Wolfram S Kunz

List of Publications by Year
in descending order

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227 papers	13,687 citations	20817 60 h-index	29157 104 g-index
244 all docs	244 docs citations	244 times ranked	19823 citing authors

#	Article	IF	Citations
1	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	12.6	1,085
2	Analysis of mitochondrial function in situ in permeabilized muscle fibers, tissues and cells. Nature Protocols, 2008, 3, 965-976.	12.0	666
3	Characterization of Superoxide-producing Sites in Isolated Brain Mitochondria. Journal of Biological Chemistry, 2004, 279, 4127-4135.	3.4	443
4	Parkinson Phenotype in Aged PINK1-Deficient Mice Is Accompanied by Progressive Mitochondrial Dysfunction in Absence of Neurodegeneration. PLoS ONE, 2009, 4, e5777.	2.5	305
5	Permeabilized cell and skinned fiber techniques in studies of mitochondrial function in vitro. Molecular and Cellular Biochemistry, 1998, 184, 81-100.	3.1	302
6	Genetic determinants of common epilepsies: a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2014, 13, 893-903.	10.2	264
7	Recombination of Human Mitochondrial DNA. Science, 2004, 304, 981-981.	12.6	253
8	Impairment of mitochondrial function in skeletal muscle of patients with amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 1998, 156, 65-72.	0.6	252
9	Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals. American Journal of Human Genetics, 2019, 105, 267-282.	6.2	237
10	Mitochondrial complex I deficiency in the epileptic focus of patients with temporal lobe epilepsy. Annals of Neurology, 2000, 48, 766-773.	5.3	201
11	Loss-of-function mutations in MGME1 impair mtDNA replication and cause multisystemic mitochondrial disease. Nature Genetics, 2013, 45, 214-219.	21.4	198
12	Mitochondrial dysfunction, peroxidation damage and changes in glutathione metabolism in PARK6. Neurobiology of Disease, 2007, 25, 401-411.	4.4	180
13	Mitochondrial dysfunction and seizures: the neuronal energy crisis. Lancet Neurology, The, 2015, 14, 956-966.	10.2	176
14	Impaired mitochondrial oxidative phosphorylation in skeletal muscle of the dystrophin-deficient mdx mouse. Molecular and Cellular Biochemistry, 1998, 183, 87-96.	3.1	174
15	Epilepsy, hippocampal sclerosis and febrile seizures linked by common genetic variation around SCN1A. Brain, 2013, 136, 3140-3150.	7.6	168
16	Neurobehavioral Outcome Prediction After Cardiac Surgery. Stroke, 2000, 31, 645-650.	2.0	155
17	Mitochondrial potassium channels. IUBMB Life, 2009, 61, 134-143.	3.4	153
18	Linear mitochondrial DNA is rapidly degraded by components of the replication machinery. Nature Communications, 2018, 9, 1727.	12.8	151

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19	Mitochondrial dysfunction in epilepsy. Mitochondrion, 2012, 12, 35-40.	3.4	143
20	Seizure-dependent modulation of mitochondrial oxidative phosphorylation in rat hippocampus. European Journal of Neuroscience, 2002, 15, 1105-1114.	2.6	142
21	Myofiber integrity depends on desmin network targeting to Z-disks and costameres via distinct plectin isoforms. Journal of Cell Biology, 2008, 181, 667-681.	5.2	138
22	Genome-wide association analysis of genetic generalized epilepsies implicates susceptibility loci at 1q43, 2p16.1, 2q22.3 and 17q21.32. Human Molecular Genetics, 2012, 21, 5359-5372.	2.9	134
23	Fear Processing and Social Networking in the Absence of a Functional Amygdala. Biological Psychiatry, 2012, 72, 70-77.	1.3	123
