

# Haktan BaÄÄ±Å Erdem

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4498009/publications.pdf>

Version: 2024-02-01

13

papers

146

citations

1478505

6

h-index

1199594

12

g-index

13

all docs

13

docs citations

13

times ranked

393

citing authors

#	ARTICLE	IF	CITATIONS
1	The Genomics of Arthrogryposis, a Complex Trait: Candidate Genes and Further Evidence for Oligogenic Inheritance. American Journal of Human Genetics, 2019, 105, 132-150.	6.2	74
2	Spectrum of BRCA1/BRCA2 variants in 1419 Turkish breast and ovarian cancer patients: a single center study. Turkish Journal of Biochemistry, 2020, 45, 83-90.	0.5	12
3	Correlation with Platelet Parameters and Genetic Markers of Thrombophilia Panel (Factor II) Tj ETQq1 1 0.784314 rgBT /Overlock 10 T 5	3.4	11
4	SMN1 gene copy number analysis for spinal muscular atrophy (SMA) in a Turkish cohort by CODE-SEQ technology, an integrated solution for detection of SMN1 and SMN2 copy numbers and the $\alpha\text{2+0}$ genotype. Neurological Sciences, 2020, 41, 2575-2584.	1.9	10
5	Function of telomere in aging and age related diseases. Environmental Toxicology and Pharmacology, 2021, 85, 103641.	4.0	10
6	Importance and usage of chromosomal microarray analysis in diagnosing intellectual disability, global developmental delay, and autism; and discovering new loci for these disorders. Molecular Cytogenetics, 2018, 11, 54.	0.9	9
7	Association Between Human Hair Loss and the Expression Levels of Nucleolin, Nucleophosmin, and UBTF Genes. Genetic Testing and Molecular Biomarkers, 2016, 20, 197-202.	0.7	6
8	The Frequency of Monocyte Chemoattractant Protein-1 Gene Polymorphism in Obstructive Sleep Apnea Syndrome. Lung, 2019, 197, 585-592.	3.3	4
9	A rare cause of dyspnea in emergency medicine: Keutel syndrome. American Journal of Emergency Medicine, 2016, 34, 935.e3-935.e5.	1.6	3
10	Spectrum of germline cancer susceptibility gene mutations in Turkish colorectal cancer patients: a single center study. Turkish Journal of Medical Sciences, 2020, 50, 1015-1021.	0.9	3
11	TEKRARLAYAN DÄZÄK VE TAM KAN SAYIMI DEÄZERLERÄ ARASINDA HERHANGÄ BÄR Ä°LÄÅZKÄ VAR MI? BÄR VAKÄ KONTROL Ã‡ALIÅMASI. Konuralp Tip Dergisi, 2020, 12, 39-43.	0.3	
12	Mitochondrial DNA deletions in patients with chronic suppurative otitis media. European Archives of Oto-Rhino-Laryngology, 2016, 273, 2473-2479.	1.6	1
13	An Investigation of Genetic Polymorphism In The Rs35521 Serotonin Transporter Gene In Allergic Rhinitis. Ent Updates, 0, . .	0.0	0