Fiorella Piemonte

List of Publications by Year
in descending order

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139 papers	7,476 citations	61857 43 h-index	56606 83 g-index
142 all docs	142 docs citations	142 times ranked	11077 citing authors

#	Article	IF	Citations
1	Personalized profiles of antioxidant signaling pathway in patients with tuberculosis. Journal of Microbiology, Immunology and Infection, 2022, 55, 405-412.	1.5	3
2	The Effects of Exercise Training on Cardiopulmonary Exercise Testing and Cardiac Biomarkers in Adult Patients with Hypoplastic Left Heart Syndrome and Fontan Circulation. Journal of Cardiovascular Development and Disease, 2022, 9, 171.	0.8	8
3	The Nrf2 induction prevents ferroptosis in Friedreich's Ataxia. Redox Biology, 2021, 38, 101791.	3.9	78
4	Nuclear Factor Erythroid 2-Related Factor 2 Activation Might Mitigate Clinical Symptoms in Friedreichâ€™s Ataxia: Clues of an â€œOut-Brain Originâ€•of the Disease From a Family Study. Frontiers in Neuroscience, 2021, 15, 638810.	1.4	5
5	Redox Homeostasis in Muscular Dystrophies. Cells, 2021, 10, 1364.	1.8	16
6	Imbalance of Systemic Redox Biomarkers in Children with Epilepsy: Role of Ferroptosis. Antioxidants, 2021, 10, 1267.	2.2	18
7	Molecular and histological traits of reduced lysosomal acid lipase activity in the fatty liver. Cell Death and Disease, 2021, 12, 1092.	2.7	5
8	Systemic Activation of Nrf2 Pathway in Parkinson's Disease. Movement Disorders, 2020, 35, 180-184.	2.2	66
9	Antioxidant Amelioration of Riboflavin Transporter Deficiency in Motoneurons Derived from Patient-Specific Induced Pluripotent Stem Cells. International Journal of Molecular Sciences, 2020, 21, 7402.	1.8	8
10	Plasma methylcitric acid and its correlations with other disease biomarkers: The impact in the follow up of patients with propionic and methylmalonic acidemia. Journal of Inherited Metabolic Disease, 2020, 43, 1173-1185.	1.7	19
11	Ferroptosis in Friedreichâ€™s Ataxia: A Metal-Induced Neurodegenerative Disease. Biomolecules, 2020, 10, 1551.	1.8	21
12	Oxidative Stress in DNA Repeat Expansion Disorders: A Focus on NRF2 Signaling Involvement. Biomolecules, 2020, 10, 702.	1.8	17
13	Compound heterozygosity for an expanded (GAA) and a (GAAGGA) repeat at FXN locus: from a diagnostic pitfall to potential clues to the pathogenesis of Friedreich ataxia. Neurogenetics, 2020, 21, 279-287.	0.7	2
14	The NRF2 Signaling Network Defines Clinical Biomarkers and Therapeutic Opportunity in Friedreichâ€™s Ataxia. International Journal of Molecular Sciences, 2020, 21, 916.	1.8	27
15	Frataxin deficiency induces lipid accumulation and affects thermogenesis in brown adipose tissue. Cell Death and Disease, 2020, 11, 51.	2.7	47
16	Nrf2 Induction Re-establishes a Proper Neuronal Differentiation Program in Friedreichâ€™s Ataxia Neural Stem Cells. Frontiers in Cellular Neuroscience, 2019, 13, 356.	1.8	36
17	Targeting NRF2 for the Treatment of Friedreichâ€™s Ataxia: A Comparison among Drugs. International Journal of Molecular Sciences, 2019, 20, 5211.	1.8	45
18	Serum uric acid in Friedreich Ataxia. Clinical Biochemistry, 2018, 54, 139-141.	0.8	7

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19	Progressive reduction of blood lysosomal acid lipase activity according to stage of adult chronic liver disease and altered enzymatic cellular distribution in cirrhosis. Digestive and Liver Disease, 2018, 50, 37.	0.4	0
20	ISCA1 mutation in a patient with infantile-onset leukodystrophy causes defects in mitochondrial [4Feâ€“4S] proteins. Human Molecular Genetics, 2018, 27, 3650-3650.	1.4	6
21	ISCA1 mutation in a patient with infantile-onset leukodystrophy causes defects in mitochondrial [4Feâ€“4S] proteins. Human Molecular Genetics, 2018, 27, 2739-2754.	1.4	25