24	Primary Skin Fibroblasts as a Model of Parkinson's Disease. Molecular Neurobiology, 2012, 46, 20-27.	4.0	121
25	The Mechanism of Neuroprotection by Topiramate in an Animal Model of Epilepsy. Epilepsia, 2004, 45, 1478-1487.	5.1	115
26	Contribution of different enzymes to flavoprotein fluorescence of isolated rat liver mitochondria. Biochimica Et Biophysica Acta - General Subjects, 1985, 841, 237-246.	2.4	113
27	Functional Imaging of Mitochondria in Saponin-permeabilized Mice Muscle Fibers. Journal of Cell Biology, 1998, 140, 1091-1099.	5.2	113
28	Permeabilized cell and skinned fiber techniques in studies of mitochondrial function in vivo. , 1998, , 81-100.		112
29	Visualization of defective mitochondrial function in skeletal muscle fibers of patients with sporadic amyotrophic lateral sclerosis. Journal of the Neurological Sciences, 1999, 169, 133-139.	0.6	111
30	BK channel openers inhibit ROS production of isolated rat brain mitochondria. Experimental Neurology, 2008, 212, 543-547.	4.1	109
31	The role of mitochondria in epileptogenesis. Current Opinion in Neurology, 2002, 15, 179-184.	3.6	103
32	Functional characterization of mitochondrial oxidative phosphorylation in saponin-skinned human muscle fibers. Biochimica Et Biophysica Acta - Bioenergetics, 1993, 1144, 46-53.	1.0	102
33	Mitochondrial involvement in temporal lobe epilepsy. Experimental Neurology, 2009, 218, 326-332.	4.1	102
34	Disorganization of the Desmin Cytoskeleton and Mitochondrial Dysfunction in Plectin-Related Epidermolysis Bullosa Simplex with Muscular Dystrophy. Journal of Neuropathology and Experimental Neurology, 2002, 61, 520-530.	1.7	96
35	On noxious desmin: functional effects of a novel heterozygous desmin insertion mutation on the extrasarcomeric desmin cytoskeleton and mitochondria. Human Molecular Genetics, 2003, 12, 657-669.	2.9	91
36	Burden Analysis of Rare Microdeletions Suggests a Strong Impact of Neurodevelopmental Genes in Genetic Generalised Epilepsies. PLoS Genetics, 2015, 11, e1005226.	3.5	91

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37	Opening of potassium channels modulates mitochondrial function in rat skeletal muscle. Biochimica Et Biophysica Acta - Bioenergetics, 2002, 1556, 97-105.	1.0	89
38	Impaired complex I repair causes recessive Leberâ€™s hereditary optic neuropathy. Journal of Clinical Investigation, 2021, 131, .	8.2	89
39	Microglial CD33-Related Siglec-E Inhibits Neurotoxicity by Preventing the Phagocytosis-Associated Oxidative Burst. Journal of Neuroscience, 2013, 33, 18270-18276.	3.6	87
40	Potassium channel openers depolarize hippocampal mitochondria. Brain Research, 2001, 892, 42-50.	2.2	86
41	N-acetyl Cysteine Treatment Rescues Cognitive Deficits Induced by Mitochondrial Dysfunction in G72/G30 Transgenic Mice. Neuropsychopharmacology, 2011, 36, 2233-2243.	5.4	84
42	Flux Control of Cytochrome c Oxidase in Human Skeletal Muscle. Journal of Biological Chemistry, 2000, 275, 27741-27745.	3.4	81
43	Mitochondrial potassium channels and reactive oxygen species. FEBS Letters, 2010, 584, 2043-2048.	2.8	80
44	Mitochondrial dysfunction due to Leber's hereditary optic neuropathy as a cause of visual loss during assessment for epilepsy surgery. Epilepsy and Behavior, 2011, 20, 38-43.	1.7	79
45	Mitochondrial dysfunction in neurodegenerative disorders. Biochemical Society Transactions, 2007, 35, 1228-1231.	3.4	77
46	Mosaic Deficiency in Mitochondrial Oxidative Metabolism Promotes Cardiac Arrhythmia during Aging. Cell Metabolism, 2015, 21, 667-677.	16.2	73
