Thinh Huy Tran

List of Publications by Year
in descending order

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17	221	1478505 6	996975 15
papers	citations	h-index	g-index

17	17	17	397
all docs	docs citations	times ranked	citing authors

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