22	A novel mutation in <scp>NDUFB11</scp> unveils a new clinical phenotype associated with lactic acidosis and sideroblastic anemia. Clinical Genetics, 2017, 91, 441-447.	1.0	24
23	Platelet count may impact on lysosomal acid lipase activity determination in dried blood spot. Clinical Biochemistry, 2017, 50, 726-728.	0.8	14
24	Oxidative stress in Duchenne muscular dystrophy: focus on the NRF2 redox pathway. Human Molecular Genetics, 2017, 26, 2781-2790.	1.4	71
25	Novel mutations in <scp>KARS</scp> cause hypertrophic cardiomyopathy and combined mitochondrial respiratory chain defect. Clinical Genetics, 2017, 91, 918-923.	1.0	27
26	<scp>DJ</scp>â€1 modulates mitochondrial response to oxidative stress: clues from a novel diagnosis of <scp>PARK7</scp>. Clinical Genetics, 2017, 92, 18-25.	1.0	34
27	Nrf2-Inducers Counteract Neurodegeneration in Frataxin-Silenced Motor Neurons: Disclosing New Therapeutic Targets for Friedreichâ€™s Ataxia. International Journal of Molecular Sciences, 2017, 18, 2173.	1.8	58
28	The cytoskeletal arrangements necessary to neurogenesis. Oncotarget, 2016, 7, 19414-19429.	0.8	44
29	Biallelic Mutations in TBCD , Encoding the Tubulin Folding Cofactor D, Perturb Microtubule Dynamics and Cause Early-Onset Encephalopathy. American Journal of Human Genetics, 2016, 99, 962-973.	2.6	66
30	Liver Cirrhosis is Characterized by an Acquired Lysosomal Acid Lipase Deficiency Independent from the Etiology of Liver Disease. Journal of Hepatology, 2016, 64, S290.	1.8	0
31	Frataxin silencing alters microtubule stability in motor neurons: implications for Friedreichâ€™s ataxia. Human Molecular Genetics, 2016, 25, 4288-4301.	1.4	27
32	Cytoskeletal dynamics during in vitro neurogenesis of induced pluripotent stem cells (iPSCs). Molecular and Cellular Neurosciences, 2016, 77, 113-124.	1.0	9
33	Experimental violation of Bell inequalities for multi-dimensional systems. Scientific Reports, 2016, 6, 22088.	1.6	16
34	Lysosomal acid lipase activity is reduced in patients with cirrhosis and associated with surrogate indices of portal hypertension. Digestive and Liver Disease, 2016, 48, e34.	0.4	0
35	LYRM7mutations cause a multifocal cavitating leukoencephalopathy with distinct MRI appearance. Brain, 2016, 139, 782-794.	3.7	51
36	High concentrations of H2O2 trigger hypertrophic cascade and phosphatase and tensin homologue (PTEN) glutathionylation in H9c2 cardiomyocytes. Experimental and Molecular Pathology, 2016, 100, 199-206.	0.9	7

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37	Activation of the transcription factor EB rescues lysosomal abnormalities in cystinotic kidney cells. Kidney International, 2016, 89, 862-873.	2.6	85
38	A novel AIFM1 mutation expands the phenotype to an infantile motor neuron disease. European Journal of Human Genetics, 2016, 24, 463-466.	1.4	51
39	Lysosomal Acid Lipase Activity Is Reduced Both in Cryptogenic Cirrhosis and in Cirrhosis of Known Etiology. PLoS ONE, 2016, 11, e0156113.	1.1	21
40	Systemic Redox Biomarkers in Neurodegenerative Diseases. Current Drug Metabolism, 2015, 16, 46-70.	0.7	6
41	P1049 : Lysosomal Acid Lipase activity in patients with non-alcoholic fatty liver disease. Journal of Hepatology, 2015, 62, S742.	1.8	1
42	Protracted late infantile ceroid lipofuscinosis due to TPP1 mutations: Clinical, molecular and biochemical characterization in three sibs. Journal of the Neurological Sciences, 2015, 356, 65-71.	0.3	27
43	Additive effect of nuclear and mitochondrial mutations in a patient with mitochondrial encephalomyopathy. Human Molecular Genetics, 2015, 24, 3248-3256.	1.4	18
44	Reduced Lysosomal Acid Lipase Activity in Adult Patients With Non-alcoholic Fatty Liver Disease. EBioMedicine, 2015, 2, 750-754.	2.7	51
