Hannu Laaksovirta

List of Publications by Year
in descending order

13	6,010	840776	1125743
papers	citations	11 h-index	13 g-index

14	14	14	7559
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	ALS in Finland. Neurology: Genetics, 2022, 8, e665.	1.9	11
2	Association of Variants in the SPTLC1 Gene With Juvenile Amyotrophic Lateral Sclerosis. JAMA Neurology, 2021, 78, 1236.	9.0	46
3	Carriership of two copies of C9orf72 hexanucleotide repeat intermediate-length alleles is a risk factor for ALS in the Finnish population. Acta Neuropathologica Communications, 2020, 8, 187.	5.2	16
4	Home invasive mechanical ventilation in Finland in 2015â€“2019. ERJ Open Research, 2020, 6, 00223-2020.	2.6	4
5	Oligogenic basis of sporadic ALS. Neurology: Genetics, 2019, 5, e335.	1.9	15
6	C9orf72 hexanucleotide repeat length in older population: normal variation and effects on cognition. Neurobiology of Aging, 2019, 84, 242.e7-242.e12.	3.1	16
7	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. Neuron, 2018, 97, 1268-1283.e6.	8.1	517
8	July 2017 ENCALS statement on edaravone. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 471-474.	1.7	41
9	FGF21 is a biomarker for mitochondrial translation and mtDNA maintenance disorders. Neurology, 2016, 87, 2290-2299.	1.1	167
10	Genome-Wide Analysis of the Heritability of Amyotrophic Lateral Sclerosis. JAMA Neurology, 2014, 71, 1123.	9.0	69
11	Frequency of the C9orf72 hexanucleotide repeat expansion in patients with amyotrophic lateral sclerosis and frontotemporal dementia: a cross-sectional study. Lancet Neurology, The, 2012, 11, 323-330.	10.2	1,039
12	A Hexanucleotide Repeat Expansion in C9ORF72 Is the Cause of Chromosome 9p21-Linked ALS-FTD. Neuron, 2011, 72, 257-268.	8.1	3,833
13	Chromosome 9p21 in amyotrophic lateral sclerosis in Finland: a genome-wide association study. Lancet Neurology, The, 2010, 9, 978-985.	10.2	236