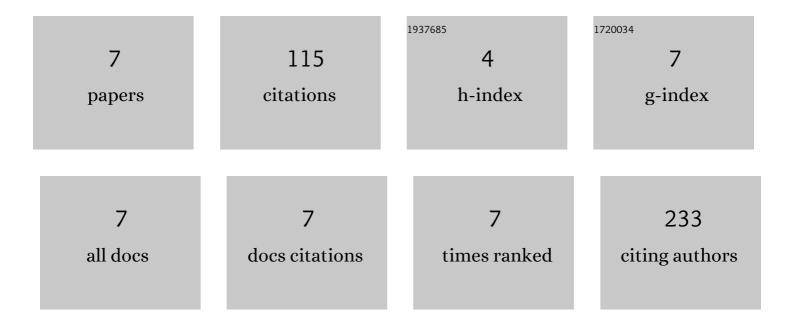
## Ã−zlem Yalçın Ã**‡**pan

List of Publications by Year in descending order

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

Version: 2024-02-01



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