

# Dorota Dziewulska

## List of Publications by Year in descending order

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Version: 2024-02-01

38  
papers

362  
citations

1039880

9  
h-index

887953

17  
g-index

40  
all docs

40  
docs citations

40  
times ranked

592  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pericytes as a new target for pathological processes in CADASIL. <i>Neuropathology</i> , 2012, 32, 515-521.	0.7	72
2	ELISA reveals a difference in the structure of substantia nigra ferritin in Parkinson's disease and incidental Lewy body compared to control. <i>Parkinsonism and Related Disorders</i> , 2007, 13, 214-218.	1.1	51
3	Dynactin Deficiency in the CNS of Humans with Sporadic ALS and Mice with Genetically Determined Motor Neuron Degeneration. <i>Neurochemical Research</i> , 2013, 38, 2463-2473.	1.6	31
4	A Study of Glutathione S-transferase pi Expression in Central Nervous System of Subjects with Amyotrophic Lateral Sclerosis Using RNA Extraction from Formalin-Fixed, Paraffin-Embedded Material. <i>Neurochemical Research</i> , 2005, 30, 1003-1007.	1.6	19
5	CADASIL or CADVaSIL?. <i>Neuropathology</i> , 2004, 24, 16-20.	0.7	13
6	CADASIL: new cases and new questions. <i>Acta Neuropathologica</i> , 2003, 106, 569-574.	3.9	11
7	Role of endoglin and transforming growth factor-beta in progressive white matter damage after an ischemic stroke. <i>Neuropathology</i> , 2006, 26, 298-306.	0.7	11
8	Kinesin Expression in the Central Nervous System of Humans and Transgenic hSOD1G93AMice with Amyotrophic Lateral Sclerosis. <i>Neurodegenerative Diseases</i> , 2013, 12, 71-80.	0.8	11
9	CADASIL: what component of the vessel wall is really a target for Notch 3 gene mutations?. <i>Neurological Research</i> , 2004, 26, 558-562.	0.6	10
10	Is the increased expression of ubiquitin in CADASIL syndrome a manifestation of aberrant endocytosis in the vascular smooth muscle cells?. <i>Journal of Clinical Neuroscience</i> , 2008, 15, 535-540.	0.8	9
11	Olfactory impairment and pathology in neurodegenerative disorders with brain iron accumulation. <i>Acta Neuropathologica</i> , 2013, 126, 151-153.	3.9	9
12	What factors determine phenotype of cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy (CADASIL)? Considerations in the context of a novel pathogenic R110C mutation in the NOTCH3 gene. <i>Folia Neuropathologica</i> , 2017, 55, 295-300.	0.5	8
13	The prevalence and types of oral- and pharyngeal-stage dysphagia in patients with demyelinating diseases based on subjective assessment by the study subjects. <i>Multiple Sclerosis and Related Disorders</i> , 2020, 37, 101484.	0.9	8
14	Enlargement of the Nissl substance as a manifestation of early damage to spinal cord motoneurons in amyotrophic lateral sclerosis. , 2013, 32, 480-485.		8
15	Progression of morphological changes within CNS in a transgenic rat model of familial amyotrophic lateral sclerosis. <i>Folia Neuropathologica</i> , 2006, 44, 162-74.	0.5	8
16	Mysteries of CADASIL - the contribution of neuropathology to understanding of the disease. <i>Folia Neuropathologica</i> , 2009, 47, 1-10.	0.5	8
17	Cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy (CADASIL) – literature review apropos an autopsy case. <i>Polish Journal of Pathology</i> , 2015, 3, 323-329.	0.1	7
18	Disturbed integrin expression in the vascular media in CADASIL. <i>Folia Neuropathologica</i> , 2016, 4, 375-381.	0.5	6

