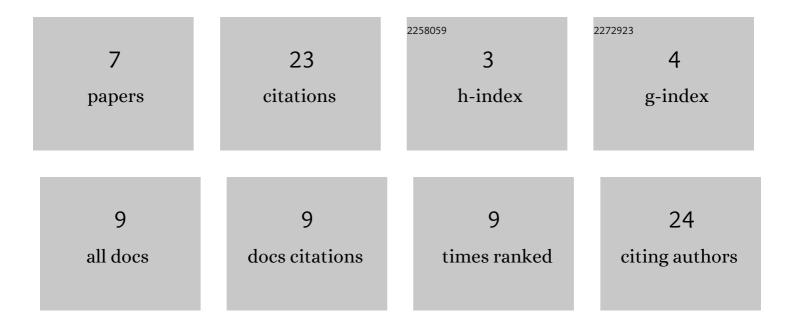
## Mustafa Dogan

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7403067/publications.pdf Version: 2024-02-01



#	Article	IF	Citations
1	Biallelic Novel USP53 Splicing Variant Disrupting the Gene Function that Causes Cholestasis Phenotype and Review of the Literature. Molecular Syndromology, 2022, 13, 471-484.	0.8	4
2	Deficiency of alkaline ceramidase 3 with infancy-onset progressive leukoencephalopathy: a second case report. Acta Neurologica Belgica, 2021, 121, 1867-1870.	1.1	2
3	Clinical and molecular findings in a Turkish family with an ultra-rare condition, ELP2-related neurodevelopmental disorder. Molecular Biology Reports, 2021, 48, 701-708.	2.3	4
4	Clinical, radiological and computational studies on two novel GNPTG variants causing mucolipidosis III gamma phenotypes with varying severity. Molecular Biology Reports, 2021, 48, 1465-1474.	2.3	2
5	Chorioretinal dystrophy, hypogonadotropic hypogonadism, and cerebellar ataxia: Boucher-Neuhauser syndrome due to a homozygous (c.3524C>G (p.Ser1175Cys)) variant in <i>PNPLA6</i> gene. Ophthalmic Genetics, 2021, 42, 276-282.	1.2	7
6	A case with Turner syndrome and alopecia areata. Ortadoğu Tıp Dergisi, 2019, 11, 631-633.	0.1	0
7	Study of ten causal genes in Turkish patients with clinically suspected maturity-onset diabetes of the young (MODY) using a targeted next-generation sequencing panel. Molecular Biology Reports, 0, , .	2.3	4