
14	Genome-wide study of DNA methylation shows alterations in metabolic, inflammatory, and cholesterol pathways in ALS. <i>Science Translational Medicine</i> , 2022, 14, eabj0264.	12.4	38
15	A standard operating procedure for King's ALS clinical staging. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019, 20, 159-164.	1.7	26
16	Structural variation analysis of 6,500 whole genome sequences in amyotrophic lateral sclerosis. <i>Npj Genomic Medicine</i> , 2022, 7, 8.	3.8	23
17	Telomere length is greater in ALS than in controls: a whole genome sequencing study. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019, 20, 229-234.	1.7	18
18	Whole-genome sequencing reveals that variants in the Interleukin 18 Receptor Accessory Protein 3'UTR protect against ALS. <i>Nature Neuroscience</i> , 2022, 25, 433-445.	14.8	16

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19	<i>SCFD1</i> expression quantitative trait loci in amyotrophic lateral sclerosis are differentially expressed. Brain Communications, 2021, 3, fcab236.	3.3	14
20	Cross-reactive probes on Illumina DNA methylation arrays: a large study on ALS shows that a cautionary approach is warranted in interpreting epigenome-wide association studies. NAR Genomics and Bioinformatics, 2020, 2, lqaa105.	3.2	13
21	ALSgeneScanner: a pipeline for the analysis and interpretation of DNA sequencing data of ALS patients. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 207-215.	1.7	11
22	Relationship between smoking and ALS: Mendelian randomisation interrogation of causality. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1312-1315.	1.9	11
23	UK case control study of smoking and risk of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 222-227.	1.7	10
24	Does genetic anticipation occur in familial Alexander disease?. Neurogenetics, 2021, 22, 215-219.	1.4	4
25	An international virtual hackathon to build tools for the analysis of structural variants within species ranging from coronaviruses to vertebrates. F1000Research, 2021, 10, 246.	1.6	3
26	Stage of prolonged survival in ALS – Author's reply. Lancet Neurology, The, 2018, 17, 579-580.	10.2	2
27	Intuitive Staging Correlates With King's Clinical Stage. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 336-340.	1.7	2
28	An international virtual hackathon to build tools for the analysis of structural variants within species ranging from coronaviruses to vertebrates. F1000Research, 2021, 10, 246.	1.6	2
29	The third international hackathon for applying insights into large-scale genomic composition to use cases in a wide range of organisms. F1000Research, 0, 11, 530.	1.6	1