Peter Bross

List of Publications by Year
in descending order

123 papers	6,996 citations	47 h-index	79 g-index
		47006	64796
125 all docs	125 docs citations	125 times ranked	7277 citing authors

#	Article	IF	Citations
1	Î±-cardiac actin is a novel disease gene in familial hypertrophic cardiomyopathy. Journal of Clinical Investigation, 1999, 103, R39-R43.	8.2	353
2	Hereditary Spastic Paraplegia SPG13 Is Associated with a Mutation in the Gene Encoding the Mitochondrial Chaperonin Hsp60. American Journal of Human Genetics, 2002, 70, 1328-1332.	6.2	347
3	Clear Correlation of Genotype with Disease Phenotype in Veryâ€"Long-Chain Acyl-CoA Dehydrogenase Deficiency. American Journal of Human Genetics, 1999, 64, 479-494.	6.2	285
4	Protein Misfolding and Human Disease. Annual Review of Genomics and Human Genetics, 2006, 7, 103-124.	6.2	258
5	Medium-Chain Acyl-CoA Dehydrogenase (MCAD) Mutations Identified by MS/MS-Based Prospective Screening of Newborns Differ from Those Observed in Patients with Clinical Symptoms: Identification and Characterization of a New, Prevalent Mutation That Results in Mild MCAD Deficiency*. American Journal of Human Genetics, 2001, 68, 1408-1418.	6.2	219
6	Clear relationship betweenETF/ETFDH genotype and phenotype in patients with multiple acyl-CoA dehydrogenation deficiency. Human Mutation, 2003, 22, 12-23.	2.5	196
7	Mitochondrial Hsp60 Chaperonopathy Causes an Autosomal-Recessive Neurodegenerative Disorder Linked to Brain Hypomyelination and Leukodystrophy. American Journal of Human Genetics, 2008, 83, 30-42.	6.2	195
8	Protein misfolding and degradation in genetic diseases. Human Mutation, 1999, 14, 186-198.	2.5	184
9	Mutation analysis in mitochondrial fatty acid oxidation defects: Exemplified by acyl-CoA dehydrogenase deficiencies, with special focus on genotype-phenotype relationship. Human Mutation, 2001, 18, 169-189.	2.5	178
10	Genomic structure of the human mitochondrial chaperonin genes: HSP60 and HSP10 are localised head to head on chromosome 2 separated by a bidirectional promoter. Human Genetics, 2003, 112, 71-77.	3.8	131
11	Protein Misfolding and Cellular Stress: An Overview. Methods in Molecular Biology, 2010, 648, 3-23.	0.9	129
12	Mitochondrial fatty acid oxidation defectsâ€"remaining challenges. Journal of Inherited Metabolic Disease, 2008, 31, 643-657.	3.6	123
13	The Molecular Basis of Medium-Chain Acyl-CoA Dehydrogenase (MCAD) Deficiency in Compound Heterozygous Patients: Is There Correlation between Genotype and Phenotype?. Human Molecular Genetics, 1997, 6, 695-707.	2.9	119
14	Dissection of functional domains in phage fd adsorption protein. Journal of Molecular Biology, 1990, 212, 143-149.	4.2	115
15	Identification of four new mutations in the short-chain acyl-CoA dehydrogenase (SCAD) gene in two patients: one of the variant alleles, 511C-->T, is present at an unexpectedly high frequency in the general population, as was the case for 625G-->A, together conferring susceptibility to ethylmalonic aciduria. Human Molecular Genetics, 1998, 7, 619-627.	2.9	109
16	Cloning and characterization of human very-long-chain acyl-CoA dehydrogenase cDNA, chromosomal assignment of the gene and identification in four patients of nine different mutations within the VLCAD gene [published erratum appears in Hum Mol Genet 1996 Sep;5(9):1390]. Human Molecular Genetics, 1996, 5, 461-472.	2.9	106
17	The ACADS gene variation spectrum in 114 patients with short-chain acyl-CoA dehydrogenase (SCAD) deficiency is dominated by missense variations leading to protein misfolding at the cellular level. Human Genetics, 2008, 124, 43-56.	3.8	101
18	Leptin regulation of Hsp60 impacts hypothalamic insulin signaling. Journal of Clinical Investigation, 2013, 123, 4667-4680.	8.2	101

