Roza Kucharczyk

List of Publications by Year in descending order

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201674 161849 5,781 54 27 54 citations h-index g-index papers 60 60 60 12361 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Assembly-dependent translation of subunits <i>6</i> (Atp6) and <i>9</i> (Atp9) of ATP synthase in yeast mitochondria. Genetics, 2022, 220, .	2.9	5
2	The pathogenic m.8993ÂT > C mutation in mitochondrial <i>ATP6</i> gene prevents proton releation the subunit <i>c</i> -ring rotor of ATP synthase. Human Molecular Genetics, 2021, 30, 381-392.	ase 2.9	8
3	5,6-diiodo-1H-benzotriazole: new TBBt analogue that minutely affects mitochondrial activity. Scientific Reports, 2021, 11, 23701.	3.3	2
4	The Unique Cysteine of F-ATP Synthase OSCP Subunit Participates in Modulation of the Permeability Transition Pore. Cell Reports, 2020, 32, 108095.	6.4	35
5	Case Report: Identification of a Novel Variant (m.8909T>C) of Human Mitochondrial ATP6 Gene and Its Functional Consequences on Yeast ATP Synthase. Life, 2020, 10, 215.	2.4	7
6	Molecular Basis of the Pathogenic Mechanism Induced by the m.9191T>C Mutation in Mitochondrial ATP6 Gene. International Journal of Molecular Sciences, 2020, 21, 5083.	4.1	12
7	ATP Synthase Subunit a Supports Permeability Transition in Yeast Lacking Dimerization Subunits and Modulates yPTP Conductance. Cellular Physiology and Biochemistry, 2020, 54, 211-229.	1.6	9
8	Assigning mitochondrial localization of dual localized proteins using a yeast Bi-Genomic Mitochondrial-Split-GFP. ELife, 2020, 9, .	6.0	20
9	Deregulating mitochondrial metabolite and ion transport has beneficial effects in yeast and human cellular models for NARP syndrome. Human Molecular Genetics, 2019, 28, 3792-3804.	2.9	12
10	Functional investigation of an universally conserved leucine residue in subunit a of ATP synthase targeted by the pathogenic m.9176â€T>G mutation. Biochimica Et Biophysica Acta - Bioenergetics, 2019, 1860, 52-59.	1.0	8
11	The pathogenic MT-ATP6 m.8851T>C mutation prevents proton movements within the n-side hydrophilic cleft of the membrane domain of ATP synthase. Biochimica Et Biophysica Acta - Bioenergetics, 2019, 1860, 562-572.	1.0	12
12	Perturbation of the yeast mitochondrial lipidome and associated membrane proteins following heterologous expression of Artemia-ANT. Scientific Reports, 2018, 8, 5915.	3.3	3
13	Two mutations in mitochondrial ATP6 gene of ATP synthase, related to human cancer, affect ROS, calcium homeostasis and mitochondrial permeability transition in yeast. Biochimica Et Biophysica Acta - Molecular Cell Research, 2018, 1865, 117-131.	4.1	36
14	High-Conductance Channel Formation in Yeast Mitochondria is Mediated by F-ATP Synthase e and g Subunits. Cellular Physiology and Biochemistry, 2018, 50, 1840-1855.	1.6	57
15	Protein AMPylation by an Evolutionarily Conserved Pseudokinase. Cell, 2018, 175, 809-821.e19.	28.9	149
16	Ancestral State Reconstruction of the Apoptosis Machinery in the Common Ancestor of Eukaryotes. G3: Genes, Genomes, Genetics, 2018, 8, 2121-2134.	1.8	32
17	Molecular basis of diseases caused by the mtDNA mutation m.8969G> A in the subunit a of ATP synthase. Biochimica Et Biophysica Acta - Bioenergetics, 2018, 1859, 602-611.	1.0	19
18	ATP Synthase Diseases of Mitochondrial Genetic Origin. Frontiers in Physiology, 2018, 9, 329.	2.8	88

