Seda TaÅÄ±r YÄ±lmaz

List of Publications by Year
in descending order

7 papers	197 citations	1684188 5 h-index	1872680 6 g-index
7 all docs	7 docs citations	7 times ranked	443 citing authors

#	Article	IF	Citations
1	Homozygous Mutations in Fibroblast Growth Factor 3 Are Associated with a New Form of Syndromic Deafness Characterized by Inner Ear Agenesis, Microtia, and Microdontia. American Journal of Human Genetics, 2007, 80, 338-344.	6.2	97
2	Mutations in TMC1 contribute significantly to nonsyndromic autosomal recessive sensorineural hearing loss: A report of five novel mutations. International Journal of Pediatric Otorhinolaryngology, 2009, 73, 699-705.	1.0	51
3	A founder TMIE mutation is a frequent cause of hearing loss in southeastern Anatolia. Clinical Genetics, 2009, 75, 562-567.	2.0	20
4	BehÃ§et's: A Disease or a Syndrome? Answer from an Expression Profiling Study. PLoS ONE, 2016, 11, e0149052.	2.5	12
5	Matrilin-3 as a putative effector of C-type natriuretic peptide signaling during TGF-Î² induced chondrogenic differentiation of mesenchymal stem cells. Molecular Biology Reports, 2014, 41, 5549-5555.	2.3	11
6	C-type lectin domain family 12, member A: A common denominator in BehÃ§etâ€™s syndrome and acute gouty arthritis. Medical Hypotheses, 2015, 85, 186-191.	1.5	6
7	Transcriptomic Profile of Bone Marrow-Derived Mesenchymal Stromal Cells of Pediatric Pre-B Acute Lymphoblastic Leukemia Patients and Healthy Donors after Interaction with Leukemic Cells. Blood, 2019, 134, 3957-3957.	1.4	0