MiloÅ; BrkuÅ;anin

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4462616/publications.pdf

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10	155	7	9
papers	citations	h-index	g-index
10	10	10	473 citing authors
all docs	docs citations	times ranked	

#	Article	IF	CITATIONS
1	Fatigue in myotonic dystrophy type 1: a seven-year prospective study. Acta Myologica, 2019, 38, 239-244.	1.5	4
2	Trichome-specific and developmentally regulated biosynthesis of nepetalactones in leaves of cultivated Nepeta rtanjensis plants. Industrial Crops and Products, 2018, 117, 347-358.	5. 2	16
3	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
4	Myotonic Dystrophy Type 2 – Data from the Serbian Registry. Journal of Neuromuscular Diseases, 2018, 5, 461-469.	2.6	10
5	Effect of the introgression of Atlantic brown trout, Salmo trutta, into Adriatic trout, Salmo farioides in a stream at the drainage area of the Adriatic Sea basin of Montenegro. Acta Ichthyologica Et Piscatoria, 2018, 48, 363-372.	0.7	3
6	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
7	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
8	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
9	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
10	Molecular Genetics and Genetic Testing in Myotonic Dystrophy Type 1. BioMed Research International, 2013, 2013, 1-13.	1.9	42