45	Glutathione imbalance in blood of patients with Duchenne muscular dystrophy. Neuromuscular Disorders, 2015, 25, S250.	0.3	0
46	Launching the first clinical trial in SEPN1-related myopathy: The SELNAC study. Neuromuscular Disorders, 2015, 25, S270.	0.3	1
47	Intracellular Distribution of Glutathionylated Proteins in Cultured Dermal Fibroblasts by Immunofluorescence. Methods in Molecular Biology, 2015, 1208, 395-408.	0.4	2
48	Friedreichâ€™s Ataxia: A Neuronal Point of View on the Oxidative Stress Hypothesis. Antioxidants, 2014, 3, 592-603.	2.2	8
49	Frataxin Silencing Inactivates Mitochondrial Complex I in NSC34 Motoneuronal Cells and Alters Glutathione Homeostasis. International Journal of Molecular Sciences, 2014, 15, 5789-5806.	1.8	22
50	Glutathione metabolism in cobalamin deficiency type C (cblC). Journal of Inherited Metabolic Disease, 2014, 37, 125-129.	1.7	46
51	G.P.209. Neuromuscular Disorders, 2014, 24, 879-880.	0.3	1
52	Riboflavin responsive mitochondrial myopathy is a new phenotype of dihydrolipoamide dehydrogenase deficiency. The chaperon-like effect of vitamin B2. Mitochondrion, 2014, 18, 49-57.	1.6	39
53	Effects of levosimendan on mitochondrial function in patients withÂseptic shock: A randomized trial. Biochimie, 2014, 102, 166-173.	1.3	41
54	Pontocerebellar hypoplasia type 6 caused by mutations in RARS2: definition of the clinical spectrum and molecular findings in five patients. Journal of Inherited Metabolic Disease, 2013, 36, 43-53.	1.7	70

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55	Glutathione: A redox signature in monitoring EPI-743 therapy in children with mitochondrial encephalomyopathies. Molecular Genetics and Metabolism, 2013, 109, 208-214.	0.5	49
56	Glutathione imbalance in patients with X-linked adrenoleukodystrophy. Molecular Genetics and Metabolism, 2013, 109, 366-370.	0.5	39
57	Frataxin Deficiency Leads to Reduced Expression and Impaired Translocation of NF-E2-Related Factor (Nrf2) in Cultured Motor Neurons. International Journal of Molecular Sciences, 2013, 14, 7853-7865.	1.8	75
58	Protein Glutathionylation in Cardiovascular Diseases. International Journal of Molecular Sciences, 2013, 14, 20845-20876.	1.8	81
59	Emodin Prevents Intrahepatic Fat Accumulation, Inflammation and Redox Status Imbalance During Diet-Induced Hepatosteatosis in Rats. International Journal of Molecular Sciences, 2012, 13, 2276-2289.	1.8	48
60	Protein glutathionylation in cellular compartments: A constitutive redox signal. Redox Report, 2012, 17, 63-71.	1.4	8
61	EPI-743 reverses the progression of the pediatric mitochondrial diseaseâ€"Genetically defined Leigh Syndrome. Molecular Genetics and Metabolism, 2012, 107, 383-388.	0.5	163
62	Glutathione Status in MMACHC Patients. Free Radical Biology and Medicine, 2012, 53, S69-S70.	1.3	0
63	TMEM70: a mutational hot spot in nuclear ATP synthase deficiency with a pivotal role in complex V biogenesis. Neurogenetics, 2012, 13, 375-386.	0.7	25
64	Intermittentâ€relapsing pyruvate dehydrogenase complex deficiency: a case with clinical, biochemical, and neuroradiological reversibility. Developmental Medicine and Child Neurology, 2012, 54, 472-476.	1.1	18
65	Pediatric reference intervals for muscle coenzyme Q10. Biomarkers, 2012, 17, 764-766.	0.9	2
66	Brownâ€"Vialettoâ€"van Laere and Fazioâ€"Londe overlap syndromes: A clinical, biochemical and genetic study. Neuromuscular Disorders, 2012, 22, 1075-1082.	0.3	36
67	The use of muscle biopsy in the diagnosis of undefined ataxia with cerebellar atrophy in children. European Journal of Paediatric Neurology, 2012, 16, 248-256.	0.7	39
68	S-Glutathionylation signaling in cell biology: Progress and prospects. European Journal of Pharmaceutical Sciences, 2012, 46, 279-292.	1.9	152
