Bhoom Suktitipat

List of Publications by Year
in descending order

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43 papers	1,445 citations	394421 19 h-index	345221 36 g-index
43 all docs	43 docs citations	43 times ranked	3723 citing authors

#	Article	IF	Citations
1	Genome Analyses of >200,000 Individuals Identify 58 Loci for Chronic Inflammation and Highlight Pathways that Link Inflammation and Complex Disorders. American Journal of Human Genetics, 2018, 103, 691-706.	6.2	326
2	The impact of FADS genetic variants on Ï‰6 polyunsaturated fatty acid metabolism in African Americans. BMC Genetics, 2011, 12, 50.	2.7	116
3	Identification of a specific intronic PEAR1 gene variant associated with greater platelet aggregability and protein expression. Blood, 2011, 118, 3367-3375.	1.4	95
4	Identification, Replication, and Fine-Mapping of Loci Associated with Adult Height in Individuals of African Ancestry. PLoS Genetics, 2011, 7, e1002298.	3.5	93
5	Heritability Analysis of Spherical Equivalent, Axial Length, Corneal Curvature, and Anterior Chamber Depth in the Beaver Dam Eye Study. JAMA Ophthalmology, 2009, 127, 649.	2.4	91
6	Lack of Association Between the Trp719Arg Polymorphism in Kinesin-Like Protein-6 and Coronary Artery Disease in 19 Case-Control Studies. Journal of the American College of Cardiology, 2010, 56, 1552-1563.	2.8	84
7	Genome-wide association study of platelet aggregation in African Americans. BMC Genetics, 2015, 16, 58.	2.7	50
8	A common variant in the CDKN2B gene on chromosome 9p21 protects against coronary artery disease in Americans of African ancestry. Journal of Human Genetics, 2011, 56, 224-229.	2.3	43
9	Risks and cancer associations of metachronous and synchronous multiple primary cancers: a 25-year retrospective study. BMC Cancer, 2021, 21, 1045.	2.6	43
10	HLA-DRB1 and HLA-DQB1 Are Associated with Adult-Onset Immunodeficiency with Acquired Anti-Interferon-Gamma Autoantibodies. PLoS ONE, 2015, 10, e0128481.	2.5	41
11	Protective Effects of Mangosteen Extract on H2O2-Induced Cytotoxicity in SK-N-SH Cells and Scopolamine-Induced Memory Impairment in Mice. PLoS ONE, 2013, 8, e85053.	2.5	39
12	Genetic history of Southeast Asian populations as revealed by ancient and modern human mitochondrial DNA analysis. American Journal of Physical Anthropology, 2008, 137, 425-440.	2.1	34
13	Geneâ€Gene Interaction Among WNT Genes for Oral Cleft in Trios. Genetic Epidemiology, 2015, 39, 385-394.	1.3	30
14	Factors associated with acquired Anti IFN-Î³ autoantibody in patients with nontuberculous mycobacterial infection. PLoS ONE, 2017, 12, e0176342.	2.5	30
15	The unique characteristics of Thai Leber hereditary optic neuropathy: analysis of 30 G11778A pedigrees. Journal of Human Genetics, 2006, 51, 298-304.	2.3	28
16	Targeted Deep Resequencing Identifies Coding Variants in the PEAR1 Gene That Play a Role in Platelet Aggregation. PLoS ONE, 2013, 8, e64179.	2.5	28
17	Platelet Kainate Receptor Signaling Promotes Thrombosis by Stimulating Cyclooxygenase Activation. Circulation Research, 2009, 105, 595-603.	4.5	27
18	Copy Number Variation in Thai Population. PLoS ONE, 2014, 9, e104355.	2.5	27