47	Estimation of flux control coefficients from inhibitor titrations by non-linear regression. FEBS Letters, 1990, 274, 167-170.	2.8	72
48	Subfieldâ€specific Loss of Hippocampal Nâ€acetyl Aspartate in Temporal Lobe Epilepsy. Epilepsia, 2008, 49, 40-50.	5.1	72
49	Sites of generation of reactive oxygen species in homogenates of brain tissue determined with the use of respiratory substrates and inhibitors. Biochimica Et Biophysica Acta - Bioenergetics, 2008, 1777, 689-695.	1.0	72
50	Effect of coenzyme Q10 on the mitochondrial function of skin fibroblasts from Parkinson patients. Journal of the Neurological Sciences, 2004, 220, 41-48.	0.6	71
51	A novel potassium channel in skeletal muscle mitochondria. Biochimica Et Biophysica Acta - Bioenergetics, 2008, 1777, 651-659.	1.0	70
52	Complex III-dependent superoxide production of brain mitochondria contributes to seizure-related ROS formation. Biochimica Et Biophysica Acta - Bioenergetics, 2010, 1797, 1163-1170.	1.0	70
53	Correlation of Hippocampal Glucose Oxidation Capacity and Interictal FDGâ€PET in Temporal Lobeâ€ƒEpilepsy. Epilepsia, 2003, 44, 193-199.	5.1	69
54	Mitochondrial potassium channels: From pharmacology to function. Biochimica Et Biophysica Acta - Bioenergetics, 2006, 1757, 715-720.	1.0	69

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55	Calcium Ions Regulate K+ Uptake into Brain Mitochondria: The Evidence for a Novel Potassium Channel. International Journal of Molecular Sciences, 2009, 10, 1104-1120.	4.1	69
56	Sonography of the median nerve in CMT1A, CMT2A, CMTX, and HNPP. Muscle and Nerve, 2013, 47, 385-395.	2.2	69
57	Characterization of superoxide production sites in isolated rat brain and skeletal muscle mitochondria. Biomedicine and Pharmacotherapy, 2005, 59, 163-168.	5.6	68
58	Quantification of the Content of Fluorescent Flavoproteins in Mitochondria from Liver, Kidney Cortex, Skeletal Muscle, and Brain. Biochemical Medicine and Metabolic Biology, 1993, 50, 103-110.	0.7	67
59	Rare coding variants in genes encoding GABAA receptors in genetic generalised epilepsies: an exome-based case-control study. Lancet Neurology, The, 2018, 17, 699-708.	10.2	67
60	Mitofusin 2 mutations affect mitochondrial function by mitochondrial DNA depletion. Acta Neuropathologica, 2013, 125, 245-256.	7.7	65
61	Linear mtDNA fragments and unusual mtDNA rearrangements associated with pathological deficiency of MGME1 exonuclease. Human Molecular Genetics, 2014, 23, 6147-6162.	2.9	64
62	Behavioral changes in G72/G30 transgenic mice. European Neuropsychopharmacology, 2009, 19, 339-348.	0.7	63
63	Loss of UCP2 Attenuates Mitochondrial Dysfunction without Altering ROS Production and Uncoupling Activity. PLoS Genetics, 2014, 10, e1004385.	3.5	63
64	Mitochondrial DNA damage and the aging processâ€“facts and imaginations. Free Radical Research, 2006, 40, 1284-1294.	3.3	62
65	Metabolic progression markers of neurodegeneration in the transgenic G93A-SOD1 mouse model of amyotrophic lateral sclerosis. European Journal of Neuroscience, 2007, 25, 1669-1677.	2.6	61
66	Genetic variation in dopaminergic activity is associated with the risk for psychiatric side effects of levetiracetam. Epilepsia, 2013, 54, 36-44.	5.1	61
67	16p11.2 600 kb Duplications confer risk for typical and atypical Rolandic epilepsy. Human Molecular Genetics, 2014, 23, 6069-6080.	2.9	61
