ArmaÄan Ä°ncesulu

List of Publications by Year
in descending order

26 papers	1,247 citations	687363 13 h-index	713466 21 g-index
27 all docs	27 docs citations	27 times ranked	1663 citing authors

#	ARTICLE	IF	CITATIONS
1	GJB2 Mutations and Degree of Hearing Loss: A Multicenter Study. American Journal of Human Genetics, 2005, 77, 945-957.	6.2	455
2	EAONO/JOS Joint Consensus Statements on the Definitions, Classification and Staging of Middle Ear Cholesteatoma. Journal of International Advanced Otology, 2017, 13, 1-8.	1.0	181
3	Comprehensive analysis via exome sequencing uncovers genetic etiology in autosomal recessive nonsyndromic deafness in a large multiethnic cohort. Genetics in Medicine, 2016, 18, 364-371.	2.4	124
4	Revision Stapedectomy: Intraoperative Findings, Results, and Review of the Literature. Laryngoscope, 1997, 107, 1185-1192.	2.0	75
5	Spectrum of GJB2 mutations in Turkey comprises both Caucasian and Oriental variants: Roles of parental consanguinity and assortative mating. Human Mutation, 2003, 21, 552-553.	2.5	61
6	Cochlear implantation in cases with incomplete partition type III (X-linked anomaly). European Archives of Oto-Rhino-Laryngology, 2008, 265, 1425-1430.	1.6	56
7	A novel missense mutation in a C2 domain ofOTOF results in autosomal recessive auditory neuropathy. American Journal of Medical Genetics, Part A, 2005, 138A, 6-10.	1.2	50
8	Phenotypic variability of patients homozygous for the GJB2 mutation 35delG cannot be explained by the influence of one major modifier gene. European Journal of Human Genetics, 2009, 17, 517-524.	2.8	46
9	International Otology Outcome Group and the International Consensus on the Categorization of Tympanomastoid Surgery. Journal of International Advanced Otology, 2018, 14, 216-226.	1.0	46
10	Recurrent and Private MYO15A Mutations Are Associated with Deafness in the Turkish Population. Genetic Testing and Molecular Biomarkers, 2010, 14, 543-550.	0.7	45
11	FOXF2is required for cochlear development in humans and mice. Human Molecular Genetics, 2019, 28, 1286-1297.	2.9	20
12	MPZL2 is a novel gene associated with autosomal recessive nonsyndromic moderate hearing loss. Human Genetics, 2018, 137, 479-486.	3.8	19
13	Effects of GJB2 genotypes on the audiological phenotype: Variability is present for all genotypes. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 1687-1694.	1.0	18
14	Computed Tomographic Findings of X-Linked Deafness. Journal of Computer Assisted Tomography, 2014, 38, 20-24.	0.9	14
15	Novel pathogenic variants underlie SLC26A4 -related hearing loss in a multiethnic cohort. International Journal of Pediatric Otorhinolaryngology, 2017, 101, 167-171.	1.0	11
16	Long-range cis-regulatory elements controlling GDF6 expression are essential for ear development. Journal of Clinical Investigation, 2020, 130, 4213-4217.	8.2	9
17	Women in otolaryngology in Turkey: Insight of gender equality, career development and work-life balance. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2020, 41, 102305.	1.3	7
18	The Effect of Nasal Obstruction after Different Nasal Surgeries Using Acoustic Rhinometry and Nasal Obstruction Symptom Evaluation Scale. World Journal of Plastic Surgery, 2016, 5, 236-243.	0.6	6

#	Article	IF	Citations
19	Perception of male otolaryngologists on gender discrimination: a comparative study. European Archives of Oto-Rhino-Laryngology, 2021, 278, 1257-1264.	1.6	2
20	Association Between Incomplete Partition Type III and Abnormal Hypothalamic Morphology: Further Imaging Evidence. Journal of Computer Assisted Tomography, 2020, 44, 704-707.	0.9	1
21	Irregular Contour of Inner Ear Structures and Hypomineralized Areas at Otic Capsule. Journal of Computer Assisted Tomography, 2020, 44, 386-388.	0.9	1
22	Cochlear Implantation in Patients with Keratitis-Ichthyosis-Deafness Syndrome: A Report of Two Cases. Case Reports in Otolaryngology, 2017, 2017, 1-5.	0.2	0
23	Anesthesia for a child with congenital long QT syndrome, a case report and literature review. Anesthesia: Essays and Researches, 2021, 15, 149.	0.5	0
24	From Past to Present: The Journey of Female Doctors in Medicine and Otorhinolaryngology in Turkey. Turkish Archives of Otorhinolaryngology, 2021, 59, 166-171.	0.5	0
25	Diagnosis and Differential Diagnosis of Disorders of Hearing Development. European Manual of Medicine, 2020, , 857-961.	0.1	0
26	Effects of Oxygen Therapies in Experimental Acute Acoustic Trauma. , 2021, 17, 508-513.		0