Qing Fang

List of Publications by Year
in descending order

11	324	1163117 8	1372567 10
papers	citations	h-index	g-index

12	12	12	615
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetics of Combined Pituitary Hormone Deficiency: Roadmap into the Genome Era. Endocrine Reviews, 2016, 37, 636-675.	20.1	147
2	The 133-kDa N-terminal domain enables myosin 15 to maintain mechanotransducing stereocilia and is essential for hearing. ELife, 2015, 4, .	6.0	67
3	<scp>HESX</scp>1 mutations in patients with congenital hypopituitarism: variable phenotypes with the same genotype. Clinical Endocrinology, 2016, 85, 408-414.	2.4	24
4	High-throughput splicing assays identify missense and silent splice-disruptive POU1F1 variants underlying pituitary hormone deficiency. American Journal of Human Genetics, 2021, 108, 1526-1539.	6.2	23
5	Thyroid hormone is required for pruning, functioning and longâ€term maintenance of afferent inner hair cell synapses. European Journal of Neuroscience, 2016, 43, 148-161.	2.6	19
6	The phenotypic spectrum associated with OTX2 mutations in humans. European Journal of Endocrinology, 2021, 185, 121-135.	3.7	15
7	Genetic Background of Prop1 df Mutants Provides Remarkable Protection Against Hypothyroidism-Induced Hearing Impairment. JARO - Journal of the Association for Research in Otolaryngology, 2012, 13, 173-184.	1.8	14
8	A Modifier Gene Alleviates Hypothyroidism-Induced Hearing Impairment in Pou1f1dw Dwarf Mice. Genetics, 2011, 189, 665-673.	2.9	9
9	Growth hormone deficiency with advanced bone age: phenotypic interaction between GHRH receptor and CYP21A2 mutations diagnosed by sanger and whole exome sequencing. Archives of Endocrinology and Metabolism, 2017, 61, 633-636.	0.6	4
10	Genetic variation in thyroid folliculogenesis influences susceptibility to hypothyroidism-induced hearing impairment. Mammalian Genome, 2019, 30, 5-22.	2.2	2
11	Allelic Variants in Established Hypopituitarism Genes Expand Our Knowledge of the Phenotypic Spectrum. Genes, 2021, 12, 1128.	2.4	0