Pornprot Limprasert

List of Publications by Year
in descending order

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42 papers	755 citations	623188 14 h-index	552369 26 g-index
43 all docs	43 docs citations	43 times ranked	1229 citing authors

#	Article	IF	Citations
1	Î²-Synuclein gene alterations in dementia with Lewy bodies. Neurology, 2004, 63, 805-811.	1.5	159
2	Î²-synuclein modulates Î±-synuclein neurotoxicity by reducing Î±-synuclein protein expression. Human Molecular Genetics, 2006, 15, 3002-3011.	1.4	75
3	Familial Dementia With Lewy Bodies. Archives of Neurology, 2002, 59, 1622.	4.9	64
4	HLA-B*15:21 and carbamazepine-induced Stevens-Johnson syndrome: pooled-data and in silico analysis. Scientific Reports, 2017, 7, 45553.	1.6	46
5	Analysis of CAG Repeat of the Machado-Joseph Gene in Human, Chimpanzee and Monkey Populations: A Variant Nucleotide is Associated with the Number of CAG Repeats. Human Molecular Genetics, 1996, 5, 207-213.	1.4	42
6	Novel Compound Heterozygous Mutations in the TRAPPC9 Gene in Two Siblings With Autism and Intellectual Disability. Frontiers in Genetics, 2019, 10, 61.	1.1	28
7	Familial Dementia with Lewy Bodies with an Atypical Clinical Presentation. Journal of Geriatric Psychiatry and Neurology, 2003, 16, 59-64.	1.2	27
8	De novo subtelomeric deletion of 15q associated with satellite translocation in a child with developmental delay and severe growth retardation. American Journal of Medical Genetics, Part A, 2007, 143A, 271-276.	0.7	24
9	Two Thai families with Norrie disease (ND): Association of two novel missense mutations with severe ND phenotype, seizures, and a manifesting carrier. American Journal of Medical Genetics Part A, 2001, 100, 52-55.	2.4	22
10	Chromosomal microarray analysis in a cohort of underrepresented population identifies SERINC2 as a novel candidate gene for autism spectrum disorder. Scientific Reports, 2017, 7, 12096.	1.6	21
11	A Case with a Ring Chromosome 13 in a Cohort of 203 Children with Non-Syndromic Autism and Review of the Cytogenetic Literature. Cytogenetic and Genome Research, 2014, 144, 1-8.	0.6	20
12	Significant Changes in Plasma Alpha-Synuclein and Beta-Synuclein Levels in Male Children with Autism Spectrum Disorder. BioMed Research International, 2018, 2018, 1-7.	0.9	20
13	Mutation screening of the CDKL5 gene in cryptogenic infantile intractable epilepsy and review of clinical sensitivity. European Journal of Paediatric Neurology, 2011, 15, 432-438.	0.7	15
14	A New Method for FMR1 Gene Methylation Screening by Multiplex Methylation-Specific Real-Time Polymerase Chain Reaction. Genetic Testing and Molecular Biomarkers, 2011, 15, 387-393.	0.3	14
15	Mutation Screening of the Neurexin 1 Gene in Thai Patients with Intellectual Disability and Autism Spectrum Disorder. Genetic Testing and Molecular Biomarkers, 2014, 18, 510-515.	0.3	14
16	A genome wide pattern of population structure and admixture in peninsular Malaysia Malays. The HUGO Journal, 2014, 8, 5.	4.1	14
17	Mutation screening of the Aristaless-related homeobox (ARX) gene in Thai pediatric patients with delayed development: First report from Thailand. European Journal of Medical Genetics, 2007, 50, 346-354.	0.7	13
18	Whole-exome sequencing identifies a novel heterozygous missense variant of the EN2 gene in two unrelated patients with autism spectrum disorder. Psychiatric Genetics, 2016, 26, 297-301.	0.6	13

