

Yidong Bai

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

4,557
citations

159585

30
h-index

123424

61
g-index

65
all docs

65
docs citations

65
times ranked

7064
citing authors

#	ARTICLE	IF	CITATIONS
1	Mitochondrial chaperones in human health and disease. <i>Free Radical Biology and Medicine</i> , 2022, 179, 363-374.	2.9	17
2	A novel sweetpotato RING-H2 type E3 ubiquitin ligase gene <i>lbATL38</i> enhances salt tolerance in transgenic <i>Arabidopsis</i> . <i>Plant Science</i> , 2021, 304, 110802.	3.6	25
3	Melatonin inhibits lung cancer development by reversing the Warburg effect via stimulating the SIRT3/PDH axis. <i>Journal of Pineal Research</i> , 2021, 71, e12755.	7.4	48
4	Guidelines for the use and interpretation of assays for monitoring autophagy (4th) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 622 Td (edition 9.1 1,430	9.1	1,430
5	Creating Cell Model 2.0 Using Patient Samples Carrying a Pathogenic Mitochondrial DNA Mutation: iPSC Approach for LHON. <i>Methods in Molecular Biology</i> , 2021, , .	0.9	3
6	Diagnostic value of circulating cell-free mtDNA in patients with suspected thyroid cancer: ND4/ND1 ratio as a new potential plasma marker. <i>Mitochondrion</i> , 2020, 55, 145-153.	3.4	14
7	BHRF1 Enhances EBV Mediated Nasopharyngeal Carcinoma Tumorigenesis through Modulating Mitophagy Associated with Mitochondrial Membrane Permeabilization Transition. <i>Cells</i> , 2020, 9, 1158.	4.1	8
8	Redox regulation by SOD2 modulates colorectal cancer tumorigenesis through AMPK-mediated energy metabolism. <i>Molecular Carcinogenesis</i> , 2020, 59, 545-556.	2.7	28
9	Emerging model systems and treatment approaches for Leber's hereditary optic neuropathy: Challenges and opportunities. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020, 1866, 165743.	3.8	27
10	Thioredoxin overexpression in mitochondria showed minimum effects on aging and age-related diseases in male C57BL/6 mice.. <i>Aging Pathobiology and Therapeutics</i> , 2020, 2, 20-31.	0.5	30
11	The implications of mitochondria in doxorubicin treatment of cancer in the context of traditional and modern medicine. <i>Traditional Medicine and Modern Medicine</i> , 2020, 03, 239-254.	0.2	0
12	Thioredoxin and aging: What have we learned from the survival studies?. <i>Aging Pathobiology and Therapeutics</i> , 2020, 2, 126-133.	0.5	4
13	Interval and continuous exercise overcome memory deficits related to β 2-Amyloid accumulation through modulating mitochondrial dynamics. <i>Behavioural Brain Research</i> , 2019, 376, 112171.	2.2	53
14	Mitophagy activation repairs Leber's hereditary optic neuropathy-associated mitochondrial dysfunction and improves cell survival. <i>Human Molecular Genetics</i> , 2019, 28, 422-433.	2.9	26
15	Oncocytic tumors are marked by enhanced mitochondrial content and mtDNA mutations of complex I in Chinese patients. <i>Mitochondrion</i> , 2019, 45, 1-6.	3.4	8
16	Dataset of mitochondrial genome variants in oncocytic tumors. <i>Data in Brief</i> , 2018, 17, 1149-1152.	1.0	1
17	The interaction between mitochondria and oncoviruses. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2018, 1864, 481-487.	3.8	9
18	Thioredoxin overexpression in both the cytosol and mitochondria accelerates age-related disease and shortens lifespan in male C57BL/6 mice. <i>GeroScience</i> , 2018, 40, 453-468.	4.6	18

