

Shunyao Liao

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Whole-exome sequencing of de novo genetic variants in a Chinese family with a sporadic case of congenital nonsyndromic hearing loss. <i>F1000Research</i> , 2021, 10, 61.	1.6	1
2	Whole-exome sequencing of de novo genetic variants in a Chinese family with a sporadic case of congenital nonsyndromic hearing loss. <i>F1000Research</i> , 2021, 10, 61.	1.6	2
3	Heterogeneous impact of type 2 diabetes mellitus-related genetic variants on gestational glycemic traits: review and future research needs. <i>Molecular Genetics and Genomics</i> , 2019, 294, 811-847.	2.1	7
4	DNA demethylation facilitates the specific transcription of the mouse X-linked <i>Tsga8</i> gene in round spermatids. <i>Biology of Reproduction</i> , 2019, 100, 994-1007.	2.7	2
5	Novel compound heterozygous mutations of <i>ALDH1A3</i> contribute to anophthalmia in a non-consanguineous Chinese family. <i>Genetics and Molecular Biology</i> , 2017, 40, 430-435.	1.3	9
6	Epigenetic modifications promote the expression of the orphan nuclear receptor <i>NROB1</i> in human lung adenocarcinoma cells. <i>Oncotarget</i> , 2016, 7, 43162-43176.	1.8	7
7	Association of variants in <i>CDKN2A/2B</i> and <i>CDKAL1</i> genes with gestational insulin sensitivity and disposition in pregnant Han Chinese women. <i>Journal of Diabetes Investigation</i> , 2015, 6, 295-301.	2.4	10
8	The Impact of Genetic Variants for Different Physiological Characterization of Type 2 Diabetes Loci on Gestational Insulin Signaling in Nondiabetic Pregnant Chinese Women. <i>Reproductive Sciences</i> , 2015, 22, 1421-1428.	2.5	9