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19	Prenatal diagnosis of deletion of chromosome 6p presenting with hydrops fetalis. Prenatal Diagnosis, 2004, 24, 887-889.	1.1	12
20	Comparative studies of the CAG repeats in the spinocerebellar ataxia type 1 (SCA1) gene. , 1997, 74, 488-493.		11
21	Haplotype analysis at the FRAXA locus in Thai subjects. American Journal of Medical Genetics Part A, 2001, 98, 224-229.	2.4	11
22	Exome Sequencing Identifies Compound Heterozygous Mutations inSCN5AAssociated with Congenital Complete Heart Block in the Thai Population. Disease Markers, 2016, 2016, 1-10.	0.6	10
23	Common Clinical Characteristics and Rare Medical Problems of Fragile X Syndrome in Thai Patients and Review of the Literature. International Journal of Pediatrics (United Kingdom), 2017, 2017, 1-11.	0.2	10
24	Polymorphism ofFXR1 showing lack of association with autism. American Journal of Medical Genetics Part A, 1997, 74, 453-454.	2.4	7
25	Screening of NLGN3 and NLGN4X genes in Thai children with autism spectrum disorder. Psychiatric Genetics, 2014, 24, 42-43.	0.6	7
26	Comparison of a New In-House and Three Published HLA-B*15:02 Screening Methods for Prevention of Carbamazepine-Induced Severe Drug Reactions. PLoS ONE, 2016, 11, e0155907.	1.1	7
27	Screening for Subtelomeric Rearrangements in Thai Patients with Intellectual Disabilities Using FISH and Review of Literature on Subtelomeric FISH in 15,591 Cases with Intellectual Disabilities. Genetics Research International, 2016, 2016, 1-13.	2.0	6
28	Unilateral macroorchidism in fragile X syndrome. American Journal of Medical Genetics Part A, 2000, 95, 516-517.	2.4	5
29	No association of Val158Met variant in the COMT gene with autism spectrum disorder in Thai children. Psychiatric Genetics, 2014, 24, 230-231.	0.6	5
30	Unique AGG Interruption in the CGG Repeats of the FMR1 Gene Exclusively Found in Asians Linked to a Specific SNP Haplotype. Genetics Research International, 2016, 2016, 1-7.	2.0	5
31	A new structured interview for children with autism spectrum disorder based on the DSM-IV. Journal of the Medical Association of Thailand = Chotmaihet Thangphaet, 2014, 97 Suppl 8, S7-14.	0.4	5
32	Association of ABO Blood Group Phenotype and Allele Frequency with Chikungunya Fever. Advances in Hematology, 2015, 2015, 1-4.	0.6	4
33	Multiplex methylation specific PCR analysis of fragile X syndrome: experience in Songklanagarind Hospital. Journal of the Medical Association of Thailand = Chotmaihet Thangphaet, 2005, 88, 1057-61.	0.4	4
34	Possible founder effects for FRAXE alleles. , 1999, 84, 286-290.		3
35	Whole-Exome Sequencing Identifies One De Novo Variant in theFGD6Gene in a Thai Family with Autism Spectrum Disorder. International Journal of Genomics, 2018, 2018, 1-7.	0.8	3
36	Clinical abnormalities, intervention program, and school attendance of Down syndrome children in southern Thailand. Journal of the Medical Association of Thailand = Chotmaihet Thangphaet, 2004, 87, 1199-204.	0.4	3

#	Article	IF	Citations
37	Significant associations between 5-hydroxytryptaminetransporter-linked promoter region polymorphisms of the serotonin transporter (solute carrier family 6 member 4) gene and Thai patients with autism spectrum disorder. Medicine (United States), 2020, 99, e21946.	0.4	2
38	Maternal Age-Specific Rates for Trisomy 21 and Common Autosomal Trisomies in Fetuses from a Single Diagnostic Center in Thailand. PLoS ONE, 2016, 11, e0165859.	1.1	2
39	Screening for FMR1 CGG Repeat Expansion in Thai Patients with Autism Spectrum Disorder. BioMed Research International, 2021, 2021, 1-11.	0.9	1
40	Case Report: An Atypical Angelman Syndrome Case With Obesity and Fulfilled Autism Spectrum Disorder Identified by Microarray. Frontiers in Genetics, 2021, 12, 755605.	1.1	0
41	AB051. Chromosomal microarray analysis in a large cohort of Thai patients with autism spectrum disorder. Annals of Translational Medicine, 2017, 5, AB051-AB051.	0.7	0
42	No Evidence of Abnormal Expression of Beta-Catenin and Bcl-2 Proteins in Pilomatricoma as One Clinical Feature of Tetrasomy 9p Syndrome. International Journal of Pediatrics (United Kingdom), 2021, 2021, 1-8.	0.2	0