Ludmila Pawlikowska

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Assessing the association of common genetic variants in <i>EPHB4</i> and <i>RASA1</i> with phenotype severity in familial cerebral cavernous malformation. Molecular Genetics & Genomic Medicine, 2021, 9, e1794.	1.2	2
2	Association of common candidate variants with vascular malformations and intracranial hemorrhage in hereditary hemorrhagic telangiectasia. Molecular Genetics & Genomic Medicine, 2018, 6, 350-356.	1.2	19
3	Genome-wide association study of sporadic brain arteriovenous malformations. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 916-923.	1.9	29
4	Cytochrome P450 and matrix metalloproteinase genetic modifiers of disease severity in Cerebral Cavernous Malformation type 1. Free Radical Biology and Medicine, 2016, 92, 100-109.	2.9	47
5	The <i>ACVRL1</i> c.314—35A>G polymorphism is associated with organ vascular malformations in hereditary hemorrhagic telangiectasia patients with <i>ENG</i> mutations, but not in patients with <i>ACVRL1</i> mutations. American Journal of Medical Genetics, Part A, 2015, 167, 1262-1267.	1.2	19
6	Mitochondrial DNA Heteroplasmy Associations With Neurosensory and Mobility Function in Elderly Adults. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2015, 70, 1418-1424.	3.6	19
7	Expression Quantitative Trait Loci and Receptor Pharmacology Implicate Arg1 and the GABA-A Receptor as Therapeutic Targets in Neuroblastoma. Cell Reports, 2014, 9, 1034-1046.	6.4	28
8	Association of Cardiovascular Risk Factors with Disease Severity in Cerebral Cavernous Malformation Type 1 Subjects with the Common Hispanic Mutation. Cerebrovascular Diseases, 2014, 37, 57-63.	1.7	38
9	Polymorphisms in Inflammatory and Immune Response Genes Associated with Cerebral Cavernous Malformation Type 1 Severity. Cerebrovascular Diseases, 2014, 38, 433-440.	1.7	57
10	Common variants on 9p21.3 are associated with brain arteriovenous malformations with accompanying arterial aneurysms. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1280-1283.	1.9	26
11	Angiopoietin-Like 4 <i>(ANGPTL4)</i> Gene Polymorphisms and Risk of Brain Arteriovenous Malformations. Cerebrovascular Diseases, 2011, 31, 338-345.	1.7	41
12	Reduced Expression of Integrin αvβ8 Is Associated with Brain Arteriovenous Malformation Pathogenesis. American Journal of Pathology, 2010, 176, 1018-1027.	3.8	56
13	Common Variants in Interleukin-1-Beta Gene Are Associated with Intracranial Hemorrhage and Susceptibility to Brain Arteriovenous Malformation. Cerebrovascular Diseases, 2009, 27, 176-182.	1.7	84
14	<i>EPHB4</i> Gene Polymorphisms and Risk of Intracranial Hemorrhage in Patients With Brain Arteriovenous Malformations. Circulation: Cardiovascular Genetics, 2009, 2, 476-482.	5.1	33
15	Population Stratification in a Case-Control Study of Brain Arteriovenous Malformation in Latinos. Neuroepidemiology, 2008, 31, 224-228.	2.3	17
16	ASSOCIATION OF TUMOR NECROSIS FACTOR-α–238G>A AND APOLIPOPROTEIN E2 POLYMORPHISMS WIT INTRACRANIAL HEMORRHAGE AFTER BRAIN ARTERIOVENOUS MALFORMATION TREATMENT. Neurosurgery, 2007, 61, 731-740.	H 1.1	39
17	Apolipoprotein EÉ>2 Is Associated with New Hemorrhage Risk in Brain Arteriovenous Malformations. Neurosurgery, 2006, 58, 838-843.	1.1	53
18	Tumor Necrosis Factor-α–238G>A Promoter Polymorphism Is Associated With Increased Risk of New Hemorrhage in the Natural Course of Patients With Brain Arteriovenous Malformations. Stroke, 2006, 37, 231-234.	2.0	93

#	Article	IF	CITATIONS
19	Polymorphisms in Transforming Growth Factor-β-Related Genes <i>ALK1</i> and <i>ENG</i> Are Associated With Sporadic Brain Arteriovenous Malformations. Stroke, 2005, 36, 2278-2280.	2.0	90
20	Polymorphisms in Genes Involved in Inflammatory and Angiogenic Pathways and the Risk of Hemorrhagic Presentation of Brain Arteriovenous Malformations. Stroke, 2004, 35, 2294-2300.	2.0	134