68	Cryopreservation of mitochondria and mitochondrial function in cardiac and skeletal muscle fibers. Analytical Biochemistry, 2003, 319, 296-303.	2.4	60
69	Changes in mitochondrial reactive oxygen species synthesis during differentiation of skeletal muscle cells. Mitochondrion, 2012, 12, 144-148.	3.4	60
70	Evaluation of Methods for the Determination of Mitochondrial Respiratory Chain Enzyme Activities in Human Skeletal Muscle Samples. Analytical Biochemistry, 2000, 279, 55-60.	2.4	59
71	Rare exonic deletions of the <scp>RBFOX1</scp> gene increase risk of idiopathic generalized epilepsy. Epilepsia, 2013, 54, 265-271.	5.1	59
72	Exonâ€disrupting deletions of <scp>NRXN1</scp> in idiopathic generalized epilepsy. Epilepsia, 2013, 54, 256-264.	5.1	59

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73	A homozygous splice-site mutation in CARS2 is associated with progressive myoclonic epilepsy. Neurology, 2014, 83, 2183-2187.	1.1	59
74	Different Metabolic Properties of Mitochondrial Oxidative Phosphorylation in Different Cell Types - Important Implications for Mitochondrial Cytopathies. Experimental Physiology, 2003, 88, 149-154.	2.0	57
75	Mutant desmin substantially perturbs mitochondrial morphology, function and maintenance in skeletal muscle tissue. Acta Neuropathologica, 2016, 132, 453-473.	7.7	57
76	Metabolic consequences of a novel missense mutation of the mtDNA CO I gene. Human Molecular Genetics, 2002, 11, 1797-1805.	2.9	56
77	Recombination of mitochondrial DNA in skeletal muscle of individuals with multiple mitochondrial DNA heteroplasmy. Nature Genetics, 2005, 37, 873-877.	21.4	56
78	Binding of copper is a mechanism of homocysteine toxicity leading to COX deficiency and apoptosis in primary neurons, PC12 and SHSY-5Y cells. Neurobiology of Disease, 2006, 23, 725-730.	4.4	55
79	Multidrug Resistance Protein-1 Affects Oxidative Stress, Endothelial Dysfunction, and Atherogenesis via Leukotriene C ₄ Export. Circulation, 2008, 117, 2912-2918.	1.6	55
80	Carbamazepineâ€•and oxcarbazepineâ€•induced hyponatremia in people with epilepsy. Epilepsia, 2017, 58, 1227-1233.	5.1	54
81	Application of inhibitor titrations for the detection of oxidative phosphorylation defects in saponin-skinned muscle fibers of patients with mitochondrial diseases. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1997, 1360, 142-150.	3.8	53
82	Sustained seizure remission on perampanel in progressive myoclonic epilepsy (Lafora disease). Epilepsy & Behavior Case Reports, 2013, 1, 118-121.	1.5	53
83	Volumetric Magnetic Resonance Imaging of Functionally Relevant Structural Alterations in Chronic Epilepsy after Pilocarpine-induced Status Epilepticus in Rats. Epilepsia, 2005, 46, 1021-1026.	5.1	52
84	Re-evaluation of the dysfunction of mitochondrial respiratory chain in skeletal muscle of patients with Parkinson?s disease. Journal of Neural Transmission, 2005, 112, 499-518.	2.8	52
85	In vivo quantification of spinal and bulbar motor neuron degeneration in the G93A-SOD1 transgenic mouse model of ALS by T2 relaxation time and apparent diffusion coefficient. Experimental Neurology, 2006, 201, 293-300.	4.1	52
86	Exonic microdeletions of the gephyrin gene impair GABAergic synaptic inhibition in patients with idiopathic generalized epilepsy. Neurobiology of Disease, 2014, 67, 88-96.	4.4	51
87	Altered mitochondrial oxidative phosphorylation in hippocampal slices of kainate-treated rats. Brain Research, 1999, 826, 236-242.	2.2	50
