

Chariyawan Charalsawadi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4235085/publications.pdf>

Version: 2024-02-01

10
papers

126
citations

1937685

4
h-index

1588992

8
g-index

10
all docs

10
docs citations

10
times ranked

217
citing authors

#	ARTICLE	IF	CITATIONS
1	Possible association between a polymorphism of EPAS1 gene and persistent pulmonary hypertension of the newborn: a case-control study. <i>Jornal De Pediatria</i> , 2022, 98, 383-389.	2.0	2
2	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.	2.3	0
3	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.	0.8	0
4	Clinical delineation of 18q11-q12 microdeletion: Intellectual disability, speech and behavioral disorders, and conotruncal heart defects. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e896.	1.2	5
5	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.	0.8	10
6	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
7	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.	2.5	2
8	Mosaicism for trisomy 21: A review. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 26-39.	1.2	77
9	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.	1.1	20
10	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.	0.1	4