## Blažo Nikolić

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3043370/publications.pdf

Version: 2024-02-01

1684188 1720034 14 67 5 7 citations g-index h-index papers 14 14 14 119 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Multiple Sclerosis in Pediatrics: Current Concepts and Treatment Options. Neurology and Therapy, 2016, 5, 131-143.	3.2	20
2	Analysis of secondary mtDNA mutations in families with Leber's hereditary optic neuropathy: Four novel variants and their association with clinical presentation. Mitochondrion, 2020, 50, 132-138.	3.4	9
3	Oxidative Stress Profile in Genetically Confirmed Cases of Leber's Hereditary Optic Neuropathy. Journal of Molecular Neuroscience, 2021, 71, 1070-1081.	2.3	9
4	A 6-month follow-up of disability, quality of life, and depressive and anxiety symptoms in pediatric migraine with magnesium prophylaxis. Magnesium Research, 2017, 30, 133-141.	0.5	6
5	Current Use of mTOR Inhibitors for the Treatment of Subependymal Giant Cell Astrocytomas and Epilepsy in Patients with TSC. Current Medicinal Chemistry, 2016, 23, 4260-4269.	2.4	6
6	Characteristics of pediatric multiple sclerosis: A tertiary referral center study. PLoS ONE, 2020, 15, e0243031.	2.5	4
7	Leber's Hereditary Optic Neuropathy: Novel Views and Persisting Challenges. CNS and Neurological Disorders - Drug Targets, 2018, 16, 927-935.	1.4	3
8	In silico model of mtDNA mutations effect on secondary and 3D structure of mitochondrial rRNA and tRNA in Leber's hereditary optic neuropathy. Experimental Eye Research, 2020, 201, 108277.	2.6	2
9	Visual evoked potentials - current concepts and future perspectives. Vojnosanitetski Pregled, 2018, 75, 496-503.	0.2	2
10	Does handedness matter? Writing and tracing kinematic analysis in healthy adults. Psihologija, 2019, 52, 413-435.	0.6	2
11	The usefulness of visual evoked potentials in the assessment of the pediatric multiple sclerosis. European Journal of Paediatric Neurology, 2022, 36, 130-136.	1.6	2
12	Multiple Sclerosis Therapies in Pediatric Patients: Challenges and Opportunities., 0,, 39-52.		1
13	GLUT1 deficiency syndrome: A case report with a novel SLC2A1 mutation. Vojnosanitetski Pregled, 2019, 76, 543-546.	0.2	1
14	Child Neurology: <i>Bartonella henselae</i> Neuroretinitis in 2 Patients. Neurology, 2022, , 10.1212/WNL.00000000000572.	1.1	0