88	The influence of the cytosolic oncotic pressure on the permeability of the mitochondrial outer membrane for ADP: implications for the kinetic properties of mitochondrial creatine kinase and for ADP channelling into the intermembrane space. Molecular and Cellular Biochemistry, 1994, 133-134, 85-104.	3.1	49
89	Increase of Flux Control of Cytochrome c Oxidase in Copper-deficient Mottled Brindled Mice. Journal of Biological Chemistry, 1996, 271, 283-288.	3.4	49
90	Repeats, longevity and the sources of mtDNA deletions: evidence from â€˜deletional spectraâ€™. Trends in Genetics, 2010, 26, 340-343.	6.7	48

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91	Bloodâ€“brain barrier dysfunction can contribute to pharmacoresistance of seizures. Epilepsia, 2014, 55, 1255-1263.	5.1	47
92	Is There Still Any Role for Oxidative Stress in Mitochondrial DNA-Dependent Aging?. Genes, 2018, 9, 175.	2.4	47
93	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17â€‰458 subjects. Brain, 2020, 143, 2106-2118.	7.6	47
94	Spectral properties of fluorescent flavoproteins of isolated rat liver mitochondria. FEBS Letters, 1986, 195, 92-96.	2.8	45
95	Effect of Creatine Supplementation on Metabolite Levels in ALS Motor Cortices. Experimental Neurology, 2001, 172, 377-382.	4.1	45
96	Biallelic Mutations of VAC14 in Pediatric-Onset Neurological Disease. American Journal of Human Genetics, 2016, 99, 188-194.	6.2	45
97	Loss of the smallest subunit of cytochrome c oxidase, COX8A, causes Leigh-like syndrome and epilepsy. Brain, 2016, 139, 338-345.	7.6	44
98	Severe epilepsy as the major symptom of new mutations in the mitochondrial tRNA ^{Phe} gene. Neurology, 2010, 74, 507-512.	1.1	43
99	The contribution of thioredoxin-2 reductase and glutathione peroxidase to H2O2 detoxification of rat brain mitochondria. Biochimica Et Biophysica Acta - Bioenergetics, 2012, 1817, 1901-1906.	1.0	43
100	Genetic variation in CFH predicts phenytoin-induced maculopapular exanthema in European-descent patients. Neurology, 2018, 90, e332-e341.	1.1	43
101	Cytosolic, but not matrix, calcium is essential for adjustment of mitochondrial pyruvate supply. Journal of Biological Chemistry, 2020, 295, 4383-4397.	3.4	43
102	New Insights into the Metabolic Consequences of Large-Scale mtDNA Deletions: A Quantitative Analysis of Biochemical, Morphological, and Genetic Findings in Human Skeletal Muscle. Journal of Neuropathology and Experimental Neurology, 2000, 59, 353-360.	1.7	42
103	Effect of 1-methyl-4-phenylpyridinium on glutathione in rat pheochromocytoma PC 12 cells. Neurochemistry International, 2000, 36, 489-497.	3.8	41
104	Automated CD61 immunoplatelet analysis of thrombocytopenic samples. British Journal of Haematology, 2001, 112, 584-592.	2.5	41
105	Doseâ€dependent memory effects and cerebral volume changes after in utero exposure to valproate in the rat. Epilepsia, 2009, 50, 1432-1441.	5.1	41
106	Amelioration of water maze performance deficits by topiramate applied during pilocarpine-induced status epilepticus is negatively dose-dependent. Epilepsy Research, 2007, 73, 173-180.	1.6	39
107	Clonally Expanded Mitochondrial DNA Mutations in Epileptic Individuals With Mutated DNA Polymerase Î³. Journal of Neuropathology and Experimental Neurology, 2008, 67, 857-866.	1.7	39
108	Evaluation of a procedure for the simultaneous determination of oxidized and reduced pyridine nucleotides and adenylates in organic phenol extracts from mitochondria. Analytical Biochemistry, 1992, 202, 162-165.	2.4	38

