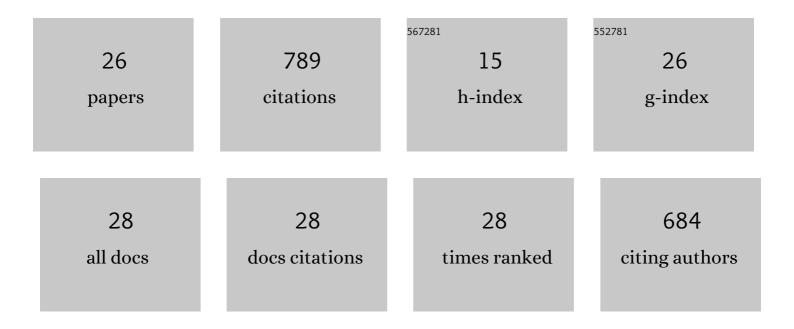
Tu Chao-Feng

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Biallelic <i>CFAP61</i> variants cause male infertility in humans and mice with severe oligoasthenoteratozoospermia. Journal of Medical Genetics, 2023, 60, 144-153.	3.2	12
2	Meiotic recombination: insights into its mechanisms and its role in human reproduction with a special focus on non-obstructive azoospermia. Human Reproduction Update, 2022, 28, 763-797.	10.8	27
3	Bi-allelic variants in <i>SHOC1</i> cause non-obstructive azoospermia with meiosis arrest in humans and mice. Molecular Human Reproduction, 2022, 28, .	2.8	7
4	Deleterious variants in X-linked CFAP47 induce asthenoteratozoospermia and primary male infertility. American Journal of Human Genetics, 2021, 108, 309-323.	6.2	74
5	TDRD7 participates in lens development and spermiogenesis by mediating autophagosome maturation. Autophagy, 2021, 17, 3848-3864.	9.1	19
6	CFAP65 is required in the acrosome biogenesis and mitochondrial sheath assembly during spermiogenesis. Human Molecular Genetics, 2021, 30, 2240-2254.	2.9	18
7	Loss-of-function missense variant of <i>AKAP4</i> induced male infertility through reduced interaction with QRICH2 during sperm flagella development. Human Molecular Genetics, 2021, 31, 219-231.	2.9	19
8	Novel variants of the PCCB gene in Chinese patients with propionic acidemia. Clinica Chimica Acta, 2021, 519, 18-25.	1.1	1
9	Bi-allelic mutations of DNAH10 cause primary male infertility with asthenoteratozoospermia in humans and mice. American Journal of Human Genetics, 2021, 108, 1466-1477.	6.2	50
10	Genetic underpinnings of asthenozoospermia. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101472.	4.7	37
11	Novel DNAAF6 variants identified by whole-exome sequencing cause male infertility and primary ciliary dyskinesia. Journal of Assisted Reproduction and Genetics, 2020, 37, 811-820.	2.5	9
12	A homozygous RPL10L missense mutation associated with maleÂfactorÂinfertility and severe oligozoospermia. Fertility and Sterility, 2020, 113, 561-568.	1.0	10
13	Novel mutations in SPEF2 causing different defects between flagella and cilia bridge: the phenotypic link between MMAF and PCD. Human Genetics, 2020, 139, 257-271.	3.8	46
14	An <i>M1AP</i> homozygous spliceâ€site mutation associated with severe oligozoospermia in a consanguineous family. Clinical Genetics, 2020, 97, 741-746.	2.0	15
15	Novel biallelic PCNT deletion causing microcephalic osteodysplastic primordial dwarfism type II with congenital heart defect. Science China Life Sciences, 2019, 62, 144-147.	4.9	4
16	<i>GPC6</i> Promotes Cell Proliferation, Migration, and Invasion in Nasopharyngeal Carcinoma. Journal of Cancer, 2019, 10, 3926-3932.	2.5	34
17	Telomerase insufficiency induced telomere erosion accumulation in successive generations in dyskeratosis congenita family. Molecular Genetics & amp; Genomic Medicine, 2019, 7, e00709.	1.2	7
18	Identification of DNAH6 mutations in infertile men with multiple morphological abnormalities of the sperm flagella. Scientific Reports, 2019, 9, 15864.	3.3	42

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19	<i>XRCC2</i> mutation causes premature ovarian insufficiency as well as nonâ€obstructive azoospermia in humans. Clinical Genetics, 2019, 95, 442-443.	2.0	35
20	Loss-of-function mutations in TDRD7 lead to a rare novel syndrome combining congenital cataract and nonobstructive azoospermia in humans. Genetics in Medicine, 2019, 21, 1209-1217.	2.4	70
21	Biallelic mutations in <i>CFAP65</i> lead to severe asthenoteratospermia due to acrosome hypoplasia and flagellum malformations. Journal of Medical Genetics, 2019, 56, 750-757.	3.2	61
22	Rare partial octosomy and hexasomy of 15q11-q13 associated with intellectual impairment and development delay: report of two cases and review of literature. Molecular Cytogenetics, 2018, 11, 15.	0.9	5
23	Identification of genomic alterations in nasopharyngeal carcinoma and nasopharyngeal carcinoma-derived Epstein–Barr virus by whole-genome sequencing. Carcinogenesis, 2018, 39, 1517-1528.	2.8	74
24	CASC2c as an unfavorable prognosis factor interacts with miR-101 to mediate astrocytoma tumorigenesis. Cell Death and Disease, 2017, 8, e2639-e2639.	6.3	30
25	Genome-Wide Analysis of 18 Epstein-Barr Viruses Isolated from Primary Nasopharyngeal Carcinoma Biopsy Specimens. Journal of Virology, 2017, 91, .	3.4	70
26	Genotype-phenotype correlation and identification of two novel SRD5A2 mutations in 33 Chinese patients with hypospadias. Steroids, 2017, 125, 61-66.	1.8	13