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19	Influence of Lewis Î±1-3/4-L-Fucosyltransferase (FUT3) Gene Mutations on Enzyme Activity, Erythrocyte Phenotyping, and Circulating Tumor Marker Sialyl-Lewis a Levels. Journal of Biological Chemistry, 1996, 271, 32260-32268.	3.4	94
20	Riboflavin Deficiencyâ€"Implications for General Human Health and Inborn Errors of Metabolism. International Journal of Molecular Sciences, 2020, 21, 3847.	4.1	92
21	Ethylmalonic Aciduria Is Associated with an Amino Acid Variant of Short Chain Acyl-Coenzyme A Dehydrogenase. Pediatric Research, 1996, 39, 1059-1066.	2.3	92
22	Genetic defects in fatty acid beta-oxidation and acyl-CoA dehydrogenases. Molecular pathogenesis and genotype-phenotype relationships. FEBS Journal, 2004, 271, 470-482.	0.2	86
23	Inactivation of the hereditary spastic paraplegia-associated Hspd1 gene encoding the Hsp60 chaperone results in early embryonic lethality in mice. Cell Stress and Chaperones, 2010, 15, 851-863.	2.9	83
24	The Hsp60-(p.V98I) Mutation Associated with Hereditary Spastic Paraplegia SPG13 Compromises Chaperonin Function Both in Vitro and in Vivo. Journal of Biological Chemistry, 2008, 283, 15694-15700.	3.4	80
25	Molecular mechanisms of riboflavin responsiveness in patients with ETF-QO variations and multiple acyl-CoA dehydrogenation deficiency. Human Molecular Genetics, 2012, 21, 3435-3448.	2.9	80
26	Effects of Two Mutations Detected in Medium Chain Acyl-CoA Dehydrogenase (MCAD)-deficient Patients on Folding, Oligomer Assembly, and Stability of MCAD Enzyme. Journal of Biological Chemistry, 1995, 270, 10284-10290.	3.4	79
27	Isolated 2-Methylbutyrylglycinuria Caused by Short/Branched-Chain Acyl-CoA Dehydrogenase Deficiency: Identification of a New Enzyme Defect, Resolution of Its Molecular Basis, and Evidence for Distinct Acyl-CoA Dehydrogenases in Isoleucine And Valine Metabolism. American Journal of Human Genetics, 2000, 67, 1095-1103.	6.2	79
28	Medium-Chain Acyl-CoA Dehydrogenase (MCAD) Deficiency: The Prevalent Mutation G985 (K304E) Is Subject to a Strong Founder Effect from Northwestern Europe. Human Heredity, 1993, 43, 342-350.	0.8	75
29	Misfolding, Degradation, and Aggregation of Variant Proteins. Journal of Biological Chemistry, 2003, 278, 47449-47458.	3.4	74
30	Decreased expression of the mitochondrial matrix proteases Lon and ClpP in cells from a patient with hereditary spastic paraplegia (SPG13). Neuroscience, 2008, 153, 474-482.	2.3	74
31	Emerging Roles for Riboflavin in Functional Rescue of Mitochondrial β-Oxidation Flavoenzymes. Current Medicinal Chemistry, 2010, 17, 3842-3854.	2.4	73
32	Actin mutations in hypertrophic and dilated cardiomyopathy cause inefficient protein folding and perturbed filament formation. FEBS Journal, 2005, 272, 2037-2049.	4.7	71
33	A human homologue of Escherichia coli ClpP caseinolytic protease: recombinant expression, intracellular processing and subcellular localization. Biochemical Journal, 1998, 331, 309-316.	3.7	67
34	Role of Flavinylation in a Mild Variant of Multiple Acyl-CoA Dehydrogenation Deficiency. Journal of Biological Chemistry, 2009, 284, 4222-4229.	3.4	67
35	Molecular characterization of medium-chain acyl-CoA dehydrogenase (MCAD) deficiency: identification of a lys329 to glu mutation in the MCAD gene, and expression of inactive mutant enzyme protein in E. coli. Human Genetics, 1991, 86, 545-51.	3.8	66
36	Reduced heat shock response in human mononuclear cells during aging and its association with polymorphisms in HSP70 genes. Cell Stress and Chaperones, 2006, 11, 208.	2.9	66

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37	The role of chaperone-assisted folding and quality control in inborn errors of metabolism: Protein folding disorders. Journal of Inherited Metabolic Disease, 2001, 24, 189-212.	3.6	65
