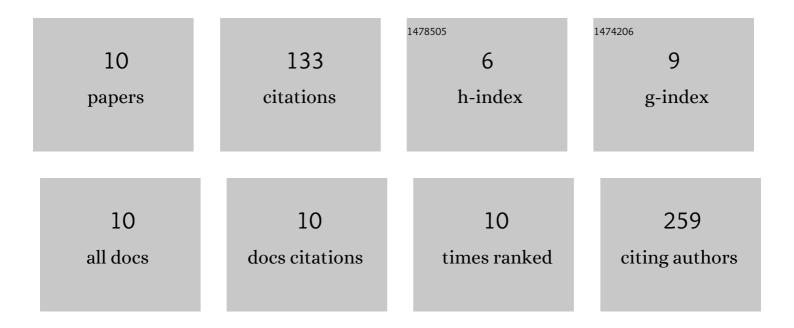
Adrijan Sarajlija

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/10715253/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Acute disseminated encephalomyelitis in children and adolescents - 20-year single-center experience in Serbia. Srpski Arhiv Za Celokupno Lekarstvo, 2022, 150, 544-550.	0.2	0
2	Clinical and genetic characteristics of patients with congenital hyperinsulinism in 21 non-consanguineous families from Serbia. European Journal of Pediatrics, 2021, 180, 2815-2821.	2.7	1
3	Impact of genotype on neutropenia in a large cohort of Serbian patients with glycogen storage disease type Ib. European Journal of Medical Genetics, 2020, 63, 103767.	1.3	8
4	Early prediction of phenotypic severity in Citrullinemia Type 1. Annals of Clinical and Translational Neurology, 2019, 6, 1858-1871.	3.7	26
5	Transatlantic combined and comparative data analysis of 1095 patients with urea cycle disorders—A successful strategy for clinical research of rare diseases. Journal of Inherited Metabolic Disease, 2019, 42, 93-106.	3.6	35
6	<i>WDR45</i> mutations may cause a <i>MECP2</i> mutation-negative Rett syndrome phenotype. Neurology: Genetics, 2018, 4, e227.	1.9	14
7	Appendiceal involvement in a patient with Gaucher disease. Blood Cells, Molecules, and Diseases, 2018, 68, 109-111.	1.4	2
8	Early Presentation of Hyperinsulinism/Hyperammonemia Syndrome in Three Serbian Patients. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 228-231.	0.9	4
9	Health-related quality of life and depression in Rett syndrome caregivers. Vojnosanitetski Pregled, 2013, 70, 842-847.	0.2	21
10	Molecular Genetics and Genotype-Based Estimation of BH4-Responsiveness in Serbian PKU Patients: Spotlight on Phenotypic Implications of p.L48S. JIMD Reports, 2012, 9, 49-58.	1.5	22