

Van Khanh Tran

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

223
citations

1163117

8
h-index

1058476

14
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25
all docs

25
docs citations

25
times ranked

364
citing authors

#	ARTICLE	IF	CITATIONS
1	Splicing analysis disclosed a determinant single nucleotide for exon skipping caused by a novel intraexonic four-nucleotide deletion in the dystrophin gene. <i>Journal of Medical Genetics</i> , 2006, 43, 924-930.	3.2	34
2	Co-occurrence of mutations in both dystrophin- and androgen-receptor genes is a novel cause of female Duchenne muscular dystrophy. <i>Human Genetics</i> , 2006, 119, 516-519.	3.8	28
3	SMN2 and NAIP gene dosages in Vietnamese patients with spinal muscular atrophy. <i>Pediatrics International</i> , 2008, 50, 346-351.	0.5	27
4	A novel cryptic exon identified in the 3' region of intron 2 of the human dystrophin gene. <i>Journal of Human Genetics</i> , 2005, 50, 425-433.	2.3	21
5	A nonsense mutation-created intraexonic splice site is active in the lymphocytes, but not in the skeletal muscle of a DMD patient. <i>Human Genetics</i> , 2006, 120, 737-742.	3.8	19
6	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
7	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
8	Novel variants of <i>CYP21A2</i> in Vietnamese patients with congenital adrenal hyperplasia. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e623.	1.2	8
9	Cryptic splice activation but not exon skipping is observed in minigene assays of dystrophin c.9361+1G>A mutation identified by NGS. <i>Journal of Human Genetics</i> , 2017, 62, 531-537.	2.3	7
10	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
11	Acromesomelic dysplasia Maroteaux type in patients from Vietnam. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1420-1422.	1.2	6
12	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
13	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
14	A case of self-improving collodion ichthyosis in Vietnam. <i>Pediatric Dermatology</i> , 2020, 37, 574-575.	0.9	4
15	Dentinogenesis Imperfecta and Caries in Osteogenesis Imperfecta among Vietnamese Children. <i>Dentistry Journal</i> , 2021, 9, 49.	2.3	4
16	Genetic analysis of 55 northern Vietnamese patients with Wilson disease: seven novel mutations in ATP7B. <i>Journal of Genetics</i> , 2017, 96, 933-939.	0.7	3
17	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
18	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

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19	Primary congenital glaucoma in Vietnam: analysis and identification of novel CYP1B1 variants. <i>Ophthalmic Genetics</i> , 2019, 40, 286-287.	1.2	3
20	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
21	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
22	Mutation characteristic of 103 haemophilia A patients in Vietnam: Identification of novel mutations. <i>Haemophilia</i> , 2019, 25, e274-e277.	2.1	2
23	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
24	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
25	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