

# Svetlana Konovalova

## List of Publications by Year in descending order

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Version: 2024-02-01

14  
papers

434  
citations

840776

11  
h-index

1058476

14  
g-index

15  
all docs

15  
docs citations

15  
times ranked

969  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mitochondrial aminoacyl-tRNA synthetases in human disease. <i>Molecular Genetics and Metabolism</i> , 2013, 108, 206-211.	1.1	133
2	Structural modeling of tissue-specific mitochondrial alanyl-tRNA synthetase (AARS2) defects predicts differential effects on aminoacylation. <i>Frontiers in Genetics</i> , 2015, 6, 21.	2.3	46
3	Metabolic determination of cell fate through selective inheritance of mitochondria. <i>Nature Cell Biology</i> , 2022, 24, 148-154.	10.3	46
4	Targeted next-generation sequencing reveals further genetic heterogeneity in axonal Charcot-Marie-Tooth neuropathy and a mutation in HSPB1. <i>European Journal of Human Genetics</i> , 2014, 22, 522-527.	2.8	33
5	Editing activity for eliminating mischarged tRNAs is essential in mammalian mitochondria. <i>Nucleic Acids Research</i> , 2018, 46, 849-860.	14.5	30
6	Truncated HSPB1 causes axonal neuropathy and impairs tolerance to unfolded protein stress. <i>BBA Clinical</i> , 2015, 3, 233-242.	4.1	26
7	Redox regulation of GRPEL2 nucleotide exchange factor for mitochondrial HSP70 chaperone. <i>Redox Biology</i> , 2018, 19, 37-45.	9.0	25
8	ALS and Parkinson's disease genes CHCHD10 and CHCHD2 modify synaptic transcriptomes in human iPSC-derived motor neurons. <i>Neurobiology of Disease</i> , 2020, 141, 104940.	4.4	24
9	Instability of the mitochondrial alanyl-tRNA synthetase underlies fatal infantile-onset cardiomyopathy. <i>Human Molecular Genetics</i> , 2019, 28, 258-268.	2.9	19
10	Exposure to arginine analog canavanine induces aberrant mitochondrial translation products, mitoribosome stalling, and instability of the mitochondrial proteome. <i>International Journal of Biochemistry and Cell Biology</i> , 2015, 65, 268-274.	2.8	16
11	Y <sup>3+</sup> , La <sup>3+</sup> , and some bivalent metals inhibited the opening of the Tl <sup>+</sup> -induced permeability transition pore in Ca <sup>2+</sup> -loaded rat liver mitochondria. <i>Journal of Inorganic Biochemistry</i> , 2014, 141, 1-9.	3.5	15
12	A patient with pontocerebellar hypoplasia type 6: Novel RARS2 mutations, comparison to previously published patients and clinical distinction from PEHO syndrome. <i>European Journal of Medical Genetics</i> , 2020, 63, 103766.	1.3	10
13	Analysis of Mitochondrial Respiratory Chain Complexes in Cultured Human Cells using Blue Native Polyacrylamide Gel Electrophoresis and Immunoblotting. <i>Journal of Visualized Experiments</i> , 2019, , .	0.3	9
14	Analysis of Mitochondrial Protein Synthesis: De Novo Translation, Steady-State Levels, and Assembled OXPHOS Complexes. <i>Current Protocols in Toxicology</i> / Editorial Board, Mahin D Maines (editor-in-chief) [et Al ], 2018, 77, e56.	1.1	2