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109	Mitochondrial dysfunction in myofibrillar myopathy. Neuropathology and Applied Neurobiology, 2003, 29, 45-51.	3.2	38
110	Human Epidermal Keratinocytes Accumulate Superoxide Due to Low Activity of Mn-SOD, Leading to Mitochondrial Functional Impairment. Journal of Investigative Dermatology, 2007, 127, 1084-1093.	0.7	38
111	Extending the phenotypic spectrum of <scp>RBFOX</scp>1 deletions: Sporadic focal epilepsy. Epilepsia, 2015, 56, e129-33.	5.1	38
112	Neuropathological signs of inflammation correlate with mitochondrial DNA deletions in mesial temporal lobe epilepsy. Acta Neuropathologica, 2016, 132, 277-288.	7.7	37
113	Clonal expansions of mitochondrial genomes: implications for in vivo mutational spectra. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2003, 522, 13-19.	1.0	36
114	Complete failure of insulin-transmitted signaling, but not obesity-induced insulin resistance, impairs respiratory chain function in muscle. Journal of Molecular Medicine, 2012, 90, 1145-1160.	3.9	36
115	Neuroimaging characteristics in mitochondrial encephalopathies associated with the m.3243A>G MTTL1 mutation. Journal of Neurology, 2013, 260, 1071-1080.	3.6	36
116	Sub-genic intolerance, ClinVar, and the epilepsies: A whole-exome sequencing study of 29,165 individuals. American Journal of Human Genetics, 2021, 108, 965-982.	6.2	35
117	Control of oxidative phosphorylation in skeletal muscle. Biochimica Et Biophysica Acta - Bioenergetics, 2001, 1504, 12-19.	1.0	34
118	Mitochondrial involvement in neurodegenerative diseases. IUBMB Life, 2013, 65, 263-272.	3.4	34
119	Comparative effectiveness of antiepileptic drugs in juvenile myoclonic epilepsy. Epilepsia Open, 2019, 4, 420-430.	2.4	34
120	Distribution of Flux Control among the Enzymes of Mitochondrial Oxidative Phosphorylation in Calcium-activated Saponin-skinned Rat Musculus Soleus Fibers. FEBS Journal, 1995, 230, 549-554.	0.2	33
121	POLG mutations cause decreased mitochondrial DNA repopulation rates following induced depletion in human fibroblasts. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2011, 1812, 321-325.	3.8	33
122	Mitochondrial oxidative phosphorylation in saponin-skinned human muscle fibers is stimulated by caffeine. FEBS Letters, 1993, 323, 188-190.	2.8	32
123	Measurement of Fluorescence Changes of NAD(P)H and of Fluorescent Flavoproteins in Saponin-Skinned Human Skeletal Muscle Fibers. Analytical Biochemistry, 1994, 216, 322-327.	2.4	32
124	Genomeâ€wide linkage metaâ€analysis identifies susceptibility loci at 2q34 and 13q31.3 for genetic generalized epilepsies. Epilepsia, 2012, 53, 308-318.	5.1	32
125	Oxygen dependence of flux control of cytochromecoxidase - implications for mitochondrial diseases. FEBS Letters, 1998, 422, 33-35.	2.8	31
126	A VASP-Racâ€“Soluble Guanylyl Cyclase Pathway Controls cGMP Production in Adipocytes. Science Signaling, 2012, 5, ra62.	3.6	31

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127	Hemin inhibits the large conductance potassium channel in brain mitochondria: A putative novel mechanism of neurodegeneration. Experimental Neurology, 2014, 257, 70-75.	4.1	31
128	Homozygous mutation in TXNRD1 is associated with genetic generalized epilepsy. Free Radical Biology and Medicine, 2017, 106, 270-277.	2.9	31
129	Mitochondrial dysfunction in neurological disorders with epileptic phenotypes. Journal of Bioenergetics and Biomembranes, 2010, 42, 443-448.	2.3	30
