

# Yeliz Guven

## List of Publications by Year in descending order

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Version: 2024-02-01

29  
papers

472  
citations

687363

13  
h-index

713466

21  
g-index

29  
all docs

29  
docs citations

29  
times ranked

1095  
citing authors

#	ARTICLE	IF	CITATIONS
1	Expanding genotypic and phenotypic spectrums of <sc><i>LTBP3</i></sc> variants in dental anomalies and short stature syndrome. <i>Clinical Genetics</i> , 2022, 102, 66-71.	2.0	5
2	Characteristic dental pattern with hypodontia and short roots in Fraser syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 1681-1689.	1.2	7
3	Antimicrobial Effect of Newly Formulated Toothpastes and a Mouthrinse on Specific Microorganisms: An In Vitro Study. <i>European Journal of Dentistry</i> , 2019, 13, 172-177.	1.7	12
4	Turkish Ectodermal Dysplasia Cohort: From Phenotype to Genotype in 17 Families. <i>Cytogenetic and Genome Research</i> , 2019, 157, 189-196.	1.1	7
5	Oral Bacteria of Children with Turner Syndrome. <i>Journal of Pediatric Research</i> , 2019, 6, 44-50.	0.2	1
6	A biallelic <i>ANTXR1</i> variant expands the anthrax toxin receptor associated phenotype to tooth agenesis. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 1015-1022.	1.2	11
7	Identification of likely pathogenic and known variants in TSPEAR, LAMB3, BCOR, and WNT10A in four Turkish families with tooth agenesis. <i>Human Genetics</i> , 2018, 137, 689-703.	3.8	24
8	Prevalence of ectopic eruption of first permanent molars in a Turkish population. <i>European Oral Research</i> , 2018, 52, 1-5.	0.9	14
9	Assessment of the endodontic microbiota of abscessed primary teeth using microarray technology. <i>Indian Journal of Dental Research</i> , 2018, 29, 781.	0.4	5
10	Periodontal disease and FAM20A mutations. <i>Journal of Human Genetics</i> , 2017, 62, 679-686.	2.3	19
11	Cleidocranial dysplasia: Clinical, endocrinologic and molecular findings in 15 patients from 11 families. <i>European Journal of Medical Genetics</i> , 2017, 60, 163-168.	1.3	31
12	Success Rates of Pulpotomies in Primary Molars Using Calcium Silicate-Based Materials: A Randomized Control Trial. <i>BioMed Research International</i> , 2017, 2017, 1-7.	1.9	11
13	Long-Term Fracture Resistance of Simulated Immature Teeth Filled with Various Calcium Silicate-Based Materials. <i>BioMed Research International</i> , 2016, 2016, 1-6.	1.9	13
14	Prevalence and characteristics of talon cusps in Turkish population. <i>Dental Research Journal</i> , 2016, 13, 145.	0.6	8
15	A Complex Facial Trauma Case with Multiple Mandibular Fractures and Dentoalveolar Injuries. <i>Case Reports in Dentistry</i> , 2015, 2015, 1-6.	0.5	2
16	THE PREVALENCE OF DENTAL ANOMALIES IN A TURKISH POPULATION. <i>Journal of Istanbul University Faculty of Dentistry</i> , 2015, 49, 23.	0.2	14
17	Shear bond strength and ultrastructural interface analysis of different adhesive systems to Er:YAG laser-prepared dentin. <i>Lasers in Medical Science</i> , 2015, 30, 769-778.	2.1	23
18	Preservation of Involved Teeth Associated with Large Dentigerous Cysts. <i>International Scholarly Research Notices</i> , 2014, 2014, 1-5.	0.9	1

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19	Enamelâ€Renalâ€Gingival syndrome, hypodontia, and a novel <i>FAM20A</i> mutation. American Journal of Medical Genetics, Part A, 2014, 164, 2124-2128.	1.2	21
20	X-ray diffraction analysis of MTA-Plus, MTA-Angelus and DiaRoot BioAggregate. European Journal of Dentistry, 2014, 08, 211-215.	1.7	29
21	C5orf42 is the major gene responsible for OFD syndrome type VI. Human Genetics, 2014, 133, 367-377.	3.8	71
22	Clinical manifestations of 17 patients affected with mucopolysaccharidosis type VI and eight novel <i>ARSB</i> mutations. American Journal of Medical Genetics, Part A, 2014, 164, 1443-1453.	1.2	20
23	Oral manifestations of 17 patients affected with mucopolysaccharidosis type VI. Journal of Inherited Metabolic Disease, 2014, 37, 263-268.	3.6	32
24	Twins with hereditary sensory and autonomic neuropathy type IV with preserved periodontal sensation. European Journal of Medical Genetics, 2014, 57, 240-246.	1.3	9
25	A novel c.1255G>T (p.D419Y) mutation in SH3BP2 gene causes cherubism in a Turkish family. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2012, 114, e42-e46.	0.4	8
26	Success rates of mineral trioxide aggregate, ferric sulfate, and formocresol pulpotomies: a 24-month study. Pediatric Dentistry (discontinued), 2011, 33, 165-70.	0.4	49
27	Assessment of dental features in 16 children with hypohidrotic ectodermal dysplasia. Pediatric Dental Journal, 2009, 19, 106-111.	0.7	2
28	Oro dental findings of a family with lacrimo-auriculo-dento digital (LADD) syndrome. Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics, 2008, 106, e33-e44.	1.4	20
29	Effect of three different remineralizing agents on artificial erosive lesions of primary teeth. Australian Dental Journal, 0, , .	1.5	3