

Xiaofen Jin

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7218666/publications.pdf>

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2258059

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242
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#	ARTICLE	IF	CITATIONS
1	The exome sequencing identified the mutation in YARS2 encoding the mitochondrial tyrosyl-tRNA synthetase as a nuclear modifier for the phenotypic manifestation of Leber's hereditary optic neuropathy-associated mitochondrial DNA mutation. <i>Human Molecular Genetics</i> , 2016, 25, 584-596.	2.9	89
2	Leber's hereditary optic neuropathy caused by the homoplasmic ND1 m.3635G>A mutation in nine Han Chinese families. <i>Mitochondrion</i> , 2014, 18, 18-26.	3.4	29
3	Loss of MED1 triggers mitochondrial biogenesis in C2C12 cells. <i>Mitochondrion</i> , 2014, 14, 18-25.	3.4	12