Shao-Hua Tang

List of Publications by Year
in descending order

8	74	1937685 4	1588992 8
papers	citations	h-index	g-index

11	11	11	128
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A pregnancy with discordant fetal and placental chromosome 18 aneuploidies revealed by invasive and noninvasive prenatal diagnosis. Reproductive BioMedicine Online, 2014, 29, 136-139.	2.4	18
2	Mutation analysis of common deafness-causing genes among 506 patients with nonsyndromic hearing loss from Wenzhou city, China. International Journal of Pediatric Otorhinolaryngology, 2019, 122, 185-190.	1.0	16
3	Skeleton Genetics: a comprehensive database for genes and mutations related to genetic skeletal disorders. Database: the Journal of Biological Databases and Curation, 2016, 2016, baw127.	3.0	12
4	Nextâ€generation sequencing identifies rare pathogenic and novel candidate variants in a cohort of Chinese patients with syndromic or nonsyndromic hearing loss. Molecular Genetics & Genomic Medicine, 2020, 8, e1539.	1.2	11
5	Novel mutation in <scp>MBTPS</scp>2 causes keratosis follicularis spinulosa decalvans in a large Chinese family. International Journal of Dermatology, 2019, 58, 493-496.	1.0	7
6	The variations in human orphan G proteinâ€coupled receptor QRFPR affect PI3Kâ€AKTâ€mTOR signaling. Journal of Clinical Laboratory Analysis, 2021, 35, e23822.	2.1	6
7	Genetic analysis and prenatal diagnosis of 20 Chinese families with oculocutaneous albinism. Journal of Clinical Laboratory Analysis, 2021, 35, e23647.	2.1	3
8	The distal arthrogryposisâ€linked p.R63C variant promotes the stability and nuclear accumulation of TNNT3. Journal of Clinical Laboratory Analysis, 2021, 35, e24089.	2.1	1