Xiaoli Wei

List of Publications by Year
in descending order

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10 papers	137 citations	1478505 6 h-index	1372567 10 g-index
10 all docs	10 docs citations	10 times ranked	209 citing authors

#	Article	IF	Citations
1	Biallelic mutations of CFAP74 may cause human primary ciliary dyskinesia and MMAF phenotype. Journal of Human Genetics, 2020, 65, 961-969.	2.3	36
2	Lossâ€ofâ€function mutations in centrosomal protein 112 is associated with human acephalic spermatozoa phenotype. Clinical Genetics, 2020, 97, 321-328.	2.0	28
3	Biallelic mutations in Sperm flagellum 2 cause human multiple morphological abnormalities of the sperm flagella (MMAF) phenotype. Clinical Genetics, 2019, 96, 385-393.	2.0	24
4	Mutations in ZP4 are associated with abnormal zona pellucida and female infertility. Journal of Clinical Pathology, 2022, 75, 201-204.	2.0	13
5	Bi-allelic mutations in DNAH7 cause asthenozoospermia by impairing the integrality of axoneme structure. Acta Biochimica Et Biophysica Sinica, 2021, 53, 1300-1309.	2.0	13
6	Biallelic mutations in <scp>KATNAL2</scp> cause male infertility due to oligoâ€asthenoâ€teratozoospermia. Clinical Genetics, 2021, 100, 376-385.	2.0	10
7	Pathogenic Variants in ACTRT1 Cause Acephalic Spermatozoa Syndrome. Frontiers in Cell and Developmental Biology, 2021, 9, 676246.	3.7	5
8	Pathogenic variants of ATG4D in infertile men with nonâ€obstructive azoospermia identified using wholeâ€exome sequencing. Clinical Genetics, 2021, 100, 280-291.	2.0	4
9	Impact of STAT4 gene silencing on the expression profile of proteins in EL-4 cells. Science Bulletin, 2009, 54, 3265-3270.	1.7	2
10	Biallelic mutations inARMC12cause asthenozoospermia and multiple midpiece defects in humans and mice. Journal of Medical Genetics, 2023, 60, 154-162.	3.2	2