

# Danijela Radivojevic

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7340716/publications.pdf>

Version: 2024-02-01

9  
papers

212  
citations

1937685

4  
h-index

1720034

7  
g-index

9  
all docs

9  
docs citations

9  
times ranked

377  
citing authors

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