

Ludmila Pawlikowska

List of Publications by Year in descending order

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papers

924
citations

471509

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#	ARTICLE	IF	CITATIONS
1	Polymorphisms in Genes Involved in Inflammatory and Angiogenic Pathways and the Risk of Hemorrhagic Presentation of Brain Arteriovenous Malformations. <i>Stroke</i> , 2004, 35, 2294-2300.	2.0	134
2	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. <i>Stroke</i> , 2006, 37, 231-234.	2.0	93
3	Polymorphisms in Transforming Growth Factor- β -Related Genes <i>ALK1</i> and <i>ENG</i> Are Associated With Sporadic Brain Arteriovenous Malformations. <i>Stroke</i> , 2005, 36, 2278-2280.	2.0	90
4	Common Variants in Interleukin-1-Beta Gene Are Associated with Intracranial Hemorrhage and Susceptibility to Brain Arteriovenous Malformation. <i>Cerebrovascular Diseases</i> , 2009, 27, 176-182.	1.7	84
5	Polymorphisms in Inflammatory and Immune Response Genes Associated with Cerebral Cavernous Malformation Type 1 Severity. <i>Cerebrovascular Diseases</i> , 2014, 38, 433-440.	1.7	57
6	Reduced Expression of Integrin α 28 Is Associated with Brain Arteriovenous Malformation Pathogenesis. <i>American Journal of Pathology</i> , 2010, 176, 1018-1027.	3.8	56
7	Apolipoprotein E ϵ 2 Is Associated with New Hemorrhage Risk in Brain Arteriovenous Malformations. <i>Neurosurgery</i> , 2006, 58, 838-843.	1.1	53
8	Cytochrome P450 and matrix metalloproteinase genetic modifiers of disease severity in Cerebral Cavernous Malformation type 1. <i>Free Radical Biology and Medicine</i> , 2016, 92, 100-109.	2.9	47
9	Angiopoietin-Like 4 & (ANGPTL4) Gene Polymorphisms and Risk of Brain Arteriovenous Malformations. <i>Cerebrovascular Diseases</i> , 2011, 31, 338-345.	1.7	41
10	ASSOCIATION OF TUMOR NECROSIS FACTOR- α 238G>A AND APOLIPOPROTEIN E2 POLYMORPHISMS WITH INTRACRANIAL HEMORRHAGE AFTER BRAIN ARTERIOVENOUS MALFORMATION TREATMENT. <i>Neurosurgery</i> , 2007, 61, 731-740.	1.1	39
11	Association of Cardiovascular Risk Factors with Disease Severity in Cerebral Cavernous Malformation Type 1 Subjects with the Common Hispanic Mutation. <i>Cerebrovascular Diseases</i> , 2014, 37, 57-63.	1.7	38
12	<i>EPHB4</i> Gene Polymorphisms and Risk of Intracranial Hemorrhage in Patients With Brain Arteriovenous Malformations. <i>Circulation: Cardiovascular Genetics</i> , 2009, 2, 476-482.	5.1	33
13	Genome-wide association study of sporadic brain arteriovenous malformations. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 916-923.	1.9	29
14	Expression Quantitative Trait Loci and Receptor Pharmacology Implicate Arg1 and the GABA-A Receptor as Therapeutic Targets in Neuroblastoma. <i>Cell Reports</i> , 2014, 9, 1034-1046.	6.4	28
15	Common variants on 9p21.3 are associated with brain arteriovenous malformations with accompanying arterial aneurysms. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, 1280-1283.	1.9	26
16	The <i>ACVRL1</i> c.314A>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. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 1262-1267.	1.2	19
17	Mitochondrial DNA Heteroplasmy Associations With Neurosensory and Mobility Function in Elderly Adults. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2015, 70, 1418-1424.	3.6	19
18	Association of common candidate variants with vascular malformations and intracranial hemorrhage in hereditary hemorrhagic telangiectasia. <i>Molecular Genetics & Genomic Medicine</i> , 2018, 6, 350-356.	1.2	19

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19	Population Stratification in a Case-Control Study of Brain Arteriovenous Malformation in Latinos. <i>Neuroepidemiology</i> , 2008, 31, 224-228.	2.3	17
20	Assessing the association of common genetic variants in <i>EPHB4</i> and <i>RASA1</i> with phenotype severity in familial cerebral cavernous malformation. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1794.	1.2	2