

ArmaÄan Ä°ncesulu

List of Publications by Year in descending order

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Version: 2024-02-01

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	Perception of male otolaryngologists on gender discrimination: a comparative study. <i>European Archives of Oto-Rhino-Laryngology</i> , 2021, 278, 1257-1264.	1.6	2
2	Anesthesia for a child with congenital long QT syndrome, a case report and literature review. <i>Anesthesia: Essays and Researches</i> , 2021, 15, 149.	0.5	0
3	From Past to Present: The Journey of Female Doctors in Medicine and Otorhinolaryngology in Turkey. <i>Turkish Archives of Otorhinolaryngology</i> , 2021, 59, 166-171.	0.5	0
4	Effects of Oxygen Therapies in Experimental Acute Acoustic Trauma. , 2021, 17, 508-513.		0
5	Women in otolaryngology in Turkey: Insight of gender equality, career development and work-life balance. <i>American Journal of Otolaryngology - Head and Neck Medicine and Surgery</i> , 2020, 41, 102305.	1.3	7
6	Association Between Incomplete Partition Type III and Abnormal Hypothalamic Morphology: Further Imaging Evidence. <i>Journal of Computer Assisted Tomography</i> , 2020, 44, 704-707.	0.9	1
7	Irregular Contour of Inner Ear Structures and Hypomineralized Areas at Otic Capsule. <i>Journal of Computer Assisted Tomography</i> , 2020, 44, 386-388.	0.9	1
8	Long-range cis-regulatory elements controlling GDF6 expression are essential for ear development. <i>Journal of Clinical Investigation</i> , 2020, 130, 4213-4217.	8.2	9
9	Diagnosis and Differential Diagnosis of Disorders of Hearing Development. <i>European Manual of Medicine</i> , 2020, , 857-961.	0.1	0
10	FOXF2 is required for cochlear development in humans and mice. <i>Human Molecular Genetics</i> , 2019, 28, 1286-1297.	2.9	20
11	MPZL2 is a novel gene associated with autosomal recessive nonsyndromic moderate hearing loss. <i>Human Genetics</i> , 2018, 137, 479-486.	3.8	19
12	International Otology Outcome Group and the International Consensus on the Categorization of Tympanomastoid Surgery. <i>Journal of International Advanced Otology</i> , 2018, 14, 216-226.	1.0	46
13	Novel pathogenic variants underlie SLC26A4 -related hearing loss in a multiethnic cohort. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2017, 101, 167-171.	1.0	11
14	Cochlear Implantation in Patients with Keratitis-Ichthyosis-Deafness Syndrome: A Report of Two Cases. <i>Case Reports in Otolaryngology</i> , 2017, 2017, 1-5.	0.2	0
15	EAONO/JOS Joint Consensus Statements on the Definitions, Classification and Staging of Middle Ear Cholesteatoma. <i>Journal of International Advanced Otology</i> , 2017, 13, 1-8.	1.0	181
16	Comprehensive analysis via exome sequencing uncovers genetic etiology in autosomal recessive nonsyndromic deafness in a large multiethnic cohort. <i>Genetics in Medicine</i> , 2016, 18, 364-371.	2.4	124
17	The Effect of Nasal Obstruction after Different Nasal Surgeries Using Acoustic Rhinometry and Nasal Obstruction Symptom Evaluation Scale. <i>World Journal of Plastic Surgery</i> , 2016, 5, 236-243.	0.6	6
18	Computed Tomographic Findings of X-Linked Deafness. <i>Journal of Computer Assisted Tomography</i> , 2014, 38, 20-24.	0.9	14

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19	Recurrent and Private <i>MYO15A</i> Mutations Are Associated with Deafness in the Turkish Population. <i>Genetic Testing and Molecular Biomarkers</i> , 2010, 14, 543-550.	0.7	45
20	Phenotypic variability of patients homozygous for the GJB2 mutation 35delG cannot be explained by the influence of one major modifier gene. <i>European Journal of Human Genetics</i> , 2009, 17, 517-524.	2.8	46
21	Cochlear implantation in cases with incomplete partition type III (X-linked anomaly). <i>European Archives of Oto-Rhino-Laryngology</i> , 2008, 265, 1425-1430.	1.6	56
22	Effects of GJB2 genotypes on the audiological phenotype: Variability is present for all genotypes. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2006, 70, 1687-1694.	1.0	18
23	A novel missense mutation in a C2 domain of OTOF results in autosomal recessive auditory neuropathy. <i>American Journal of Medical Genetics, Part A</i> , 2005, 138A, 6-10.	1.2	50
24	GJB2 Mutations and Degree of Hearing Loss: A Multicenter Study. <i>American Journal of Human Genetics</i> , 2005, 77, 945-957.	6.2	455
25	Spectrum of GJB2 mutations in Turkey comprises both Caucasian and Oriental variants: Roles of parental consanguinity and assortative mating. <i>Human Mutation</i> , 2003, 21, 552-553.	2.5	61
26	Revision Stapedectomy: Intraoperative Findings, Results, and Review of the Literature. <i>Laryngoscope</i> , 1997, 107, 1185-1192.	2.0	75