

Hiroyuki Akagawa

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

159
citations

1478505

6
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

263
citing authors

#	ARTICLE	IF	CITATIONS
1	A haplotype spanning two genes, ELN and LIMK1 , decreases their transcripts and confers susceptibility to intracranial aneurysms. <i>Human Molecular Genetics</i> , 2006, 15, 1722-1734.	2.9	62
2	Systematic screening of lysyl oxidase-like (LOXL) family genes demonstrates that LOXL2 is a susceptibility gene to intracranial aneurysms. <i>Human Genetics</i> , 2007, 121, 377-387.	3.8	36
3	Unusual case of cerebral small vessel disease with a heterozygous nonsense mutation in HTRA1. <i>Journal of the Neurological Sciences</i> , 2016, 362, 144-146.	0.6	16
4	Exome Sequencing Identified CCER2 as a Novel Candidate Gene for Moyamoya Disease. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2017, 26, 150-161.	1.6	13
5	Is there any evidence for linkage on chromosome 17cen in affected Japanese sib-pairs with an intracranial aneurysm?. <i>Journal of Human Genetics</i> , 2006, 51, 491-494.	2.3	10
6	Novel RASA1 mutations in Japanese pedigrees with capillary malformation-arteriovenous malformation. <i>Brain and Development</i> , 2019, 41, 812-816.	1.1	6
7	Rare and Low-Frequency Variants in RNF213 Confer Susceptibility to Moyamoya Syndrome Associated with Hyperthyroidism. <i>World Neurosurgery</i> , 2019, 127, e460-e466.	1.3	5
8	Association of Rare Nonsynonymous Variants in PKD1 and PKD2 with Familial Intracranial Aneurysms in a Japanese Population. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2016, 25, 2900-2906.	1.6	4
9	Deep intronic deletion in intron 3 of PLP1 is associated with a severe phenotype of Pelizaeus-Merzbacher disease. <i>Human Genome Variation</i> , 2021, 8, 14.	0.7	4
10	Novel SLC20A2 mutation in a Japanese pedigree with primary familial brain calcification. <i>Journal of the Neurological Sciences</i> , 2019, 399, 183-185.	0.6	2
11	Whole-exome sequencing in a Japanese multiplex family identifies new susceptibility genes for intracranial aneurysms. <i>PLoS ONE</i> , 2022, 17, e0265359.	2.5	1