Vidosava Rakocevic-Stojanovic

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2695460/publications.pdf

Version: 2024-02-01

24	347	840776 11	839539 18
papers	citations	h-index	g-index

26	26	26	553
all docs	docs citations	times ranked	citing authors

#	ARTICLE	IF	CITATIONS
1	Employment status of patients with Charcot-Marie-Tooth type 1A. Acta Neurologica Belgica, 2021, , 1.	1.1	1
2	Cerebral involvement and related aspects in myotonic dystrophy type 2. Neuromuscular Disorders, 2021, 31, 681-694.	0.6	5
3	Yield of the PMP22 deletion analysis in patients with compression neuropathies. Journal of Neurology, 2020, 267, 3617-3623.	3.6	0
4	Analysis of duplications versus deletions in the dystrophin gene in Serbian cohort with dystrophinopathies. Vojnosanitetski Pregled, 2020, 77, 387-394.	0.2	0
5	Body composition analysis in patients with myotonic dystrophy types 1 and 2. Neurological Sciences, 2019, 40, 1035-1040.	1.9	11
6	Quality of life in patients with multifocal motor neuropathy from Serbia. Journal of the Neurological Sciences, 2019, 399, 151-154.	0.6	6
7	Phenotypic and genetic spectrum of patients with limb-girdle muscular dystrophy type 2A from Serbia. Acta Myologica, 2019, 38, 163-171.	1.5	5
8	Survival and mortality of adultâ€onset myasthenia gravis in the population of Belgrade, Serbia. Muscle and Nerve, 2018, 58, 708-712.	2.2	12
9	Neuromuscular diseaseâ€specific questionnaire to assess quality of life in patients with chronic inflammatory demyelinating polyradiculoneuropathy. Journal of the Peripheral Nervous System, 2018, 23, 11-16.	3.1	15
10	Quality of life in adult patients with limbâ€“girdle muscular dystrophies. Acta Neurologica Belgica, 2018, 118, 243-250.	1.1	15
11	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
12	Myotonic Dystrophy Type 2 â€“ Data from the Serbian Registry. Journal of Neuromuscular Diseases, 2018, 5, 461-469.	2.6	10
13	Eight years after an international workshop on myotonic dystrophy patient registries: case study of a global collaboration for a rare disease. Orphanet Journal of Rare Diseases, 2018, 13, 155.	2.7	19
14	Metabolic impairments in patients with myotonic dystrophy type 2. Acta Myologica, 2018, 37, 252-256.	1.5	4
15	Diabetes mellitus may affect shortâ€term outcome of Guillainâ€BarrÃ© syndrome. Journal of the Peripheral Nervous System, 2017, 22, 127-130.	3.1	8
16	Magnetic resonance imaging of leg muscles in patients with myotonic dystrophies. Journal of Neurology, 2017, 264, 1899-1908.	3.6	21
17	Personality traits in patients with myotonic dystrophy type 2. Acta Myologica, 2017, 36, 14-18.	1.5	7
18	Brain sonography insight into the midbrain in myotonic dystrophy type 2. Muscle and Nerve, 2016, 53, 700-704.	2.2	14

#	ARTICLE	IF	CITATIONS
19	Prognostic factors and survival of ALS patients from Belgrade, Serbia. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2016, 17, 508-514.	1.7	15
20	Metabolic syndrome in patients with myotonic dystrophy type 1. Muscle and Nerve, 2015, 52, 273-277.	2.2	33
21	Variability of multisystemic features in myotonic dystrophy type 1 â€“ lessons from Serbian registry. Neurological Research, 2015, 37, 939-944.	1.3	12
22	Multidimensional aspects of pain in myotonic dystrophies. Acta Myologica, 2015, 34, 126-32.	1.5	20
23	Cognitive Impairment in Myotonic Dystrophy Type 1 Is Associated with White Matter Damage. PLoS ONE, 2014, 9, e104697.	2.5	76
24	Quantitative analysis of the dystrophin gene by real-time PCR. Archives of Biological Sciences, 2012, 64, 787-792.	0.5	3