69	Synergistic phage-antibiotic combinations for the control ofEscherichia colibiofilmsin vitro. FEMS Immunology and Medical Microbiology, 2012, 65, 395-398.	2.7	182
70	Redox homeostasis and posttranslational modifications/activity of phosphatase and tensin homolog in hepatocytes from rats with diet-induced hepatosteatosis. Journal of Nutritional Biochemistry, 2012, 23, 169-178.	1.9	14
71	All glutathione forms are depleted in blood of obese and type 1 diabetic children. Pediatric Diabetes, 2012, 13, 272-277.	1.2	23
72	Novel large-range mitochondrial DNA deletions and fatal multisystemic disorder with prominent hepatopathy. Biochemical and Biophysical Research Communications, 2011, 415, 300-304.	1.0	7

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73	Effect of protein glutathionylation on neuronal cytoskeleton: a potential link to neurodegeneration. Neuroscience, 2011, 192, 285-294.	1.1	29
74	P3.1 Brownâ€“Vialettoâ€“Van Laere and Fazio Londe overlap sindromes: A clinical, biochemical and genetic study in 6 patients. Neuromuscular Disorders, 2011, 21, 682.	0.3	1
75	Mirnome analysis reveals novel molecular determinants in the pathogenesis of diet-induced nonalcoholic fatty liver disease. Laboratory Investigation, 2011, 91, 283-293.	1.7	176
76	Progressive cavitating leukoencephalopathy associated with respiratory chain complex I deficiency and a novel mutation in NDUFS1. Neurogenetics, 2011, 12, 9-17.	0.7	43
77	Neuroprotection: The Emerging Concept of Restorative Neural Stem Cell Biology for the Treatment of Neurodegenerative Diseases. Current Neuropharmacology, 2011, 9, 313-317.	1.4	53
78	Susceptibility of isolated myofibrils to in vitro glutathionylation: Potential relevance to muscle functions. Cytoskeleton, 2010, 67, 81-89.	1.0	20
79	Endotoxin and Plasminogen Activator Inhibitorâ€1 Serum Levels Associated With Nonalcoholic Steatohepatitis in Children. Journal of Pediatric Gastroenterology and Nutrition, 2010, 50, 645-649.	0.9	126
80	Oxidative stress parameters in paediatric non-alcoholic fatty liver disease. International Journal of Molecular Medicine, 2010, 26, 471-6.	1.8	78
81	GSSG-mediated Complex I defect in isolated cardiac mitochondria. International Journal of Molecular Medicine, 2010, 26, 95-9.	1.8	26
82	Assaying ATP synthesis in cultured cells: A valuable tool for the diagnosis of patients with mitochondrial disorders. Biochemical and Biophysical Research Communications, 2009, 383, 58-62.	1.0	26
83	Late-onset MNGIE without peripheral neuropathy due to incomplete loss of thymidine phosphorylase activity. Neuromuscular Disorders, 2009, 19, 837-840.	0.3	34
84	Variant Late Infantile Neuronal Ceroid Lipofuscinosis Because of CLN1 Mutations. Pediatric Neurology, 2009, 40, 271-276.	1.0	29
85	Friedreich's ataxia: Oxidative stress and cytoskeletal abnormalities. Journal of the Neurological Sciences, 2009, 287, 111-118.	0.3	75
86	694 MICRORNA EXPRESSION PROFILES IN LIVER TISSUES FROM RAT FED HIGH FAT/HIGH CARBOYDRATE DIET MAY HELP TO ELUCIDATE MOLECULAR PATHOGENESIS OF NON-ALCOHOLIC FATTY LIVER DISEASE. Journal of Hepatology, 2009, 50, S254-S255.	1.8	0
87	695 EMODIN PROTECTS PRIMARY RAT HEPATOCYTES FROM PRO-OXIDATIVE EFFECTS AND AKT PATHWAY DYSREGULATION INDUCED BY A HIGH-FAT/LOW CARBOHYDRATE DIET. Journal of Hepatology, 2009, 50, S255.	1.8	0
88	Glutathionylation of p65NF-Î²B correlates with proliferating/apoptotic hepatoma cells exposed to pro- and anti-oxidants. International Journal of Molecular Medicine, 2009, 24, 319-26.	1.8	18
89	Myosin as a potential redox-sensor: an inÂvitro study. Journal of Muscle Research and Cell Motility, 2008, 29, 119-126.	0.9	37
90	Lifestyle intervention and antioxidant therapy in children with nonalcoholic fatty liver disease: A randomized, controlled trial. Hepatology, 2008, 48, 119-128.	3.6	362