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19	Decreasing cytosolic translation is beneficial to yeast and human Tafazzin-deficient cells. Microbial Cell, 2018, 5, 220-232.	3.2	13
20	Regulation of Aerobic Energy Metabolism in Podospora anserina by Two Paralogous Genes Encoding Structurally Different c-Subunits of ATP Synthase. PLoS Genetics, 2016, 12, e1006161.	3.5	6
21	Revisiting Mitochondrial pH with an Improved Algorithm for Calibration of the Ratiometric 5(6)-carboxy-SNARF-1 Probe Reveals Anticooperative Reaction with H+ Ions and Warrants Further Studies of Organellar pH. PLoS ONE, 2016, 11, e0161353.	2.5	18
22	Identification of G8969>A in mitochondrial ATP6 gene that severely compromises ATP synthase function in a patient with IgA nephropathy. Scientific Reports, 2016, 6, 36313.	3.3	28
23	Yeast models of mutations in the mitochondrial ATP6 gene found in human cancer cells. Mitochondrion, 2016, 29, 7-17.	3.4	14
24	Biallelic PPA2 Mutations Cause Sudden Unexpected Cardiac Arrest in Infancy. American Journal of Human Genetics, 2016, 99, 666-673.	6.2	39
25	Yeast as a system for modeling mitochondrial disease mechanisms and discovering therapies. DMM Disease Models and Mechanisms, 2015, 8, 509-526.	2.4	115
26	Mitochondrial protein sorting as a therapeutic target for ATP synthase disorders. Nature Communications, 2014, 5, 5585.	12.8	29
27	Expression of Nuclear and Mitochondrial Genes Encoding ATP Synthase Is Synchronized by Disassembly of a Multisynthetase Complex. Molecular Cell, 2014, 56, 763-776.	9.7	43
28	Defining the impact on yeast ATP synthase of two pathogenic human mitochondrial DNA mutations, T9185C and T9191C. Biochimie, 2014, 100, 200-206.	2.6	28
29	The depletion of F ₁ subunit $\hat{l}\mu$ in yeast leads to an uncoupled respiratory phenotype that is rescued by mutations in the proton-translocating subunits of F ₀ . Molecular Biology of the Cell, 2014, 25, 791-799.	2.1	10
30	Defining the pathogenesis of human mtDNA mutations using a yeast model: The case of T8851C. International Journal of Biochemistry and Cell Biology, 2013, 45, 130-140.	2.8	22
31	The Suppressor of AAC2 Lethality SAL1 Modulates Sensitivity of Heterologously Expressed Artemia ADP/ATP Carrier to Bongkrekate in Yeast. PLoS ONE, 2013, 8, e74187.	2.5	7
32	Experimental Relocation of the Mitochondrial ATP9 Gene to the Nucleus Reveals Forces Underlying Mitochondrial Genome Evolution. PLoS Genetics, 2012, 8, e1002876.	3.5	48
33	Guidelines for the use and interpretation of assays for monitoring autophagy. Autophagy, 2012, 8, 445-544.	9.1	3,122
34	Two Nuclear Life Cycle-Regulated Genes Encode Interchangeable Subunits c of Mitochondrial ATP Synthase in Podospora anserina. Molecular Biology and Evolution, 2011, 28, 2063-2075.	8.9	43
35	A Genetic Screen Targeted on the FO Component of Mitochondrial ATP Synthase in Saccharomyces cerevisiae. Journal of Biological Chemistry, 2011, 286, 18181-18189.	3.4	18
36	Energetic requirements and bioenergetic modulation of mitochondrial morphology and dynamics. Seminars in Cell and Developmental Biology, 2010, 21, 558-565.	5.0	87

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37	Mutants of the Saccharomyces cerevisiae VPS genes CCZ1 and YPT7 are blocked in different stages of sporulation. European Journal of Cell Biology, 2010, 89, 780-787.	3.6	17
