

# Jialing Yu

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7102817/publications.pdf>

Version: 2024-02-01

7  
papers

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1478505

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#	ARTICLE	IF	CITATIONS
1	Leber's Hereditary Optic Neuropathy Arising From the Synergy Between ND1 3635G&gt;A Mutation and Mitochondrial YARS2 Mutations. , 2021, 62, 22.		10
2	PRICKLE3 linked to ATPase biogenesis manifested Leber's hereditary optic neuropathy. Journal of Clinical Investigation, 2020, 130, 4935-4946.	8.2	43
3	Contribution of mitochondrial ND1 3394T&gt;C mutation to the phenotypic manifestation of Leber's hereditary optic neuropathy. Human Molecular Genetics, 2019, 28, 1515-1529.	2.9	26
4	Mitochondrial DNA depletion, mitochondrial mutations and high TFAM expression in hepatocellular carcinoma. Oncotarget, 2017, 8, 84373-84383.	1.8	25
5	A Hypertension-Associated tRNA <sup>Ala</sup> Mutation Alters tRNA Metabolism and Mitochondrial Function. Molecular and Cellular Biology, 2016, 36, 1920-1930.	2.3	48
6	Aminoglycoside Stress Together with the 12S rRNA 1494C&gt;T Mutation Leads to Mitophagy. PLoS ONE, 2014, 9, e114650.	2.5	14
7	Loss of MED1 triggers mitochondrial biogenesis in C2C12 cells. Mitochondrion, 2014, 14, 18-25.	3.4	12