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19	Neuropathological Findings in Ephedrone Encephalopathy. <i>Movement Disorders</i> , 2020, 35, 1858-1863.	2.2	6
20	Is the spinal cord motoneuron exclusively a target in ALS? Comparison between astroglial reactivity in a rat model of familial ALS and in human sporadic ALS cases. <i>Neurological Research</i> , 2010, 32, 867-872.	0.6	5
21	Intraosseous Lipoma of the Sphenoid: A Case Study. <i>Case Reports in Neurological Medicine</i> , 2013, 2013, 1-3.	0.3	5
22	Nuclear abnormalities in vascular myocytes in cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy (CADASIL). <i>Neuropathology</i> , 2018, 38, 601-608.	0.7	5
23	Remote morphological changes in the white matter after ischaemic stroke. <i>Folia Neuropathologica</i> , 2004, 42, 75-80.	0.5	5
24	Original article Non-woven nanofiber mats – a new perspective for experimental studies of the central nervous system?. <i>Folia Neuropathologica</i> , 2014, 4, 407-416.	0.5	4
25	Nanofiber mat spinal cord dressing-released glutamate impairs blood-spinal cord barrier. <i>Folia Neuropathologica</i> , 2016, 4, 392-404.	0.5	4
26	Cerebral amyloid angiopathy-related inflammation – A case report presenting diagnostic difficulties. <i>Neurologia i Neurochirurgia Polska</i> , 2018, 52, 298-305.	0.6	4
27	Clinical presentation of Y189C mutation of the NOTCH3 gene in the Polish family with CADASIL. <i>Folia Neuropathologica</i> , 2020, 58, 83-92.	0.5	4
28	Diverse Expression of Selected SMN Complex Proteins in Humans with Sporadic Amyotrophic Lateral Sclerosis and in a Transgenic Rat Model of Familial Form of the Disease. <i>PLoS ONE</i> , 2014, 9, e104614.	1.1	3
29	Changes in the Vascular Extracellular Matrix as a Potential Cause of Myocyte Loss via Anoikis in Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy. , 2017, 07, .		3
30	Dementia means number of things - the overlap of neurodegeneration with brain iron accumulation (NBIA) and Alzheimer changes: an autopsy case. , 2010, 48, 129-33.		3
31	NF- $\kappa$ B deficit in spinal motoneurons in patients with sporadic amyotrophic lateral sclerosis – a pilot study. <i>Folia Neuropathologica</i> , 2015, 4, 367-376.	0.5	2
32	Identification of the first in Poland CACNA1A gene mutation in familial hemiplegic migraine. Case report. <i>Neurologia i Neurochirurgia Polska</i> , 2017, 51, 184-189.	0.6	2
33	Proteinaceous intracellular inclusions in neurodegenerative disorders. <i>Folia Neuropathologica</i> , 2005, 43, 51-63.	0.5	2
34	Original article Immunoexpression of gemins 2 and 4 in the rat spinal cord. Is the SMN complex a new target in investigations of sporadic amyotrophic lateral sclerosis pathogenesis?. <i>Folia Neuropathologica</i> , 2012, 4, 390-396.	0.5	1
35	Diplomyelia in a patient with a clinical suspicion of autosomal recessive spastic ataxia of Charlevoix-Saguenay type (ARSACS). <i>Folia Neuropathologica</i> , 2020, 58, 94-99.	0.5	1
36	The spectrum of microvascular ultrastructural changes in the subpopulation of patients with migraine and cerebral white matter hyperintensities on MRI. <i>Folia Neuropathologica</i> , 2021, 59, 262-270.	0.5	1

#	ARTICLE	IF	CITATIONS
37	Survival motor neuron: motor neuron insurance for a whole lifespan?. , 2011, 49, 301-10.		1
38	Paraneoplastic brainstem encephalomyelitis and atypical form of chronic inflammatory demyelinating polyneuropathy in patient with testicular germinal tumorâ€”Is this an overlap syndrome? A case report. Neurologia I Neurochirurgia Polska, 2015, 49, 129-133.	0.6	0