Dusanka L Savic-Pavicevic

List of Publications by Year
in descending order

71	1,040	430874 18	526287 27
papers	citations	h-index	g-index

72	72	72	1730
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Loss-of-function mutations in HINT1 cause axonal neuropathy with neuromyotonia. Nature Genetics, 2012, 44, 1080-1083.	21.4	102
2	Molecular Genetics and Genetic Testing in Myotonic Dystrophy Type 1. BioMed Research International, 2013, 2013, 1-13.	1.9	42
3	Assessment of association between genetic variants in microRNA genes hsa-miR-499, hsa-miR-196a2 and hsa-miR-27a and prostate cancer risk in Serbian population. Experimental and Molecular Pathology, 2015, 99, 145-150.	2.1	41
4	Novel GNB1 mutations disrupt assembly and function of G protein heterotrimers and cause global developmental delay in humans. Human Molecular Genetics, 2017, 26, ddx018.	2.9	41
5	Frontostriatal dysexecutive syndrome: a core cognitive feature of myotonic dystrophy type 2. Journal of Neurology, 2015, 262, 142-148.	3.6	38
6	Clusters of cognitive impairment among different phenotypes of myotonic dystrophy type 1 and type 2. Neurological Sciences, 2017, 38, 415-423.	1.9	37
7	Repeat Interruptions Modify Age at Onset in Myotonic Dystrophy Type 1 by Stabilizing DMPK Expansions in Somatic Cells. Frontiers in Genetics, 2018, 9, 601.	2.3	35
8	Joint effect ofADARB1gene,HTR2Cgene and stressful life events on suicide attempt risk in patients with major psychiatric disorders. World Journal of Biological Psychiatry, 2015, 16, 261-271.	2.6	33
9	250 CTG repeats in DMPK is a threshold for correlation of expansion size and age at onset of juvenile-adult DM1. Human Mutation, 2002, 19, 131-139.	2.5	30
10	Intellectual Ability in the Duchenne Muscular Dystrophy and Dystrophin Gene Mutation Location. Balkan Journal of Medical Genetics, 2014, 17, 25-35.	0.5	30
11	Molecular genetic and clinical characterization of myotonic dystrophy type 1 patients carrying variant repeats within DMPK expansions. Neurogenetics, 2017, 18, 207-218.	1.4	27
12	Intergenerational changes of CTG repeat depending on the sex of the transmitting parent in myotonic dystrophy type 1. European Journal of Neurology, 2005, 12, 236-237.	3.3	24
13	Survival and mortality of myotonic dystrophy type 1 (Steinert's disease) in the population of Belgrade. European Journal of Neurology, 2006, 13, 451-454.	3.3	24
14	Quality of life in patients with myotonic dystrophy type 2. Journal of the Neurological Sciences, 2016, 365, 158-161.	0.6	24
15	Endothelial nitric oxide synthase gene polymorphisms and prostate cancer risk in <scp>S</scp>erbian population. International Journal of Experimental Pathology, 2013, 94, 355-361.	1.3	22
16	Magnetic resonance imaging of leg muscles in patients with myotonic dystrophies. Journal of Neurology, 2017, 264, 1899-1908.	3.6	21
17	Genetic variants in RNA-induced silencing complex genes and prostate cancer. World Journal of Urology, 2017, 35, 613-624.	2.2	21
18	Genetic and clinical analysis of spinocerebellar ataxia typeâ€ƒ8 repeat expansion in Yugoslavia. Clinical Genetics, 2002, 62, 321-324.	2.0	20

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19	Comparison of temporal and stride characteristics in myotonic dystrophies type 1 and 2 during dual-task walking. Gait and Posture, 2016, 44, 194-199.	1.4	20
20	Multidimensional aspects of pain in myotonic dystrophies. Acta Myologica, 2015, 34, 126-32.	1.5	20
21	A novel recessive TTN founder variant is a common cause of distal myopathy in the Serbian population. European Journal of Human Genetics, 2017, 25, 572-581.	2.8	18
22	Epidemiology of myotonic dystrophy type 1 (Steinert disease) in Belgrade (Serbia). Clinical Neurology and Neurosurgery, 2006, 108, 757-760.	1.4	17
23	Spinocerebellar ataxia type 17 in the Yugoslav population. Acta Neurologica Scandinavica, 2004, 109, 185-187.	2.1	15
24	Effect of childhood general traumas on suicide attempt depends on TPH2 and ADARB1 variants in psychiatric patients. Journal of Neural Transmission, 2017, 124, 621-629.	2.8	15
25	Comparison of the number of triplets inSCA1, MJD/SCA3, HD, SBMA, DRPLA, MD, FRAXA andFRDA genes in schizophrenic patients and a healthy population. American Journal of Medical Genetics Part A, 2000, 96, 884-887.	2.4	14
26	Association between genetic variant in hsa-miR-146a gene and prostate cancer progression: evidence from Serbian population. Cancer Causes and Control, 2014, 25, 1571-1575.	1.8	14
27	Brain sonography insight into the midbrain in myotonic dystrophy type 2. Muscle and Nerve, 2016, 53, 700-704.	2.2	14
28	WDR45 mutations may cause a MECP2 mutation-negative Rett syndrome phenotype. Neurology: Genetics, 2018, 4, e227.	1.9	14
29	Yugoslav HD Phenocopies Analyzed on the Presence of Mutations in Prp, Ferritin, and Jp-3 Genes. International Journal of Neuroscience, 2005, 115, 299-301.	1.6	13
30	Joint effect of the SMN2 and SERF1A genes on childhood-onset types of spinal muscular atrophy in Serbian patients. Journal of Human Genetics, 2015, 60, 723-728.	2.3	12
31	Molecular and Clinical Implications of Variant Repeats in Myotonic Dystrophy Type 1. International Journal of Molecular Sciences, 2022, 23, 354.	4.1	12
32	CTG repeat polymorphism in DMPK gene in healthy Yugoslav population. Acta Neurologica Scandinavica, 2002, 105, 55-58.	2.1	11
33	Mutational analysis of GJB1, MPZ, PMP22, EGR2, and LITAF/SIMPLE in Serbian Charcotâ€Marieâ€Tooth patients. Journal of the Peripheral Nervous System, 2009, 14, 125-136.	3.1	11
34	Body composition analysis in patients with myotonic dystrophy types 1 and 2. Neurological Sciences, 2019, 40, 1035-1040.	1.9	11
35	Genetic Variants within Endothelial Nitric Oxide Synthase Gene and Prostate Cancer: A Meta-Analysis. Clinical and Translational Science, 2015, 8, 23-31.	3.1	10
36	Epidemiology of Rett Syndrome in Serbia: Prevalence, Incidence and Survival. Neuroepidemiology, 2015, 44, 1-5.	2.3	10