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19	The cyclin D1-CDK4 oncogenic interactome enables identification of potential novel oncogenes and clinical prognosis. Cell Cycle, 2014, 13, 2889-2900.	2.6	27
20	Mitochondrial Haplogroup Background May Influence Southeast Asian G11778A Leber Hereditary Optic Neuropathy. , 2011, 52, 4742.		21
21	Effects of Cytochrome P450 2C19 and Paraoxonase 1 Polymorphisms on Antiplatelet Response to Clopidogrel Therapy in Patients with Coronary Artery Disease. PLoS ONE, 2014, 9, e110188.	2.5	19
22	Mitochondrial DNA Haplogroup Distribution in Pedigrees of Southeast Asian G11778A Leber Hereditary Optic Neuropathy. Journal of Neuro-Ophthalmology, 2006, 26, 264-267.	0.8	18
23	Tumor mutational profile of triple negative breast cancer patients in Thailand revealed distinctive genetic alteration in chromatin remodeling gene. PeerJ, 2019, 7, e6501.	2.0	18
24	Oncoproteomic and gene expression analyses identify prognostic biomarkers for second primary malignancy in patients with head and neck squamous cell carcinoma. Modern Pathology, 2019, 32, 943-956.	5.5	17
25	Molecular investigation by whole exome sequencing revealed a high proportion of pathogenic variants among Thai victims of sudden unexpected death syndrome. PLoS ONE, 2017, 12, e0180056.	2.5	17
26	Proportion of 11778 Mutant Mitochondrial DNA and Clinical Expression in a Thai Population With Leber Hereditary Optic Neuropathy. Journal of Neuro-Ophthalmology, 2005, 25, 173-175.	0.8	13
27	Silencing of the Long Noncoding RNAMYCNOS1Suppresses Activity ofMYCN-Amplified Retinoblastoma WithoutRB1Mutation. , 2020, 61, 8.		9
28	MITF variants cause nonsyndromic sensorineural hearing loss with autosomal recessive inheritance. Scientific Reports, 2020, 10, 12712.	3.3	9
29	Infantile onset Sandhoff disease: clinical manifestation and a novel common mutation in Thai patients. BMC Pediatrics, 2021, 21, 22.	1.7	8
30	The Robustness of Generalized Estimating Equations for Association Tests in Extended Family Data. Human Heredity, 2012, 74, 17-26.	0.8	7
31	Clinical Effectiveness of Hyperbaric Oxygen Therapy in Complex Wounds. The Journal of the American College of Clinical Wound Specialists, 2014, 6, 9-13.	0.1	7
32	Validation of genotype imputation in Southeast Asian populations and the effect of single nucleotide polymorphism annotation on imputation outcome. BMC Medical Genetics, 2018, 19, 23.	2.1	6
33	Optimal cumulative dose of cisplatin for concurrent chemoradiotherapy among patients with non-metastatic nasopharyngeal carcinoma: a multicenter analysis in Thailand. BMC Cancer, 2020, 20, 518.	2.6	5
34	Identification of novel mutation in RANKL by wholeâ€exome sequencing in a Thai family with osteopetrosis; a case report and review of RANKL osteopetrosis. Molecular Genetics & Genomic Medicine, 2021, 9, e1727.	1.2	5
35	Regulatory effect of Phikud Navakot extract on HMG-CoA reductase and LDL-R: potential and alternate agents for lowering blood cholesterol. BMC Complementary and Alternative Medicine, 2018, 18, 258.	3.7	4
36	Successful treatment of arrhythmia with Î²â€blocker and flecainide combination in pregnant patients with Andersenâ€“Tawil syndrome: A case report and literature review. Annals of Noninvasive Electrocardiology, 2021, 26, e12798.	1.1	3

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37	Benefits of prophylactic percutaneous gastrostomy in patients with nasopharyngeal cancer receiving concurrent chemoradiotherapy: A multicenter analysis. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2022, 43, 103356.	1.3	3
38	Expression Pattern of Genes in Condyloma Acuminata Treated with Clinacanthus nutans Lindau Cream versus Podophyllin. Evidence-based Complementary and Alternative Medicine, 2021, 2021, 1-9.	1.2	1
39	Realâ€world evidence of cisplatin versus carboplatin in patients with locally advanced nasopharyngeal carcinoma receiving concurrent chemoradiotherapy: A multicenter analysis. Asia-Pacific Journal of Clinical Oncology, 2022, , .	1.1	1
40	Association of Mitochondrial DNA Polymorphisms With Pediatric-Onset Cyclic Vomiting Syndrome. Frontiers in Pediatrics, 2022, 10, .	1.9	1
41	Clinical expression and mitochondrial deoxyribonucleic acid study in twins with 14484 Leberâ€™s hereditary optic neuropathy: A case report. World Journal of Clinical Cases, 2022, 10, 6944-6953.	0.8	1
42	Genome-wide association analyses of expression phenotypes. Genetic Epidemiology, 2007, 31, S7-S11.	1.3	0
43	Antisense Oligonucleotide Induction of the hnRNPA1b Isoform Affects Pre-mRNA Splicing of SMN2 in SMA Type I Fibroblasts. International Journal of Molecular Sciences, 2022, 23, 3937.	4.1	0