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91	Protein glutathionylation increases in the liver of patients with nonâ€alcoholic fatty liver disease. Journal of Gastroenterology and Hepatology (Australia), 2008, 23, e457-64.	1.4	26
92	Waist circumference correlates with liver fibrosis in children with non-alcoholic steatohepatitis. Gut, 2008, 57, 1283-1287.	6.1	123
93	Collapsing glomerulopathy associated with inherited mitochondrial injury. Kidney International, 2008, 74, 237-243.	2.6	31
94	COQ2 Nephropathy. Journal of the American Society of Nephrology: JASN, 2007, 18, 2773-2780.	3.0	297
95	Chronic GM2 gangliosidosis type Sandhoff associated with a novel missense HEXB gene mutation causing a double pathogenic effect. Molecular Genetics and Metabolism, 2007, 91, 111-114.	0.5	14
96	SUCLA2 mutations are associated with mild methylmalonic aciduria, Leigh-like encephalomyopathy, dystonia and deafness. Brain, 2007, 130, 862-874.	3.7	180
97	Protein glutathionylation increases in liver of patients with nonalcoholic fatty liver disease (NAFLD). Digestive and Liver Disease, 2007, 39, A85.	0.4	0
98	Infantile Mitochondrial Disorders. Bioscience Reports, 2007, 27, 105-112.	1.1	10
99	Effect of vitamin E on aminotransferase levels and insulin resistance in children with non-alcoholic fatty liver disease. Alimentary Pharmacology and Therapeutics, 2006, 24, 1553-1561.	1.9	161
100	Novel CLN1 mutation in two Italian sibs with late infantile neuronal ceroid lipofuscinosis. European Journal of Paediatric Neurology, 2006, 10, 154-156.	0.7	15
101	NAFLD in children: A prospective clinical-pathological study and effect of lifestyle advice. Hepatology, 2006, 44, 458-465.	3.6	324
102	Protein glutathionylation in human central nervous system: Potential role in redox regulation of neuronal defense against free radicals. Journal of Neuroscience Research, 2006, 83, 256-263.	1.3	50
103	Impaired Activity of the Î³-Glutamyl Cycle in Nephropathic Cystinosis Fibroblasts. Pediatric Research, 2006, 59, 332-335.	1.1	43
104	Leptin, free leptin index, insulin resistance and liver fibrosis in children with non-alcoholic fatty liver disease. European Journal of Endocrinology, 2006, 155, 735-743.	1.9	91
105	Simultaneous determination of ubiquinol and ubiquinone in skeletal muscle of pediatric patients. Analytical Biochemistry, 2005, 342, 352-355.	1.1	15
106	Glutathione metabolism and antioxidant enzymes in patients affected by nonalcoholic steatohepatitis. Clinica Chimica Acta, 2005, 355, 105-111.	0.5	68
107	Axonal degeneration in paraplegin-deficient mice is associated with abnormal mitochondria and impairment of axonal transport. Journal of Clinical Investigation, 2004, 113, 231-242.	3.9	241
108	Determination of glutathionyl-hemoglobin in human erythrocytes by cation-exchange high-performance liquid chromatography. Analytical Biochemistry, 2003, 312, 85-90.	1.1	28

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109	Analysis of glutathione: implication in redox and detoxification. Clinica Chimica Acta, 2003, 333, 19-39.	0.5	931
110	Glutathione metabolism and antioxidant enzymes in children with down syndrome. Journal of Pediatrics, 2003, 142, 583-585.	0.9	56
111	Actin Glutathionylation Increases in Fibroblasts of Patients with Friedreich's Ataxia. Journal of Biological Chemistry, 2003, 278, 42588-42595.	1.6	142
112	Atypical Leigh syndrome associated with the D393N mutation in the mitochondrial ND5 subunit. Neurology, 2003, 61, 1017-1018.	1.5	29
113	Antioxidant enzymes in blood of patients with Friedreich's ataxia. Archives of Disease in Childhood, 2002, 86, 376-379.	1.0	59
114	Determination of superoxide dismutase and glutathione peroxidase activities in blood of healthy pediatric subjects. Clinica Chimica Acta, 2002, 322, 117-120.	0.5	34