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19	Aging-associated mitochondrial DNA mutations alter oxidative phosphorylation machinery and cause mitochondrial dysfunctions. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2017, 1863, 2266-2273.	3.8	30
20	Cyclophilin D over-expression increases mitochondrial complex III activity and accelerates supercomplex formation. <i>Archives of Biochemistry and Biophysics</i> , 2017, 613, 61-68.	3.0	12
21	Biochemical evidence for a mitochondrial genetic modifier in the phenotypic manifestation of Leber's hereditary optic neuropathy-associated mitochondrial DNA mutation. <i>Human Molecular Genetics</i> , 2016, 25, 3613-3625.	2.9	32
22	Mitochondrial metabolites extend lifespan. <i>Aging Cell</i> , 2016, 15, 336-348.	6.7	52
23	Mitochondrial common deletion is elevated in blood of breast cancer patients mediated by oxidative stress. <i>Mitochondrion</i> , 2016, 26, 104-112.	3.4	24
24	Mitochondrial DNA haplogroups modify the risk of osteoarthritis by altering mitochondrial function and intracellular mitochondrial signals. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2016, 1862, 829-836.	3.8	38
25	Respiratory supercomplexes plasticity and implications. <i>Frontiers in Bioscience - Landmark</i> , 2015, 20, 621-634.	3.0	30
26	Exercise intolerance and developmental delay associated with a novel mitochondrial ND5 mutation. <i>Scientific Reports</i> , 2015, 5, 10480.	3.3	19
27	Redefining the roles of mitochondrial DNA-encoded subunits in respiratory Complex I assembly. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2015, 1852, 1531-1539.	3.8	17
28	The Role of Mitochondria in T-2 Toxin-Induced Human Chondrocytes Apoptosis. <i>PLoS ONE</i> , 2014, 9, e108394.	2.5	49
29	A nonapoptotic role for CASP2/caspase 2. <i>Autophagy</i> , 2014, 10, 1054-1070.	9.1	63
30	Role of mtDNA Haplogroups in the Prevalence of Knee Osteoarthritis in a Southern Chinese Population. <i>International Journal of Molecular Sciences</i> , 2014, 15, 2646-2659.	4.1	44
31	Difference in apoptosis-associated genes expression profiling and immunohistology analysis between Kashin-Beck disease and primary osteoarthritis. <i>Science Bulletin</i> , 2014, 59, 833-839.	1.7	2
32	An update on complex I assembly: the assembly of players. <i>Journal of Bioenergetics and Biomembranes</i> , 2014, 46, 323-328.	2.3	25
33	Respiratory supercomplexes: structure, function and assembly. <i>Protein and Cell</i> , 2013, 4, 582-590.	11.0	70
34	Comparative bioenergetic study of neuronal and muscle mitochondria during aging. <i>Free Radical Biology and Medicine</i> , 2013, 63, 30-40.	2.9	17
35	Mitochondrial Common Deletion, a Potential Biomarker for Cancer Occurrence, Is Selected against in Cancer Background: A Meta-Analysis of 38 Studies. <i>PLoS ONE</i> , 2013, 8, e67953.	2.5	37
36	Tissue-specific implications of mitochondrial alterations in aging. <i>Frontiers in Bioscience - Elite</i> , 2013, E5, 734-747.	1.8	4

