Xianqin Zhang

List of Publications by Year
in descending order

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22	907	933447 10	794594 19
papers	citations	h-index	g-index

22	22	22	1921
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Two novel STAT1 mutations cause Mendelian susceptibility to mycobacterial disease. Biochemical and Biophysical Research Communications, 2022, 591, 124-129.	2.1	5
2	Novel mutations in ZP2 and ZP3 cause female infertility in three patients. Journal of Assisted Reproduction and Genetics, 2022, 39, 1205-1215.	2.5	3
3	Novel Heterozygous Mutations in ZP2 Cause Abnormal Zona Pellucida and Female Infertility. Reproductive Sciences, 2022, 29, 3047-3054.	2.5	5
4	A Novel Homozygous Nonsense Mutation in ZP1 Causes Female Infertility due to Empty Follicle Syndrome. Reproductive Sciences, 2022, 29, 3516-3520.	2.5	2
5	A novel mutation in ZP3 causes empty follicle syndrome and abnormal zona pellucida formation. Journal of Assisted Reproduction and Genetics, 2021, 38, 251-259.	2.5	20
6	Tsga10 is essential for arrangement of mitochondrial sheath and male fertility in mice. Andrology, 2021, 9, 368-375.	3.5	16
7	TUBB8 Mutations Cause Female Infertility with Large Polar Body Oocyte and Fertilization Failure. Reproductive Sciences, 2021, 28, 2942-2950.	2.5	13
8	A novel microdeletion upstream of HOXD13 in a Chinese family with synpolydactyly. American Journal of Medical Genetics, Part A, 2021, , .	1.2	1
9	Novel mutations in ZP1 and ZP2 cause primary infertility due to empty follicle syndrome and abnormal zona pellucida. Journal of Assisted Reproduction and Genetics, 2020, 37, 2853-2860.	2.5	21
10	p.His16Arg of STXBP1 (MUNC18-1) Associated With Syntaxin 3B Causes Autosomal Dominant Congenital Nystagmus. Frontiers in Cell and Developmental Biology, 2020, 8, 591781.	3.7	6
11	A novel UBE2A mutation in a Chinese family with Xâ€linked intellectual disability. Journal of Gene Medicine, 2020, 22, e3191.	2.8	4
12	Novel homozygous mutations in PATL2 lead to female infertility with oocyte maturation arrest. Journal of Assisted Reproduction and Genetics, 2020, 37, 841-847.	2.5	22
13	Deficiency of SCAMP5 leads to pediatric epilepsy and dysregulation of neurotransmitter release in the brain. Human Genetics, 2020, 139, 545-555.	3.8	16
14	Novel compound heterozygous mutations in WEE2 causes female infertility and fertilization failure. Journal of Assisted Reproduction and Genetics, 2019, 36, 1957-1962.	2.5	22
15	Three novel compound heterozygous IL12RB1 mutations in Chinese patients with Mendelian susceptibility to mycobacterial disease. PLoS ONE, 2019, 14, e0215648.	2.5	7
16	A novel MIP mutation in a Chinese family with congenital cataract. Ophthalmic Genetics, 2018, 39, 473-476.	1.2	3
17	Human intracellular ISG15 prevents interferon-Î±/Î² over-amplification and auto-inflammation. Nature, 2015, 517, 89-93.	27.8	432
18	Protective effect of KCNH2 single nucleotide polymorphism K897T in LQTS families and identification of novel KCNQ1 and KCNH2mutations. BMC Medical Genetics, 2008, 9, 87.	2.1	23

#	ARTICLE	IF	CITATIONS
19	Mutation in Nuclear Pore Component NUP155 Leads to Atrial Fibrillation and Early Sudden Cardiac Death. Cell, 2008, 135, 1017-1027.	28.9	249
20	A novel DSPPmutation is associated with type II dentinogenesis Imperfecta in a chinese family. BMC Medical Genetics, 2007, 8, 52.	2.1	33
21	Juvenile-Onset Kufs Disease in a Chinese Consanguineous Family due to CLN6 Mutation. Neurodegenerative Diseases, 0, , .	1.4	0
22	Novel Compound Heterozygous Mutation in FSIP2 Causes Multiple Morphological Abnormalities of the Sperm Flagella (MMAF) and Male Infertility. Reproductive Sciences, 0, , .	2.5	4