

Thinh Huy Tran

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/12064024/publications.pdf>

Version: 2024-02-01

17
papers

221
citations

1478505

6
h-index

996975

15
g-index

17
all docs

17
docs citations

17
times ranked

397
citing authors

#	ARTICLE	IF	CITATIONS
1	B cell-specific and stimulation-responsive enhancers derepress Aicda by overcoming the effects of silencers. <i>Nature Immunology</i> , 2010, 11, 148-154.	14.5	111
2	Preventing AID, a physiological mutator, from deleterious activation: regulation of the genomic instability that is associated with antibody diversity. <i>International Immunology</i> , 2010, 22, 227-235.	4.0	41
3	Validation of ambiguous MLPA results by targeted next-generation sequencing discloses a nonsense mutation in the DMD gene. <i>Clinica Chimica Acta</i> , 2014, 436, 155-159.	1.1	16
4	Exon Deletion Patterns of the Dystrophin Gene in 82 Vietnamese Duchenne/Becker Muscular Dystrophy Patients. <i>Journal of Neurogenetics</i> , 2013, 27, 170-175.	1.4	11
5	Rapid method for targeted prenatal diagnosis of Duchenne muscular dystrophy in Vietnam. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2013, 52, 534-539.	1.3	6
6	Acromesomelic dysplasia Maroteaux type in patients from Vietnam. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1420-1422.	1.2	6
7	Association of the <i>STAT4</i> , <i>CDKN1A</i> , and <i>IRF5</i> variants with risk of lupus nephritis and renal biopsy classification in patients in Vietnam. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1648.	1.2	6
8	Mosaicism in carrier of Duchenne muscular dystrophy mutation—Implication for prenatal diagnosis. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2018, 57, 878-880.	1.3	4
9	A case of self-improving collodion ichthyosis in Vietnam. <i>Pediatric Dermatology</i> , 2020, 37, 574-575.	0.9	4
10	Prenatal diagnosis of a case with SEA-HPFH deletion thalassemia with whole HBB gene deletion. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2018, 57, 435-441.	1.3	3
11	Assessment of 6 STR loci for prenatal diagnosis of Duchenne Muscular Dystrophy. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2019, 58, 645-649.	1.3	3
12	Microcephaly primary hereditary (MCPH): Report of novel ASPM variants and prenatal diagnosis in a Vietnamese family. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2021, 60, 907-910.	1.3	3
13	Variation of Mitochondrial DNA HV1 AND HV2 of the Vietnamese Population. <i>Advances in Experimental Medicine and Biology</i> , 2018, 1292, 37-63.	1.6	2
14	Mutation characteristic of 103 haemophilia A patients in Vietnam: Identification of novel mutations. <i>Haemophilia</i> , 2019, 25, e274-e277.	2.1	2
15	Whole exome sequencing analysis in a couple with three children who died prematurely due to carnitine-acylcarnitine translocase deficiency. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2022, 61, 153-156.	1.3	2
16	Molecular Characterization and Genotype-Phenotype Correlation of G6PD Mutations in Five Ethnicities of Northern Vietnam. <i>Anemia</i> , 2022, 2022, 1-10.	1.7	1
17	A Novel Nonsense Mutation c.374C>G in CYP21A2 Gene of a Vietnamese Patient with Congenital Adrenal Hyperplasia. <i>Advances in Experimental Medicine and Biology</i> , 2018, 1292, 27-35.	1.6	0