38	Co-overexpression of bacterial GroESL chaperonins partly overcomes non-productive folding and tetramer assembly of E. coli-expressed human medium-chain acyl-CoA dehydrogenase (MCAD) carrying the prevalent disease-causing K304E mutation. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1993, 1182, 264-274.	3.8	64
39	CLPB Variants Associated with Autosomal-Recessive Mitochondrial Disorder with Cataract, Neutropenia, Epilepsy, and Methylglutaconic Aciduria. American Journal of Human Genetics, 2015, 96, 258-265.	6.2	58
40	Medium-Long-Chain Chimeric Human Acyl-CoA Dehydrogenase:Â Medium-Chain Enzyme with the Active Center Base Arrangement of Long-Chain Acyl-CoA Dehydrogenaseâ€. Biochemistry, 1996, 35, 12402-12411.	2.5	54
41	Protein Misfolding, Aggregation, and Degradation in Disease<SUP>. Molecular Biotechnology, 2005, 31, 141-150.	2.4	54
42	Defective folding and rapid degradation of mutant proteins is a common disease mechanism in genetic disorders. Journal of Inherited Metabolic Disease, 2000, 23, 441-447.	3.6	52
43	A novel mutation in the HSPD1 gene in a patient with hereditary spastic paraplegia. Journal of Neurology, 2007, 254, 897-900.	3.6	51
44	A Cellular Viability Assay to Monitor Drug Toxicity. Methods in Molecular Biology, 2010, 648, 303-311.	0.9	51
45	The Hsp60 folding machinery is crucial for manganese superoxide dismutase folding and function. Free Radical Research, 2014, 48, 168-179.	3.3	50
46	Rapid Degradation of Short-chain Acyl-CoA Dehydrogenase Variants with Temperature-sensitive Folding Defects Occurs after Import into Mitochondria. Journal of Biological Chemistry, 1998, 273, 13065-13071.	3.4	48
47	Quantitative Proteomics Reveals Cellular Targets of Celastrol. PLoS ONE, 2011, 6, e26634.	2.5	48
48	Human ClpP protease: cDNA sequence, tissue-specific expression and chromosomal assignment of the gene. FEBS Letters, 1995, 377, 249-252.	2.8	47
49	Grp78 Is Involved in Retention of Mutant Low Density Lipoprotein Receptor Protein in the Endoplasmic Reticulum. Journal of Biological Chemistry, 2000, 275, 33861-33868.	3.4	47
50	Clinical and genetic characteristics of Â cardiac actin gene mutations in hypertrophic cardiomyopathy. Journal of Medical Genetics, 2004, 41, 10e-10.	3.2	46
51	Disease-Associated Mutations in the HSPD1 Gene Encoding the Large Subunit of the Mitochondrial HSP60/HSP10 Chaperonin Complex. Frontiers in Molecular Biosciences, 2016, 3, 49.	3.5	46
52	Late onset motoneuron disorder caused by mitochondrial Hsp60 chaperone deficiency in mice. Neurobiology of Disease, 2013, 54, 12-23.	4.4	44
53	Proteomics of human mitochondria. Mitochondrion, 2017, 33, 2-14.	3.4	44
54	Heat-Shock Protein 70 Genes and Human Longevity: A View from Denmark. Annals of the New York Academy of Sciences, 2006, 1067, 301-308.	3.8	43

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55	Molecular Chaperone Disorders: Defective Hsp60 in Neurodegeneration. Current Topics in Medicinal Chemistry, 2013, 12, 2491-2503.	2.1	43
56	Structural organization of the human short-chain acyl-CoA dehydrogenase gene. Mammalian Genome, 1997, 8, 922-926.	2.2	42
57	Electron transfer flavoprotein and its role in mitochondrial energy metabolism in health and disease. Gene, 2021, 776, 145407.	2.2	42
58	The mutational spectrum in very long-chain acyl-CoA dehydrogenase deficiency. Journal of Inherited Metabolic Disease, 1996, 19, 169-172.	3.6	40
59	Assessing the relative importance of the biophysical properties of amino acid substitutions associated with human genetic disease. Human Mutation, 2002, 20, 98-109.	2.5	39
60	Enhanced genome editing in mammalian cells with a modified dual-fluorescent surrogate system. Cellular and Molecular Life Sciences, 2016, 73, 2543-2563.	5.4	39
