Nanyawan Rungroj

List of Publications by Year
in descending order

8 papers	72 citations	1684188 5 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	82 citing authors

#	Article	IF	Citations
1	Molecular mechanisms of autosomal dominant and recessive distal renal tubular acidosis caused by SLC4A1 (AE1) mutations. Journal of Molecular and Genetic Medicine: an International Journal of Biomedical Research, 2005, 01, 49-62.	0.1	33
2	Association between Human Prothrombin Variant (T165M) and Kidney Stone Disease. PLoS ONE, 2012, 7, e45533.	2.5	11
3	Prothrombin Haplotype Associated With Kidney Stone Disease in Northeastern Thai Patients. Urology, 2011, 77, 249.e17-249.e23.	1.0	7
4	Loss-of-function mutations of SCN10A encoding NaV1.8 Î± subunit of voltage-gated sodium channel in patients with human kidney stone disease. Scientific Reports, 2018, 8, 10453.	3.3	7
5	Molecular Diagnosis of Solute Carrier Family 4 Member 1 (SLC4A1) Mutationâ€“Related Autosomal Recessive Distal Renal Tubular Acidosis. Laboratory Medicine, 2019, 50, 78-86.	1.2	7
6	A novel loss-of-function mutation of PBK associated with human kidney stone disease. Scientific Reports, 2020, 10, 10282.	3.3	3
7	Association between intelectin-1 variation and human kidney stone disease in northeastern Thai population. Urolithiasis, 2021, 49, 521-532.	2.0	3
8	Correlation between genotypes of F2 rs5896 (p.Thr165Met) polymorphism and urinary prothrombin fragment 1. Urolithiasis, 2018, 46, 405-407.	2.0	1