

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	Mosaicism for trisomy 21: A review. American Journal of Medical Genetics, Part A, 2015, 167, 26-39.	1.2	77
2	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.	1.1	20
3	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.	0.8	10
4	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
5	Clinical delineation of 18q11-q12 microdeletion: Intellectual disability, speech and behavioral disorders, and conotruncal heart defects. Molecular Genetics & Genomic Medicine, 2019, 7, e896.	1.2	5
6	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.	0.1	4
7	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.	2.5	2
8	Possible association between a polymorphism of EPAS1 gene and persistent pulmonary hypertension of the newborn: a case-control study. Jornal De Pediatria, 2022, 98, 383-389.	2.0	2
9	Case Report: An Atypical Angelman Syndrome Case With Obesity and Fulfilled Autism Spectrum Disorder Identified by Microarray. Frontiers in Genetics, 2021, 12, 755605.	2.3	0
10	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.	0.8	0