130	Single channel studies of the ATP-regulated potassium channel in brain mitochondria. Journal of Bioenergetics and Biomembranes, 2009, 41, 323-334.	2.3	28
131	Evaluation of electron-transfer flavoprotein and Î±-lipoamide dehydrogenase redox states by two-channel fluorimetry and its application to the investigation of Î²-oxidation. Biochimica Et Biophysica Acta - Bioenergetics, 1988, 932, 8-16.	1.0	27
132	Guide to the Pharmacology of Mitochondrial Potassium Channels. Handbook of Experimental Pharmacology, 2016, 240, 103-127.	1.8	27
133	Effects of bc 1 -site electron transfer inhibitors on the absorption spectra of mitochondrial cytochromes b. FEBS Letters, 1985, 181, 95-99.	2.8	26
134	Hippocampal N-acetyl aspartate levels do not mirror neuronal cell densities in creatine-supplemented epileptic rats. European Journal of Neuroscience, 2003, 18, 2292-2300.	2.6	26
135	Chapter 23 Quantification of Superoxide Production by Mouse Brain and Skeletal Muscle Mitochondria. Methods in Enzymology, 2009, 456, 419-437.	1.0	26
136	Comparative effectiveness of antiepileptic drugs in patients with mesial temporal lobe epilepsy with hippocampal sclerosis. Epilepsia, 2017, 58, 1734-1741.	5.1	26
137	Mitochondrial Retinopathy. Ophthalmology Retina, 2022, 6, 65-79.	2.4	26
138	Detection of Respiratory Chain Defects in Cultivated Skin Fibroblasts and Skeletal Muscle of Patients with Parkinson's Disease. Annals of the New York Academy of Sciences, 1999, 893, 426-429.	3.8	25
139	Metabolic Consequences of the Cytochrome c Oxidase Deficiency in Brain of Copper-Deficient Movbr Mice. Journal of Neurochemistry, 2001, 72, 1580-1585.	3.9	25
140	Mitochondrial Diseases - An Expanding Spectrum of Disorders and Affected Genes. Experimental Physiology, 2003, 88, 155-166.	2.0	25
141	Antidiabetic sulphonylureas activate mitochondrial permeability transition in rat skeletal muscle. British Journal of Pharmacology, 2005, 145, 785-791.	5.4	25
142	Mitochondrial Liver Toxicity of Valproic Acid and Its Acid Derivatives Is Related to Inhibition of Î±-Lipoamide Dehydrogenase. International Journal of Molecular Sciences, 2017, 18, 1912.	4.1	25
143	Oxygraphic Evaluation of Mitochondrial Function in Digitonin-Permeabilized Mononuclear Cells and Cultured Skin Fibroblasts of Patients with Chronic Progressive External Ophthalmoplegia. Biochemical and Molecular Medicine, 1995, 54, 105-111.	1.4	24
144	Heme is required for carbon monoxide activation of mitochondrial BKCa channel. European Journal of Pharmacology, 2020, 881, 173191.	3.5	24

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145	Signaling pathways targeting mitochondrial potassium channels. International Journal of Biochemistry and Cell Biology, 2020, 125, 105792.	2.8	24
146	Effect of b-itc 1 -site inhibitors on the midpoint potentials of mitochondrial cytochromes b. FEBS Letters, 1983, 155, 237-240.	2.8	23
147	Primary carnitine deficiency: adult onset lipid storage myopathy with a mild clinical course. Journal of Clinical Neuroscience, 2004, 11, 919-924.	1.5	23
148	Distinct patterns of mitochondrial genome diversity in bonobos (Pan paniscus) and humans. BMC Evolutionary Biology, 2010, 10, 270.	3.2	23
149	Novel SCARB2 mutation in action myoclonus-renal failure syndrome and evaluation of SCARB2mutations in isolated AMRF features. BMC Neurology, 2011, 11, 134.	1.8	22
150	Mesial temporal lobe epilepsy associated with KCNT1 mutation. Seizure: the Journal of the British Epilepsy Association, 2017, 45, 181-183.	2.0	22