115	Hypertrophic cardiomyopathy and mtDNA depletion. Successful treatment with heart transplantation. Neuromuscular Disorders, 2002, 12, 56-59.	0.3	30
116	Rapid determination of mycophenolic acid in plasma by reversed-phase high-performance liquid chromatography. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2002, 776, 251-254.	1.2	22
117	Respiratory chain defects in hereditary spastic paraplegias. Neuromuscular Disorders, 2001, 11, 565-569.	0.3	15
118	Determination of Blood Total, Reduced, and Oxidized Glutathione in Pediatric Subjects. Clinical Chemistry, 2001, 47, 1467-1469.	1.5	173
119	Glutathione in blood of patients with Friedreich's ataxia. European Journal of Clinical Investigation, 2001, 31, 1007-1011.	1.7	154
120	Fatal infantile leukodystrophy. Neurology, 2001, 57, 265-270.	1.5	64
121	Cytochrome c Oxidase-deficient Patients Have Distinct Subunit Assembly Profiles. Journal of Biological Chemistry, 2001, 276, 16296-16301.	1.6	36
122	The T9176G mtDNA mutation severely affects ATP production and results in Leigh syndrome. Neurology, 2001, 56, 687-690.	1.5	79
123	Novel 7-DHCR mutation in a child with Smith-Lemli-Opitz syndrome. , 2000, 91, 138-140.		13
124	Oxidative abnormalities in Menkes disease. Journal of Inherited Metabolic Disease, 2000, 23, 349-351.	1.7	10
125	A novel SURF1 mutation results in Leigh syndrome with peripheral neuropathy caused by cytochrome c oxidase deficiency. Neuromuscular Disorders, 2000, 10, 450-453.	0.3	37
126	Steatosis in children. A case report. Journal of Hepatology, 2000, 32, 217.	1.8	0

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127	OXPHOS and mtDNA alterations in a family with spastic paraparesis. Acta Neurologica Scandinavica, 2000, 101, 255-258.	1.0	1
128	A novel 7-DHCR mutation in a lebanese child with Smith-Lemli-Opitz syndrome. Atherosclerosis, 1999, 144, 21-22.	0.4	0
129	Alternative splicing of human plasma cholesteryl ester transfer protein mRNA in Caco-2 cells and its modulation by oleic acid. Molecular and Cellular Biochemistry, 1997, 177, 107-112.	1.4	11
130	The effects of pregnancy steroids on adaptation of beta cells to pregnancy involve the pancreatic glucose sensor glucokinase. Journal of Endocrinology, 1997, 155, 247-253.	1.2	13
131	Kinetic studies on rat liver microsomal glutathione transferase: consequences of activation. BBA - Proteins and Proteomics, 1995, 1247, 277-283.	2.1	22
132	Microsomal glutathione transferase: Lipid-derived substrates and lipid dependence. Archives of Biochemistry and Biophysics, 1995, 320, 210-216.	1.4	82
133	Colorimetric and Fluorometric Assays of Glutathione Transferase Based on 7-Chloro-4-nitrobenzo-2-oxa-1,3-diazole. Analytical Biochemistry, 1994, 218, 463-465.	1.1	142
134	Aggregation of pyrene-labeled microsomal glutathione S-transferase. Effect of concentration. FEBS Journal, 1993, 217, 661-663.	0.2	6
135	Investigation of the active site of human placenta glutathione transferase Ï€ by means of a spin-labelled glutathione analogue. BBA - Proteins and Proteomics, 1992, 1122, 265-268.	2.1	9
136	Interaction of hemin with placental glutathione transferase. FEBS Journal, 1990, 189, 493-497.	0.2	36
137	Effects of Clostridium Difficile toxins A and B on cytoskeleton organization in HEp-2 cells: A comparative morphological study. Toxicon, 1989, 27, 1209-1218.	0.8	54
138	Regulation of acyl CoA: Cholesterol acyl transferase (ACAT) activity by mevalonate and cholesterol in isolated rat hepatocytes during perinatal development. Bioscience Reports, 1986, 6, 735-740.	1.1	1
139	Short term regulation of acyl CoA: Cholesterol acyl transferase (ACAT) activity in the regenerating and perinatal liver. Bioscience Reports, 1985, 5, 237-242.	1.1	3