

# Ozlem Giray Bozkaya

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7877463/publications.pdf>

Version: 2024-02-01

8  
papers

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citations

2682572

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h-index

2550090

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8  
docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium. <i>Functional and Integrative Genomics</i> , 2022, 22, 291-315.	3.5	7
2	Identification of a novel frameshift heterozygous deletion in exon 8 of the PAX6 gene in a pedigree with aniridia. <i>Molecular Medicine Reports</i> , 2016, 14, 2150-2154.	2.4	3
3	A Novel Mutation in the Mitochondrial DNA Cytochrome b Gene ( <i>MTCYB</i> ) in a Patient with Prader Willi Syndrome. <i>Journal of Child Neurology</i> , 2015, 30, 378-381.	1.4	2
4	A newborn with monosomy X in association with corpus callosum agenesis. <i>Pediatrics and Neonatology</i> , 2017, 58, 455-457.	0.9	0
5	Identification of the largest homozygous glycine decarboxylase gene deletion in a Turkish infant. <i>Pediatrics and Neonatology</i> , 2018, 59, 632-633.	0.9	0
6	Investigation of the most common clinical and imaging findings and the role of tubulin genes in the etiology of malformations of cortical development. <i>Turkish Journal of Medical Sciences</i> , 2020, 50, 1573-1579.	0.9	0
7	Glutathione S-Transferase Gene Polymorphisms in Children with Down Syndrome and Their Mothers. <i>International Journal of Human Genetics</i> , 2015, 15, 33-39.	0.1	0
8	Blended Phenotype of Pelger-Huet Anomaly with Osteochondroma and Autosomal Recessive Deafness with Enlarged Vestibular Aqueduct. <i>Molecular Syndromology</i> , 0, , 1-6.	0.8	0