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37	Analysis of mitochondrial DNA variations in a Chinese family with spinocerebellar ataxia. <i>Journal of Clinical Neuroscience</i> , 2012, 19, 60-64.	1.5	5
38	Physiology and Pathophysiology of Mitochondrial DNA. <i>Advances in Experimental Medicine and Biology</i> , 2012, 942, 39-51.	1.6	37
39	Protein Phosphorylation and Prevention of Cytochrome Oxidase Inhibition by ATP: Coupled Mechanisms of Energy Metabolism Regulation. <i>Cell Metabolism</i> , 2011, 13, 712-719.	16.2	173
40	Mitochondrial respiratory complex I dysfunction promotes tumorigenesis through ROS alteration and AKT activation. <i>Human Molecular Genetics</i> , 2011, 20, 4605-4616.	2.9	129
41	Evaluating mitochondrial DNA in patients with breast cancer and benign breast disease. <i>Journal of Cancer Research and Clinical Oncology</i> , 2011, 137, 669-675.	2.5	36
42	The mitochondrial DNA 4,977-bp deletion and its implication in copy number alteration in colorectal cancer. <i>BMC Medical Genetics</i> , 2011, 12, 8.	2.1	103
43	Cancer type-specific modulation of mitochondrial haplogroups in breast, colorectal and thyroid cancer. <i>BMC Cancer</i> , 2010, 10, 421.	2.6	88
44	Evaluating mitochondrial DNA in cancer occurrence and development. <i>Annals of the New York Academy of Sciences</i> , 2010, 1201, 26-33.	3.8	37
45	Generation and bioenergetic analysis of cybrids containing mitochondrial DNA from mouse skeletal muscle during aging. <i>Nucleic Acids Research</i> , 2010, 38, 1913-1921.	14.5	11
46	Analysis of mitochondrial DNA mutations in D-loop region in thyroid lesions. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2010, 1800, 271-274.	2.4	24
47	Association of Mitochondrial DNA Haplogroups with Exceptional Longevity in a Chinese Population. <i>PLoS ONE</i> , 2009, 4, e6423.	2.5	45
48	A heteroplasmic, not homoplasmic, mitochondrial DNA mutation promotes tumorigenesis via alteration in reactive oxygen species generation and apoptosis. <i>Human Molecular Genetics</i> , 2009, 18, 1578-1589.	2.9	205
49	Mitochondrial Respiratory Complex I: Structure, Function and Implication in Human Diseases. <i>Current Medicinal Chemistry</i> , 2009, 16, 1266-1277.	2.4	256
50	Mitochondrial DNA mutations in the D-loop region may not be frequent in cervical cancer: a discussion on pitfalls in mitochondrial DNA studies. <i>Journal of Cancer Research and Clinical Oncology</i> , 2009, 135, 649-651.	2.5	14
51	Implications of mitochondrial DNA mutations and mitochondrial dysfunction in tumorigenesis. <i>Cell Research</i> , 2009, 19, 802-815.	12.0	234
52	An Assembled Complex IV Maintains the Stability and Activity of Complex I in Mammalian Mitochondria. <i>Journal of Biological Chemistry</i> , 2007, 282, 17557-17562.	3.4	112
53	Yeast NDI1 improves oxidative phosphorylation capacity and increases protection against oxidative stress and cell death in cells carrying a Leber's hereditary optic neuropathy mutation. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2007, 1772, 533-542.	3.8	40
54	Cytochrome c oxidase subunit IV is essential for assembly and respiratory function of the enzyme complex. <i>Journal of Bioenergetics and Biomembranes</i> , 2006, 38, 283-291.	2.3	255

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55	Nuclear Suppression of Mitochondrial Defects in Cells without the ND6 Subunit. <i>Molecular and Cellular Biology</i> , 2006, 26, 1077-1086.	2.3	32
56	Restoration of Mitochondrial Function in Cells with Complex I Deficiency. <i>Annals of the New York Academy of Sciences</i> , 2005, 1042, 25-35.	3.8	12
57	Specific point mutations may not accumulate with aging in the mouse mitochondrial DNA control region. <i>Gene</i> , 2005, 350, 193-199.	2.2	19
58	Genetic and Functional Analysis of Mitochondrial DNA-Encoded Complex I Genes. <i>Annals of the New York Academy of Sciences</i> , 2004, 1011, 272-283.	3.8	37
59	Genetic and Functional Analysis of Mitochondrial DNA-Encoded Complex I Genes. , 2004, 1011, 272-283.		23
60	Revisiting the mouse mitochondrial DNA sequence. <i>Nucleic Acids Research</i> , 2003, 31, 5349-5355.	14.5	101
61	Lack of Complex I Activity in Human Cells Carrying a Mutation in MtDNA-encoded ND4 Subunit Is Corrected by the <i>Saccharomyces cerevisiae</i> NADH-Quinone Oxidoreductase (NDI1) Gene. <i>Journal of Biological Chemistry</i> , 2001, 276, 38808-38813.	3.4	104
62	Tight Control of Respiration by NADH Dehydrogenase ND5 Subunit Gene Expression in Mouse Mitochondria. <i>Molecular and Cellular Biology</i> , 2000, 20, 805-815.	2.3	110