61	Effects of a Mutation in the HSPE1 Gene Encoding the Mitochondrial Co-chaperonin HSP10 and Its Potential Association with a Neurological and Developmental Disorder. Frontiers in Molecular Biosciences, 2016, 3, 65.	3.5	38
62	Prevalent mutations in fatty acid oxidation disorders: diagnostic considerations. European Journal of Pediatrics, 2000, 159, S213-S218.	2.7	37
63	Mitochondrial proteomics on human fibroblasts for identification of metabolic imbalance and cellular stress. Proteome Science, 2009, 7, 20.	1.7	37
64	Misfolding of short-chain acyl-CoA dehydrogenase leads to mitochondrial fission and oxidative stress. Molecular Genetics and Metabolism, 2010, 100, 155-162.	1.1	37
65	Human and mouse mitochondrial orthologs of bacterial ClpX. Mammalian Genome, 2000, 11, 899-905.	2.2	36
66	Down-regulation of Hsp60 expression by RNAi impairs folding of medium-chain acyl-CoA dehydrogenase wild-type and disease-associated proteins. Molecular Genetics and Metabolism, 2005, 85, 260-270.	1.1	36
67	Cofactors and metabolites as potential stabilizers of mitochondrial acyl-CoA dehydrogenases. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2011, 1812, 1658-1663.	3.8	36
68	Truncating Plakophilin-2 Mutations in Arrhythmogenic Cardiomyopathy Are Associated With Protein Haploinsufficiency in Both Myocardium and Epidermis. Circulation: Cardiovascular Genetics, 2014, 7, 230-240.	5.1	36
69	Expression of transforming growth factor alpha and epidermal growth factor receptor in human bladder cancer. Scandinavian Journal of Clinical and Laboratory Investigation, 1999, 59, 267-277.	1.2	35
70	The C-terminal N-glycosylation sites of the human Â1,3/4-fucosyltransferase III, -V, and -VI (hFucTIII, -V and) Tj ETQq0 0 0 rgBT /Overlock	2.5	35
71	Metformin targets brown adipose tissue in vivo and reduces oxygen consumption in vitro. Diabetes, Obesity and Metabolism, 2018, 20, 2264-2273.	4.4	35
72	MCAD deficiency in Denmark. Molecular Genetics and Metabolism, 2012, 106, 175-188.	1.1	33

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73	Identification of Elements That Dictate the Specificity of Mitochondrial Hsp60 for Its Co-Chaperonin. PLoS ONE, 2012, 7, e50318.	2.5	32
74	Protein expression studies of desmoplakin mutations in cardiomyopathy patients reveal different molecular disease mechanisms. Clinical Genetics, 2013, 84, 20-30.	2.0	32
75	Mutated Desmoglein-2 Proteins are Incorporated into Desmosomes and Exhibit Dominant-Negative Effects in Arrhythmogenic Right Ventricular Cardiomyopathy. Human Mutation, 2013, 34, 697-705.	2.5	30
76	The Y42H mutation in medium-chain acyl-CoA dehydrogenase, which is prevalent in babies identified by MS/MS-based newborn screening, is temperature sensitive. FEBS Journal, 2004, 271, 4053-4063.	0.2	29
77	Association Between Low Self-Rated Health and Heterozygosity for -110A > C Polymorphism in the Promoter Region of HSP70-1 in Aged Danish Twins. Biogerontology, 2004, 5, 169-176.	3.9	29
78	Single-nucleotide variations in the genes encoding the mitochondrial Hsp60/Hsp10 chaperone system and their disease-causing potential. Journal of Human Genetics, 2007, 52, 56-65.	2.3	29
79	Biochemical Characterization of Purified, Human Recombinant Lys304Glu Medium-Chain Acyl-Coa Dehydrogenase Containing the Common Disease-Causing Mutation and Comparison with the Normal Enzyme. FEBS Journal, 1997, 246, 548-556.	0.2	27
80	APD-Containing Cyclolipodepsipeptides Target Mitochondrial Function in Hypoxic Cancer Cells. Cell Chemical Biology, 2018, 25, 1337-1349.e12.	5.2	27
81	Expression of wild-type and mutant medium-chain acyl-CoA dehydrogenase (MCAD) cDNA in eucaryotic cells. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1992, 1180, 65-72.	3.8	26
82	Prenatal diagnosis of medium-chain acyl-CoA dehydrogenase (MCAD) deficiency in a family with a previous fatal case of sudden unexpected death in childhood. Prenatal Diagnosis, 1995, 15, 82-86.	2.3	24