151	Functional variants in <scp>HCN</scp>4 and <scp>CACNA</scp>1H may contribute to genetic generalized epilepsy. Epilepsia Open, 2017, 2, 334-342.	2.4	22
152	Testing association of rare genetic variants with resistance to three common antiseizure medications. Epilepsia, 2020, 61, 657-666.	5.1	22
153	Evaluation of Presumably Disease Causing SCN1A Variants in a Cohort of Common Epilepsy Syndromes. PLoS ONE, 2016, 11, e0150426.	2.5	22
154	Caffeine and Ca2+ stimulate mitochondrial oxidative phosphorylation in saponin-skinned human skeletal muscle fibers due to activation of actomyosin ATPase. Biochimica Et Biophysica Acta - Bioenergetics, 1994, 1188, 373-379.	1.0	21
155	Epileptic Focus and Alteration of Metabolism. International Review of Neurobiology, 2014, 114, 209-243.	2.0	21
156	Pharmacoresponse in genetic generalized epilepsy: a genome-wide association study. Pharmacogenomics, 2020, 21, 325-335.	1.3	21
157	Inheritance of Mitochondrial DNA Recombinants in Double-Heteroplasmic Families: Potential Implications for Phylogenetic Analysis. American Journal of Human Genetics, 2007, 80, 298-305.	6.2	20
158	Replication fork rescue in mammalian mitochondria. Scientific Reports, 2019, 9, 8785.	3.3	20
159	Overexpression of bcl-2 Results in Reduction of Cytochrome c Content and Inhibition of Complex I Activity. Biochemical and Biophysical Research Communications, 2001, 280, 1021-1027.	2.1	19
160	Defective Mitochondrial Oxidative Phosphorylation in Myopathies with Tubular Aggregates Originating from Sarcoplasmic Reticulum. Journal of Neuropathology and Experimental Neurology, 2001, 60, 1032-1040.	1.7	19
161	Flow Cytometric Detection of Mitochondrial Dysfunction in Subpopulations of Human Mononuclear Cells. Analytical Biochemistry, 1997, 246, 218-224.	2.4	17
162	Peripheral nerve atrophy together with higher cerebrospinal fluid progranulin indicate axonal damage in amyotrophic lateral sclerosis. Muscle and Nerve, 2018, 57, 273-278.	2.2	17

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163	Mitochondrial BK Channel Openers CGS7181 and CGS7184 Exhibit Cytotoxic Properties. International Journal of Molecular Sciences, 2018, 19, 353.	4.1	17
164	Quasi-Mendelian paternal inheritance of mitochondrial DNA: A notorious artifact, or anticipated behavior?. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 14797-14798.	7.1	17
165	fibers of patients with chronic progressive external ophthalmoplegia. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1995, 1272, 181-184.	3.8	16
166	Oxyphil Cell Metaplasia in the Parathyroids Is Characterized by Somatic Mitochondrial DNA Mutations in NADH Dehydrogenase Genes and Cytochrome c Oxidase Activityâ€“Impairing Genes. American Journal of Pathology, 2014, 184, 2922-2935.	3.8	16
167	Impairment of mitochondrial oxidative phosphorylation in skin fibroblasts of SALS and FALS patients is rescued by in vitro treatment with ROS scavengers. Experimental Neurology, 2021, 339, 113620.	4.1	16
168	Assessing the role of rare genetic variants in drugâ€resistant, nonâ€lesional focal epilepsy. Annals of Clinical and Translational Neurology, 2021, 8, 1376-1387.	3.7	16
169	Quantitative investigation of mitochondrial function in single rat hippocampal slices: a novel application of high-resolution respirometry and laser-excited fluorescence spectroscopy. Brain Research Protocols, 1999, 4, 329-334.	1.6	15
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