

# Ã-zlem YalÃ§Ä±n Äapan

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5989783/publications.pdf>

Version: 2024-02-01

7  
papers

115  
citations

1937685

4  
h-index

1720034

7  
g-index

7  
all docs

7  
docs citations

7  
times ranked

233  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genes and molecular mechanisms involved in the epileptogenesis of idiopathic absence epilepsies. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2012, 21, 79-86.	2.0	74
2	SCN1A gene sequencing in 46 Turkish epilepsy patients disclosed 12 novel mutations. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2016, 39, 34-43.	2.0	13
3	A Review of Functional Characterization of Single Amino Acid Change Mutations in HNF Transcription Factors in MODY Pathogenesis. <i>Protein Journal</i> , 2021, 40, 348-360.	1.6	11
4	De novo 8p23.1 deletion in a patient with absence epilepsy. <i>Epileptic Disorders</i> , 2017, 19, 217-221.	1.3	8
5	A Common VWF Exon 28 Haplotype in the Turkish Population. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2013, 19, 550-556.	1.7	4
6	Whole exome sequencing reveals novel candidate gene variants for MODY. <i>Clinica Chimica Acta</i> , 2020, 510, 97-104.	1.1	4
7	HNF1A-MODY Mutations in Nuclear Localization Signal Impair HNF1A-Import Receptor KPNA6 Interactions. <i>Protein Journal</i> , 2021, 40, 512-521.	1.6	1