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37	Myotonic Dystrophy Type 2 â€“ Data from the Serbian Registry. Journal of Neuromuscular Diseases, 2018, 5, 461-469.	2.6	10
38	The Status of SCA1, MJD/SCA3, FRDA, DRPLA and MD Triplet Containing Genes in Patients with Huntington Disease and Healthy Controls. Journal of Neurogenetics, 2000, 14, 257-263.	1.4	9
39	The Coexistence of Myasthenia Gravis and Myotonic Dystrophy Type 2 in a Single Patient. Journal of		

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55	Assessment of Association between Common Variants at 17q12 and Prostate Cancer Riskâ€"Evidence from Serbian Population and Metaâ€Analysis. Clinical and Translational Science, 2014, 7, 307-313.	3.1	5
56	Genetic testing of individuals with preâ€senile cataract identifies patients with myotonic dystrophy type 2. European Journal of Neurology, 2017, 24, e79-e80.	3.3	5
57	Association between genetic variants in genes encoding Argonaute proteins and cancer risk: A meta-analysis. Pathology Research and Practice, 2020, 216, 152906.	2.3	5
58	Association between genetic variants in DICER1 and cancer risk: An updated meta-analysis. Gene, 2021, 766, 145132.	2.2	5
59	SCA2 and SCA3 mutations in young-onset dopa-responsive parkinsonism. European Journal of Neurology, 2003, 10, 597-597.	3.3	4
60	Tryptophan Hydroxylase 1 Variant rs1800532 is Associated with Suicide Attempt in Serbian Psychiatric Patients but does not Moderate the Effect of Recent Stressful Life Events. Suicide and Life-Threatening Behavior, 2016, 46, 664-668.	1.9	4
61	Neuromyelitis Optica in a Patient from Family with both Myotonic Dystrophy TypeÂ1 and 2. Journal of Neuromuscular Diseases, 2017, 4, 89-92.	2.6	4
62	Metabolic impairments in patients with myotonic dystrophy type 2. Acta Myologica, 2018, 37, 252-256.	1.5	4
63	Poly(A) tailing of ancient DNA: a method for reproducible microsatellite genotyping. Analytical Biochemistry, 2003, 318, 124-131.	2.4	2
64	JP-3 gene polymorphism in a healthy population of Serbia and Montenegro. Journal of Genetics, 2005, 84, 69-71.	0.7	2
65	Coexistence of Unverrichtâ€Lundborg disease and congenital deafness: Molecular resolution of a complex comorbidity. Epilepsia, 2009, 50, 1612-1615.	5.1	2
66	Retinoic acid induced 1 gene and clinical subtypes of schizophrenia: An association study. Psychiatry Research, 2011, 188, 297-298.	3.3	2
67	Cognitive assessment in patients with myotonic dystrophy type 2. Neuromuscular Disorders, 2022, 32, 743-748.	0.6	2
68	Polymorphisms of the Prion Protein Gene (PRNP) in a Serbian Population. International Journal of Neuroscience, 2010, 120, 496-501.	1.6	1
69	Assessment of possible association between rs378854 and prostate cancer risk in the Serbian population. Archives of Biological Sciences, 2013, 65, 475-486.	0.5	0
70	SMN1 copy number as a modifying factor of survival in Serbian patients with sporadic amyotrophic lateral sclerosis. Srpski Arhiv Za Celokupno Lekarstvo, 2018, 146, 646-652.	0.2	0
71	Copy number variants within AZF region of Y chromosome and their association with idiopathic male infertility in Serbian population. Andrologia, 2021, , e14297.	2.1	0