Haktan BaÄÄ±Å Erdem

List of Publications by Year
in descending order

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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	Function of telomere in aging and age related diseases. Environmental Toxicology and Pharmacology, 2021, 85, 103641.	4.0	10
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	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
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 â€œ2+0â€• genotype. Neurological Sciences, 2020, 41, 2575-2584.	1.9	10
5	TEKRARLAYAN DÃœÅžÃœK VE TAM KAN SAYIMI DEÄžERLERÄ° ARASINDA HERHANGÄ° BÄ°R Ä°LÄ°ÅžKÄ° VAR MI? BÄ°R VAKA-KONTROL Ã‡ALIÅžMASI. Konuralp Tip Dergisi, 2020, 12, 39-43.	0.3	3
6	The Frequency of Monocyte Chemoattractant Protein-1 Gene Polymorphism in Obstructive Sleep Apnea Syndrome. Lung, 2019, 197, 585-592.	3.3	4
7	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
8	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
9	Mitochondrial DNA deletions in patients with chronic suppurative otitis media. European Archives of Oto-Rhino-Laryngology, 2016, 273, 2473-2479.	1.6	1
10	Association Between Human Hair Loss and the Expression Levels of Nucleolin, Nucleophosmin, andUBTFGenes. Genetic Testing and Molecular Biomarkers, 2016, 20, 197-202.	0.7	6
11	Correlation with Platelet Parameters and Genetic Markers of Thrombophilia Panel (Factor II) Tj ETQq1 1 0.784314 rgBT /Overlock 10 T	3.4	11
12	A rare cause of dyspnea in emergency medicine: Keutel syndrome. American Journal of Emergency Medicine, 2016, 34, 935.e3-935.e5.	1.6	3
13	An Investigation of Genetic Polymorphism In The Rs35521 Serotonin Transporter Gene In Allergic Rhinitis. Ent Updates, 0, , .	0.0	0