83	Proteomic investigation of cultivated fibroblasts from patients with mitochondrial short-chain acyl-CoA dehydrogenase deficiency. Molecular Genetics and Metabolism, 2014, 111, 360-368.	1.1	24
84	Anti-Inflammatory Heat Shock Protein 70 Genes are Positively Associated with Human Survival. Current Pharmaceutical Design, 2010, 16, 796-801.	1.9	23
85	The LMNA mutation p.Arg321Ter associated with dilated cardiomyopathy leads to reduced expression and a skewed ratio of lamin A and lamin C proteins. Experimental Cell Research, 2013, 319, 3010-3019.	2.6	23
86	Comparison between medium-chain acyl-CoA dehydrogenase mutant proteins overexpressed in bacterial and mammalian cells. Human Mutation, 1995, 6, 226-231.	2.5	21
87	A Polymorphic Variant in the Human Electron Transfer Flavoprotein Î±-Chain (Î±-T171) Displays Decreased Thermal Stability and Is Overrepresented in Very-Long-Chain acyl-CoA Dehydrogenase-Deficient Patients with Mild Childhood Presentation. Molecular Genetics and Metabolism, 1999, 67, 138-147.	1.1	21
88	Mutational hotspots in electron transfer flavoprotein underlie defective folding and function in multiple acyl-CoA dehydrogenase deficiency. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2010, 1802, 1070-1077.	3.8	21
89	Do lamin A and lamin C have unique roles?. Chromosoma, 2015, 124, 1-12.	2.2	21
90	Mitochondrial Spare Respiratory Capacity Is Negatively Correlated with Nuclear Reprogramming Efficiency. Stem Cells and Development, 2017, 26, 166-176.	2.1	21

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91	Amino acid polymorphism (Gly209Ser) in the ACADS gene. Human Molecular Genetics, 1994, 3, 1711-1711.	2.9	19
92	Ethylmalonic Encephalopathy ETHE1 R163W/R163Q Mutations Alter Protein Stability and Redox Properties of the Iron Centre. PLoS ONE, 2014, 9, e107157.	2.5	19
93	An inventory of interactors of the human HSP60/HSP10 chaperonin in the mitochondrial matrix space. Cell Stress and Chaperones, 2020, 25, 407-416.	2.9	18
94	Sequence variants in SPAST, SPG3A and HSPD1 in hereditary spastic paraplegia. Journal of the Neurological Sciences, 2009, 284, 90-95.	0.6	17
95	A polymorphic position in electron transfer flavoprotein modulates kinetic stability as evidenced by thermal stress. FEBS Letters, 2011, 585, 505-510.	2.8	16
96	The adsorption protein of phage IKe. Localization by deletion mutagenesis of domains involved in infectivity. Molecular Microbiology, 1992, 6, 471-478.	2.5	15
97	A cell model to study different degrees of Hsp60 deficiency in HEK293 cells. Cell Stress and Chaperones, 2011, 16, 633-640.	2.9	14
98	Deficiency of the mitochondrial sulfide regulator ETHE1 disturbs cell growth, glutathione level and causes proteome alterations outside mitochondria. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 126-135.	3.8	14
99	Release of periplasmic proteins induced inE. coliby expression of an N-terminal proximal segment of the phage fd gene 3 protein. FEBS Letters, 1991, 280, 27-31.	2.8	13
100	Glycosylation of the N-terminal potential N-glycosylation sites in the human alpha1,3-fucosyltransferase V and -VI (hFucTV and -VI). Glycoconjugate Journal, 2000, 17, 859-865.	2.7	13
101	Characterization of Wild-Type Human Medium-Chain Acyl-CoA Dehydrogenase (MCAD) and Mutant Enzymes Present in MCAD-Deficient Patients by Two-Dimensional Gel Electrophoresis: Evidence for Posttranslational Modification of the Enzyme. Biochemical Medicine and Metabolic Biology, 1994, 52, 36-44.	0.7	12
102	Impaired Folding and Subunit Assembly as Disease Mechanism: The Example of Medium-Chain acyl-CoA Dehydrogenase Deficiency. Progress in Molecular Biology and Translational Science, 1997, 58, 301-337.	1.9	12