38	Consequences of the pathogenic T9176C mutation of human mitochondrial DNA on yeast mitochondrial ATP synthase. Biochimica Et Biophysica Acta - Bioenergetics, 2010, 1797, 1105-1112.	1.0	54
39	Introducing the human Leigh syndrome mutation T9176G into Saccharomyces cerevisiae mitochondrial DNA leads to severe defects in the incorporation of Atp6p into the ATP synthase and in the mitochondrial morphology. Human Molecular Genetics, 2009, 18, 2889-2898.	2.9	53
40	Mitochondrial ATP synthase disorders: Molecular mechanisms and the quest for curative therapeutic approaches. Biochimica Et Biophysica Acta - Molecular Cell Research, 2009, 1793, 186-199.	4.1	99
41	Biochemical consequences in yeast of the human mitochondrial DNA 8993T>C mutation in the ATPase6 gene found in NARP/MILS patients. Biochimica Et Biophysica Acta - Molecular Cell Research, 2009, 1793, 817-824.	4.1	59
42	The Saccharomyces cerevisiae protein Cczlp interacts with components of the endosomal fusion machinery. FEMS Yeast Research, 2009, 9, 565-573.	2.3	8
43	The Leader Peptide of Yeast Atp6p Is Required for Efficient Interaction with the Atp9p Ring of the Mitochondrial ATPase. Journal of Biological Chemistry, 2007, 282, 36167-36176.	3.4	26
44	Yeast Cells Lacking the Mitochondrial Gene Encoding the ATP Synthase Subunit 6 Exhibit a Selective Loss of Complex IV and Unusual Mitochondrial Morphology. Journal of Biological Chemistry, 2007, 282, 10853-10864.	3.4	106
45	A Yeast Model of the Neurogenic Ataxia Retinitis Pigmentosa (NARP) T8993G Mutation in the Mitochondrial ATP Synthase-6 Gene. Journal of Biological Chemistry, 2007, 282, 34039-34047.	3.4	59
46	The CHiPS Domain - Ancient Traces for the Hermansky-Pudlak Syndrome. Traffic, 2005, 6, 534-538.	2.7	15
47	Increasing Mitochondrial Substrate-level Phosphorylation Can Rescue Respiratory Growth of an ATP Synthase-deficient Yeast. Journal of Biological Chemistry, 2005, 280, 30751-30759.	3.4	51
48	Failure to Assemble the $\hat{l}\pm 3$ $\hat{l}^2 3$ Subcomplex of the ATP Synthase Leads to Accumulation of the $\hat{l}\pm$ and \hat{l}^2 Subunits within Inclusion Bodies and the Loss of Mitochondrial Cristae in Saccharomyces cerevisiae. Journal of Biological Chemistry, 2005, 280, 18386-18392.	3.4	67
49	Multiple functions of the vacuolar sorting protein $Ccz1p$ in Saccharomyces cerevisiae. Biochemical and Biophysical Research Communications, 2005, 329, 197-204.	2.1	13
50	F1-catalysed ATP hydrolysis is required for mitochondrial biogenesis in Saccharomyces cerevisiae growing under conditions where it cannot respire. Molecular Microbiology, 2003, 47, 1329-1339.	2.5	45
51	The two rotor components of yeast mitochondrial ATP synthase are mechanically coupled by subunit Â. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 13235-13240.	7.1	48
52	The ATP synthase is involved in generating mitochondrial cristae morphology. EMBO Journal, 2002, 21, 221-230.	7.8	686
53	Identification of a Nuclear Gene (FMC1) Required for the Assembly/Stability of Yeast Mitochondrial F1-ATPase in Heat Stress Conditions. Journal of Biological Chemistry, 2001, 276, 6789-6796.	3.4	120
54	The yeast geneYJR025cencodes a 3-hydroxyanthranilic acid dioxygenase and is involved in nicotinic acid biosynthesis. FEBS Letters, 1998, 424, 127-130.	2.8	51