

# Kanya Suphapeetiporn

## List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

12  
papers

93  
citations

7  
h-index

9  
g-index

15  
ext. papers

153  
ext. citations

3.1  
avg, IF

2.36  
L-index

#	Paper	IF	Citations
12	Genotype-phenotype correlation and expansion of orodental anomalies in LTBP3-related disorders. <i>Molecular Genetics and Genomics</i> , <b>2019</b> , 294, 773-787	3.1	17
11	A novel GJA1 mutation in oculodentodigital dysplasia with extensive loss of enamel. <i>Oral Diseases</i> , <b>2017</b> , 23, 795-800	3.5	15
10	Dental properties, ultrastructure, and pulp cells associated with a novel DSPP mutation. <i>Oral Diseases</i> , <b>2018</b> , 24, 619-627	3.5	14
9	Monoallelic FGFR3 and Biallelic ALPL mutations in a Thai girl with hypochondroplasia and hypophosphatasia. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 2747-2752	2.5	10
8	Age-Related Reference Intervals for Blood Amino Acids in Thai Pediatric Population Measured by Liquid Chromatography Tandem Mass Spectrometry. <i>Journal of Nutrition and Metabolism</i> , <b>2018</b> , 2018, 5124035	2.7	9
7	Massive parallel sequencing as a new diagnostic approach for phenylketonuria and tetrahydrobiopterin-deficiency in Thailand. <i>BMC Medical Genetics</i> , <b>2017</b> , 18, 102	2.1	7
6	A novel PITX2 mutation in non-syndromic orodental anomalies. <i>Oral Diseases</i> , <b>2018</b> , 24, 611-618	3.5	7
5	The most 5btruncating homozygous mutation of WNT1 in siblings with osteogenesis imperfecta with a variable degree of brain anomalies: a case report. <i>BMC Medical Genetics</i> , <b>2018</b> , 19, 117	2.1	5
4	Mutational and phenotypic expansion of ATP1A3-related disorders: Report of nine cases. <i>Gene</i> , <b>2020</b> , 749, 144709	3.8	4
3	Generation of two human iPSC lines (MDCUi001-A and MDCUi001-B) from dermal fibroblasts of a Thai patient with X-linked osteogenesis imperfecta using integration-free Sendai virus. <i>Stem Cell Research</i> , <b>2019</b> , 39, 101493	1.6	2
2	The Thai reference exome (T-REx) variant database. <i>Clinical Genetics</i> , <b>2021</b> , 100, 703-712	4	2
1	Clinical and molecular characteristics of Thai patients with related neutropaenia. <i>Journal of Clinical Pathology</i> , <b>2020</b> ,	3.9	1