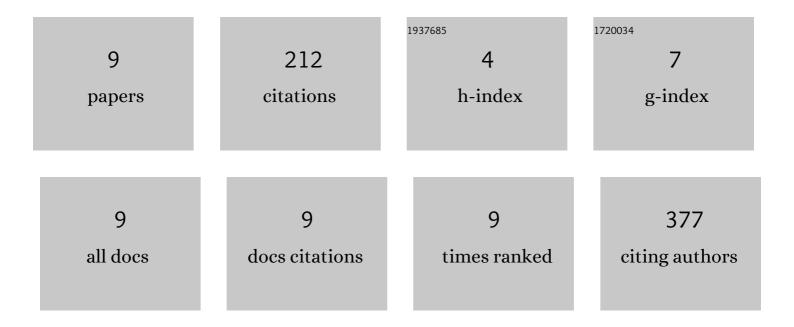
## Danijela Radivojevic

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7340716/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Prevalence of variants in DFNB1 locus in Serbian patients with autosomal recessive non-syndromic hearing loss. Genetika, 2022, 54, 447-456.	0.4	0
2	Audiological features in Serbian patients with hearing impairment identified with c.35delG in the GJB2 gene. Srpski Arhiv Za Celokupno Lekarstvo, 2021, 149, 685-690.	0.2	0
3	Novel intragenic deletions within the UBE3A gene in two unrelated patients with Angelman syndrome: case report and review of the literature. BMC Medical Genetics, 2017, 18, 137.	2.1	4
4	Ten years of experience in molecular prenatal diagnosis and carrier testing for spinal muscular atrophy among families from Serbia. International Journal of Gynecology and Obstetrics, 2014, 124, 55-58.	2.3	6
5	Newborn screening for cystic fibrosis in <scp>S</scp> erbia: A pilot study. Pediatrics International, 2013, 55, 181-184.	0.5	2
6	Lower Incidence of Deletions in the Survival of Motor Neuron Gene and the Neuronal Apoptosis Inhibitory Protein Gene in Children with Spinal Muscular Atrophy from Serbia. Tohoku Journal of Experimental Medicine, 2011, 225, 153-159.	1.2	6
7	Analysis of extra- and intragenic marker haplotypes as part of molecular diagnosis of cystic fibrosis in patients from Serbia. Archives of Biological Sciences, 2008, 60, 5-10.	0.5	3
8	Deletion and duplication screening in the DMD gene using MLPA. European Journal of Human Genetics, 2005, 13, 1231-1234.	2.8	171
9	Spectrum of Cystic Fibrosis Mutations in Serbia and Montenegro and Strategy for Prenatal Diagnosis. Genetic Testing and Molecular Biomarkers, 2004, 8, 276-280.	1.7	20