

Hiroyuki Akagawa

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

159
citations

1478505

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h-index

1281871

11
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11
all docs

11
docs citations

11
times ranked

263
citing authors

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