103	Differential degradation of variant medium-chain acyl-CoA dehydrogenase by the protein quality control proteases Lon and ClpXP. Biochemical and Biophysical Research Communications, 2005, 333, 1160-1170.	2.1	12
104	Selected reaction monitoring as an effective method for reliable quantification of diseaseâ€associated proteins in maple syrup urine disease. Molecular Genetics & Genomic Medicine, 2014, 2, 383-392.	1.2	12
105	The clinical outcome of LMNA missense mutations can be associated with the amount of mutated protein in the nuclear envelope. European Journal of Heart Failure, 2018, 20, 1404-1412.	7.1	12
106	Characterization of a disease-causing Lys329 to Glu mutation in 16 patients with medium-chain Acyl-CoA dehydrogenase deficiency. Journal of Inherited Metabolic Disease, 1991, 14, 314-316.	3.6	11
107	Molecular genetic characterization and urinary excretion pattern of metabolites in two families with MCAD deficiency due to compound heterozygosity with a 13 base pair insertion in one allele. Journal of Inherited Metabolic Disease, 1994, 17, 169-184.	3.6	11
108	Heterozygosity for an inâ€frame deletion causes glutarylâ€CoA dehydrogenase deficiency in a patient detected by newborn screening: investigation of the effect of the mutant allele. Journal of Inherited Metabolic Disease, 2012, 35, 787-796.	3.6	9

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109	NMR-Based Metabonomic Investigation of Heat Stress in Myotubes Reveals a Time-Dependent Change in the Metabolites. Journal of Agricultural and Food Chemistry, 2010, 58, 6376-6386.	5.2	8
110	A Cell Model for HSP60 Deficiencies: Modeling Different Levels of Chaperonopathies Leading to Oxidative Stress and Mitochondrial Dysfunction. Methods in Molecular Biology, 2019, 1873, 225-239.	0.9	7
111	Metabolic profiling of heat or anoxic stress in mouse C2C12 myotubes using multinuclear magnetic resonance spectroscopy. Metabolism: Clinical and Experimental, 2010, 59, 814-823.	3.4	6
112	Biochemical characterization of a variant human medium-chain acyl-CoA dehydrogenase with a disease-associated mutation localized in the active site. Biochemical Journal, 1999, 337, 225.	3.7	5
113	Characterization of mouse Clpp protease cDNA, gene, and protein. Mammalian Genome, 2000, 11, 275-280.	2.2	5
114	Oxidative Stress-Induced Metabolic Changes in Mouse C2C12 Myotubes Studied with High-Resolution ¹³C, ¹H, and ³¹P NMR Spectroscopy. Journal of Agricultural and Food Chemistry, 2010, 58, 1918-1926.	5.2	4
115	Application of an Image Cytometry Protocol for Cellular and Mitochondrial Phenotyping on Fibroblasts from Patients with Inherited Disorders. JIMD Reports, 2015, 27, 17-26.	1.5	4
116	Investigation of Folding and Degradation of In Vitro Synthesized Mutant Proteins in Mitochondria. , 2003, 232, 285-294.		3
117	Optimized High-Contrast Brightfield Microscopy Application for Noninvasive Proliferation Assays of Human Cell Cultures. Assay and Drug Development Technologies, 2020, 18, 215-225.	1.2	3
118	Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency due to heterozygosity for the common mutation and an allele resulting in low levels of MCAD mRNA. Journal of Inherited Metabolic Disease, 1994, 17, 275-278.	3.6	2
119	Measuring Consequences of Protein Misfolding and Cellular Stress Using OMICS Techniques. Methods in Molecular Biology, 2010, 648, 119-135.	0.9	2
120	77 Mutations of Human Medium-Chain Acyl-CoA Dehydrogenase. Biochemical Society Transactions, 1998, 26, S65-S65.	3.4	1
121	Biochemical Characterisation of Mutations of Human Medium-Chain Acyl-CoA Dehydrogenase. , 1999, 466, 387-393.		1
122	Basic Introduction to In Vivo Protein Folding and Its Defects. , 2003, 232, 17-26.		1
123	Mitochondrial Hsp70 and the troubles of nomenclature: leaving behind tradition to gain intuitiveness and clarity. Cell Stress and Chaperones, 2016, 21, 547-551.	2.9	0