Manit Nuinoon

List of Publications by Year
in descending order

15 papers	282 citations	1307594 7 h-index	1199594 12 g-index
16 all docs	16 docs citations	16 times ranked	566 citing authors

#	Article	IF	Citations
1	Genetic predictions of life expectancy in southern Thai patients with Î²0â€‘thalassemia/Hb E. Biomedical Reports, 2022, 16, .	2.0	1
2	Promoter polymorphism of TNFâ€Î± (rs1800629) is associated with ischemic stroke susceptibility in a southern Thai population. Biomedical Reports, 2021, 15, 78.	2.0	7
3	Simultaneous Characterization of Deletional and Nondeletional Globin Gene Mutations by Multiplex Real-Time-Polymerase Chain Reaction and High-Resolution Melting Curve Analysis. Hemoglobin, 2020, 44, 311-318.	0.8	0
4	The modified G6PD deficiency screening test. Accreditation and Quality Assurance, 2020, 25, 121-126.	0.8	1
5	Compound Heterozygote for a Novel Elongated C-Terminal Î²-Globin Variant (HBB: c.364delG) and Hb E (HBB: c.79G>A) with Heterozygous Î±-Thalassemia-2. Hemoglobin, 2019, 43, 52-55.	0.8	1
6	Quantitative Trait Loci Influencing Hb F Levels in Southern Thai Hb E (HBB: c.79G>A) Heterozygotes. Hemoglobin, 2018, 42, 23-29.	0.8	7
7	Association of vitamin D receptor gene polymorphisms with serum 25(OH)D levels and metabolic syndrome in Thai population. Gene, 2018, 659, 59-66.	2.2	16
8	Hematological and Molecular Characterization of a Novel Hb A2Variant with Homozygous Î±-Thalassemia-2 in a Southern Thai Individual. Hemoglobin, 2017, 41, 213-215.	0.8	9
9	Validation of the immunochromatographic strip for Î±-thalassemia screening: a multicenter study. Translational Research, 2015, 165, 689-695.	5.0	13
10	Studies of the CETP TaqIB and ApoE Polymorphisms in Southern Thai Subjects with the Metabolic Syndrome. Biochemical Genetics, 2015, 53, 184-199.	1.7	9
11	Thalassemia and Hemoglobin E in Southern Thai Blood Donors. Advances in Hematology, 2014, 2014, 1-6.	1.0	16
12	Insight into the Peopling of Mainland Southeast Asia from Thai Population Genetic Structure. PLoS ONE, 2013, 8, e79522.	2.5	29
13	A genome-wide association identified the common genetic variants influence disease severity in Î²0-thalassemia/hemoglobin E. Human Genetics, 2010, 127, 303-314.	3.8	167
14	Detection of a known mutation M412T in the LDL receptor in a Chinese Thai FH family. Clinica Chimica Acta, 2006, 365, 211-216.	1.1	5
15	Genomic Study in Î²-Thalassemia. , 0, , .		1