

Nanyawan Rungroj

List of Publications by Year in descending order

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Version: 2024-02-01

8
papers

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citations

1684188
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1588992
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all docs

8
docs citations

8
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82
citing authors

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