

Pornprot Limprasert

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/10808/publications.pdf>

Version: 2024-02-01

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	Case Report: An Atypical Angelman Syndrome Case With Obesity and Fulfilled Autism Spectrum Disorder Identified by Microarray. <i>Frontiers in Genetics</i> , 2021, 12, 755605.	1.1	0
2	No Evidence of Abnormal Expression of Beta-Catenin and Bcl-2 Proteins in Pilomatricoma as One Clinical Feature of Tetrasomy 9p Syndrome. <i>International Journal of Pediatrics (United Kingdom)</i> , 2021, 2021, 1-8.	0.2	0
3	Screening for FMR1 CGG Repeat Expansion in Thai Patients with Autism Spectrum Disorder. <i>BioMed Research International</i> , 2021, 2021, 1-11.	0.9	1
4	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. <i>Medicine (United States)</i> , 2020, 99, e21946.	0.4	2
5	Novel Compound Heterozygous Mutations in the TRAPPC9 Gene in Two Siblings With Autism and Intellectual Disability. <i>Frontiers in Genetics</i> , 2019, 10, 61.	1.1	28
6	Whole-Exome Sequencing Identifies One De Novo Variant in the <i>FGD6</i> Gene in a Thai Family with Autism Spectrum Disorder. <i>International Journal of Genomics</i> , 2018, 2018, 1-7.	0.8	3
7	Significant Changes in Plasma Alpha-Synuclein and Beta-Synuclein Levels in Male Children with Autism Spectrum Disorder. <i>BioMed Research International</i> , 2018, 2018, 1-7.	0.9	20
8	HLA-B*15:21 and carbamazepine-induced Stevens-Johnson syndrome: pooled-data and in silico analysis. <i>Scientific Reports</i> , 2017, 7, 45553.	1.6	46
9	Chromosomal microarray analysis in a cohort of underrepresented population identifies <i>SERINC2</i> as a novel candidate gene for autism spectrum disorder. <i>Scientific Reports</i> , 2017, 7, 12096.	1.6	21
10	Common Clinical Characteristics and Rare Medical Problems of Fragile X Syndrome in Thai Patients and Review of the Literature. <i>International Journal of Pediatrics (United Kingdom)</i> , 2017, 2017, 1-11.	0.2	10
11	AB051. Chromosomal microarray analysis in a large cohort of Thai patients with autism spectrum disorder. <i>Annals of Translational Medicine</i> , 2017, 5, AB051-AB051.	0.7	0
12	Unique AGG Interruption in the CGG Repeats of the <i>FMR1</i> Gene Exclusively Found in Asians Linked to a Specific SNP Haplotype. <i>Genetics Research International</i> , 2016, 2016, 1-7.	2.0	5
13	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. <i>Genetics Research International</i> , 2016, 2016, 1-13.	2.0	6
14	Exome Sequencing Identifies Compound Heterozygous Mutations in <i>SCN5A</i> Associated with Congenital Complete Heart Block in the Thai Population. <i>Disease Markers</i> , 2016, 2016, 1-10.	0.6	10
15	Whole-exome sequencing identifies a novel heterozygous missense variant of the <i>EN2</i> gene in two unrelated patients with autism spectrum disorder. <i>Psychiatric Genetics</i> , 2016, 26, 297-301.	0.6	13
16	Comparison of a New In-House and Three Published HLA-B*15:02 Screening Methods for Prevention of Carbamazepine-Induced Severe Drug Reactions. <i>PLoS ONE</i> , 2016, 11, e0155907.	1.1	7
17	Maternal Age-Specific Rates for Trisomy 21 and Common Autosomal Trisomies in Fetuses from a Single Diagnostic Center in Thailand. <i>PLoS ONE</i> , 2016, 11, e0165859.	1.1	2
18	Association of ABO Blood Group Phenotype and Allele Frequency with Chikungunya Fever. <i>Advances in Hematology</i> , 2015, 2015, 1-4.	0.6	4

