Farid Radmanesh

List of Publications by Year
in descending order

35 papers	1,764 citations	377584 21 h-index	406436 35 g-index
37 all docs	37 docs citations	37 times ranked	5141 citing authors

#	Article	IF	Citations
1	Large-scale analyses of common and rare variants identify 12 new loci associated with atrial fibrillation. Nature Genetics, 2017, 49, 946-952.	9.4	279
2	Meta-analysis of Genome-wide Association Studies Identifies 1q22 as a Susceptibility Locus for Intracerebral Hemorrhage. American Journal of Human Genetics, 2014, 94, 511-521.	2.6	235
3	Identification of additional risk loci for stroke and small vessel disease: a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2016, 15, 695-707.	4.9	130
4	Common variation in COL4A1/COL4A2 is associated with sporadic cerebral small vessel disease. Neurology, 2015, 84, 918-926.	1.5	106
5	GENOME-WIDE ASSOCIATION STUDY (GWAS) AND GENOME-WIDE BY ENVIRONMENT INTERACTION STUDY (GWEIS) OF DEPRESSIVE SYMPTOMS IN AFRICAN AMERICAN AND HISPANIC/LATINA WOMEN. Depression and Anxiety, 2016, 33, 265-280.	2.0	99
6	Dermal sinus tract of the spine. Child's Nervous System, 2010, 26, 349-357.	0.6	97
7	Mutations in LAMB1 Cause Cobblestone Brain Malformation without Muscular or Ocular Abnormalities. American Journal of Human Genetics, 2013, 92, 468-474.	2.6	96
8	Genome-wide meta-analysis of cerebral white matter hyperintensities in patients with stroke. Neurology, 2016, 86, 146-153.	1.5	91
9	Genetic variation at 16q24.2 is associated with small vessel stroke. Annals of Neurology, 2017, 81, 383-394.	2.8	73
10	COL4A2 is associated with lacunar ischemic stroke and deep ICH. Neurology, 2017, 89, 1829-1839.	1.5	58
11	Dual-source computed tomography angiography for diagnosis and assessment of coronary artery disease: Systematic review and meta-analysis. Journal of Cardiovascular Computed Tomography, 2012, 6, 78-90.	0.7	54
12	Exome-chip meta-analysis identifies novel loci associated with cardiac conduction, including ADAMTS6. Genome Biology, 2018, 19, 87.	3.8	47
13	Shunt complications in children with myelomeningocele: effect of timing of shunt placement. Journal of Neurosurgery: Pediatrics, 2009, 3, 516-520.	0.8	42
14	Accuracy of imputation to infer unobserved APOE epsilon alleles in genome-wide genotyping data. European Journal of Human Genetics, 2014, 22, 1239-1242.	1.4	36
15	Genetic variants inCETPincrease risk of intracerebral hemorrhage. Annals of Neurology, 2016, 80, 730-740.	2.8	33
16	APOE Îµ variants increase risk of warfarin-related intracerebral hemorrhage. Neurology, 2014, 83, 1139-1146.	1.5	29
17	Discovery of novel heart rate-associated loci using the Exome Chip. Human Molecular Genetics, 2017, 26, 2346-2363.	1.4	29
18	ExomeChip-Wide Analysis of 95 626 Individuals Identifies 10 Novel Loci Associated With QT and JT Intervals. Circulation Genomic and Precision Medicine, 2018, 11, e001758.	1.6	27

#	Article	IF	Citations
19	Risk Factors for Computed Tomography Angiography Spot Sign in Deep and Lobar Intracerebral Hemorrhage Are Shared. Stroke, 2014, 45, 1833-1835.	1.0	26
20	17p12 Influences Hematoma Volume and Outcome in Spontaneous Intracerebral Hemorrhage. Stroke, 2018, 49, 1618-1625.	1.0	26
21	Genetic Architecture of White Matter Hyperintensities Differs in Hypertensive and Nonhypertensive Ischemic Stroke. Stroke, 2015, 46, 348-353.	1.0	25
22	Severe cerebral involvement in adult-onset hemophagocytic lymphohistiocytosis. Journal of Clinical Neuroscience, 2020, 76, 236-237.	0.8	24
23	Common and Rare Coding Genetic Variation Underlying the Electrocardiographic PR Interval. Circulation Genomic and Precision Medicine, 2018, 11, e002037.	1.6	19
24	A genome-wide association study of outcome from traumatic brain injury. EBioMedicine, 2022, 77, 103933.	2.7	17
25	Spina bifida occulta: is it a predictor of underlying spinal cord abnormality in patients with lower urinary tract dysfunction?. Journal of Neurosurgery: Pediatrics, 2008, 1, 114-117.	0.8	14
26	Cerebral Infarction as the First Presentation of Tuberculosis in an Infant: A Case Report. Journal of Microbiology, Immunology and Infection, 2010, 43, 249-252.	1.5	8
27	Rare Coding Variation and Risk of Intracerebral Hemorrhage. Stroke, 2015, 46, 2299-2301.	1.0	8
28	Teratoma within an encephalocele: common etiology or coincidence. Journal of Neurosurgery: Pediatrics, 2007, 107, 263-265.	0.8	6
29	Translational Genomics in Neurocritical Care: a Review. Neurotherapeutics, 2020, 17, 563-580.	2.1	6
30	Megalencephaly-Polymicrogyria-Polydactyly-Hydrocephalus Syndrome. Journal of Child Neurology, 2013, 28, 651-657.	0.7	5
31	Diagnostic Performance of Low-Radiation-Dose Coronary Computed Tomography Angiography. Annals of Internal Medicine, 2011, 155, 278.	2.0	4
32	Congenital spinal tumor in a patient with encephalocele and hydrocephalus: a case report. Journal of Medical Case Reports, 2011, 5, 9.	0.4	4
33	Diagnostic Accuracy and Clinical Utility of Noninvasive Testing for Coronary Artery Disease. Annals of Internal Medicine, 2011, 154, 290.	2.0	3
34	Infection in myelomeningocele after VP shunt placement. Child's Nervous System, 2011, 27, 341-342.	0.6	3
35	Spina bifida occulta. Journal of Neurosurgery: Pediatrics, 2008, 1, 113.	0.8	0