

# Mustafa Dogan

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7403067/publications.pdf>

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2258059  
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#	ARTICLE	IF	CITATIONS
1	Biallelic Novel USP53 Splicing Variant Disrupting the Gene Function that Causes Cholestasis Phenotype and Review of the Literature. <i>Molecular Syndromology</i> , 2022, 13, 471-484.	0.8	4
2	Deficiency of alkaline ceramidase 3 with infancy-onset progressive leukoencephalopathy: a second case report. <i>Acta Neurologica Belgica</i> , 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. <i>Molecular Biology Reports</i> , 2021, 48, 701-708.	2.3	4
4	Clinical, radiological and computational studies on two novel GNPTG variants causing mucopolipidosis III gamma phenotypes with varying severity. <i>Molecular Biology Reports</i> , 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. <i>Ophthalmic Genetics</i> , 2021, 42, 276-282.	1.2	7
6	A case with Turner syndrome and alopecia areata. <i>OrtadoÄŸu TÄ±p Dergisi</i> , 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. <i>Molecular Biology Reports</i> , 0, , .	2.3	4