#	ARTICLE	IF	CITATIONS
19	Mutation Screening of the Neurexin 1 Gene in Thai Patients with Intellectual Disability and Autism Spectrum Disorder. <i>Genetic Testing and Molecular Biomarkers</i> , 2014, 18, 510-515.	0.3	14
20	A Case with a Ring Chromosome 13 in a Cohort of 203 Children with Non-Syndromic Autism and Review of the Cytogenetic Literature. <i>Cytogenetic and Genome Research</i> , 2014, 144, 1-8.	0.6	20
21	A genome wide pattern of population structure and admixture in peninsular Malaysia Malays. <i>The HUGO Journal</i> , 2014, 8, 5.	4.1	14
22	No association of Val158Met variant in the COMT gene with autism spectrum disorder in Thai children. <i>Psychiatric Genetics</i> , 2014, 24, 230-231.	0.6	5
23	Screening of NLGN3 and NLGN4X genes in Thai children with autism spectrum disorder. <i>Psychiatric Genetics</i> , 2014, 24, 42-43.	0.6	7
24	A new structured interview for children with autism spectrum disorder based on the DSM-IV. <i>Journal of the Medical Association of Thailand = Chotmaihet Thangphaet</i> , 2014, 97 Suppl 8, S7-14.	0.4	5
25	A New Method for FMR1 Gene Methylation Screening by Multiplex Methylation-Specific Real-Time Polymerase Chain Reaction. <i>Genetic Testing and Molecular Biomarkers</i> , 2011, 15, 387-393.	0.3	14
26	Mutation screening of the CDKL5 gene in cryptogenic infantile intractable epilepsy and review of clinical sensitivity. <i>European Journal of Paediatric Neurology</i> , 2011, 15, 432-438.	0.7	15
27	Mutation screening of the Aristaless-related homeobox (ARX) gene in Thai pediatric patients with delayed development: First report from Thailand. <i>European Journal of Medical Genetics</i> , 2007, 50, 346-354.	0.7	13
28	De novo subtelomeric deletion of 15q associated with satellite translocation in a child with developmental delay and severe growth retardation. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 271-276.	0.7	24
29	$\hat{\tau}^2$ -synuclein modulates $\hat{\tau}^1$ -synuclein neurotoxicity by reducing $\hat{\tau}^1$ -synuclein protein expression. <i>Human Molecular Genetics</i> , 2006, 15, 3002-3011.	1.4	75
30	Multiplex methylation specific PCR analysis of fragile X syndrome: experience in Songklanagarind Hospital. <i>Journal of the Medical Association of Thailand = Chotmaihet Thangphaet</i> , 2005, 88, 1057-61.	0.4	4
31	$\hat{\tau}^2$ -Synuclein gene alterations in dementia with Lewy bodies. <i>Neurology</i> , 2004, 63, 805-811.	1.5	159
32	Prenatal diagnosis of deletion of chromosome 6p presenting with hydrops fetalis. <i>Prenatal Diagnosis</i> , 2004, 24, 887-889.	1.1	12
33	Clinical abnormalities, intervention program, and school attendance of Down syndrome children in southern Thailand. <i>Journal of the Medical Association of Thailand = Chotmaihet Thangphaet</i> , 2004, 87, 1199-204.	0.4	3
34	Familial Dementia with Lewy Bodies with an Atypical Clinical Presentation. <i>Journal of Geriatric Psychiatry and Neurology</i> , 2003, 16, 59-64.	1.2	27
35	Familial Dementia With Lewy Bodies. <i>Archives of Neurology</i> , 2002, 59, 1622.	4.9	64
36	Haplotype analysis at the FRAXA locus in Thai subjects. <i>American Journal of Medical Genetics Part A</i> , 2001, 98, 224-229.	2.4	11

#	ARTICLE	IF	CITATIONS
37	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
38	Unilateral macroorchidism in fragile X syndrome. American Journal of Medical Genetics Part A, 2000, 95, 516-517.	2.4	5
39	Possible founder effects for FRAXE alleles. , 1999, 84, 286-290.		3
40	Polymorphism of FXR1 showing lack of association with autism. American Journal of Medical Genetics Part A, 1997, 74, 453-454.	2.4	7
41	Comparative studies of the CAG repeats in the spinocerebellar ataxia type 1 (SCA1) gene. , 1997, 74, 488-493.		11
42	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