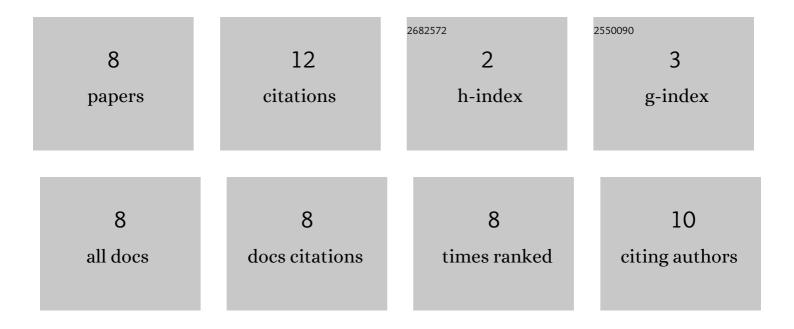
## Ozlem Giray Bozkaya

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7877463/publications.pdf Version: 2024-02-01



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