Thanakorn Theerapanon

List of Publications by Year
in descending order

19 papers	210 citations	1040056 9 h-index	1058476 14 g-index
19 all docs	19 docs citations	19 times ranked	288 citing authors

#	Article	IF	Citations
1	PTEN regulates proliferation and osteogenesis of dental pulp cells and adipogenesis of human adiposeâ€derived stem cells. Oral Diseases, 2023, 29, 735-746.	3.0	3
2	Ageâ€related dental phenotypes and tooth characteristics of FAM83Hâ€associated hypocalcified amelogenesis imperfecta. Oral Diseases, 2022, 28, 734-744.	3.0	3
3	Patterns of molar agenesis associated with p.P20L and p.R77Q variants in PAX9. European Journal of Oral Sciences, 2022, 130, e12855.	1.5	3
4	Tooth ultrastructure of a novel COL1A2 mutation expanding its genotypic and phenotypic spectra. Oral Diseases, 2021, 27, 1257-1267.	3.0	7
5	Phenotypic features of dentinogenesis imperfecta associated with osteogenesis imperfecta and COL1A2 mutations. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2021, 131, 694-701.	0.4	4
6	Compound Heterozygosity for a Novel Frameshift Variant Causing Fatal Infantile Liver Failure and Genotypeâ€“Phenotype Correlation of POLG c.3286C>T Variant. International Journal of Neonatal Screening, 2021, 7, 9.	3.2	1
7	Expanding the genotypic spectrum of <scp>PYCR2</scp> and a common ancestry in Thai patients with hypomyelinating leukodystrophy 10. American Journal of Medical Genetics, Part A, 2021, 185, 3068-3073.	1.2	3
8	Phenotypic and Genotypic Features of Thai Patients With Nonsyndromic Tooth Agenesis and WNT10A Variants. Frontiers in Physiology, 2020, 11, 573214.	2.8	9
9	Expanding phenotypic and mutational spectra of mitochondrial HMG-CoA synthase deficiency. European Journal of Medical Genetics, 2020, 63, 104086.	1.3	6
10	Four novel mutations of FAM20A in amelogenesis imperfecta type IG and review of literature for its genotype and phenotype spectra. Molecular Genetics and Genomics, 2020, 295, 923-931.	2.1	16
11	Decreased osteogenic activity and mineralization of alveolar bone cells from a patient with amelogenesis imperfecta and FAM83H 1261G>T mutation. Genes and Diseases, 2019, 6, 391-397.	3.4	7
12	A novel mutation in COL1A2 leads to osteogenesis imperfecta/Ehlers-Danlos overlap syndrome with brachydactyly. Genes and Diseases, 2019, 6, 138-146.	3.4	21
13	Genotypeâ€“phenotype correlation and expansion of orodental anomalies in LTBP3-related disorders. Molecular Genetics and Genomics, 2019, 294, 773-787.	2.1	24
14	Compromised alveolar bone cells in a patient with dentinogenesis imperfecta caused by DSPP mutation. Clinical Oral Investigations, 2019, 23, 303-313.	3.0	19
15	Novel compound heterozygous mutations in KREMEN1 confirm it as a disease gene for ectodermal dysplasia. British Journal of Dermatology, 2018, 179, 758-760.	1.5	10
16	A novel PITX2 mutation in nonâ€syndromic orodental anomalies. Oral Diseases, 2018, 24, 611-618.	3.0	21
17	Amelogenesis imperfecta: A novel FAM83H mutation and characteristics of periodontal ligament cells. Oral Diseases, 2018, 24, 1522-1531.	3.0	13
18	Coleâ€Carpenter syndrome in a patient from Thailand. American Journal of Medical Genetics, Part A, 2018, 176, 1706-1710.	1.2	11

#	Article	IF	Citations
19	Expanding the Oro-Dental and Mutational Spectra of Kabuki Syndrome and Expression of KMT2D and KDM6A in Human Tooth Germs. International Journal of Biological Sciences, 2018, 14, 381-389.	6.4	29