

# Aaron R D'souza

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1038398/publications.pdf>

Version: 2024-02-01

8  
papers

615  
citations

1307594

7  
h-index

1588992

8  
g-index

8  
all docs

8  
docs citations

8  
times ranked

1078  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mitochondrial transcription and translation: overview. <i>Essays in Biochemistry</i> , 2018, 62, 309-320.	4.7	192
2	TRMT5 Mutations Cause a Defect in Post-transcriptional Modification of Mitochondrial tRNA Associated with Multiple Respiratory-Chain Deficiencies. <i>American Journal of Human Genetics</i> , 2015, 97, 319-328.	6.2	83
3	Maturation of selected human mitochondrial tRNAs requires deadenylation. <i>ELife</i> , 2017, 6, .	6.0	72
4	METTL15 introduces N4-methylcytidine into human mitochondrial 12S rRNA and is required for mitoribosome biogenesis. <i>Nucleic Acids Research</i> , 2019, 47, 10267-10281.	14.5	70
5	Mitochondrial transcript maturation and its disorders. <i>Journal of Inherited Metabolic Disease</i> , 2015, 38, 655-680.	3.6	69
6	Defective mitochondrial rRNA methyltransferase MRM2 causes MELAS-like clinical syndrome. <i>Human Molecular Genetics</i> , 2017, 26, 4257-4266.	2.9	63
7	Biallelic C1QBP Mutations Cause Severe Neonatal-, Childhood-, or Later-Onset Cardiomyopathy Associated with Combined Respiratory-Chain Deficiencies. <i>American Journal of Human Genetics</i> , 2017, 101, 525-538.	6.2	58
8	YbeY is required for ribosome small subunit assembly and tRNA processing in human mitochondria. <i>Nucleic Acids Research</i> , 2021, 49, 5798-5812.	14.5	8