Kanya Suphapeetiporn

List of Publications by Year in Descending Order

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