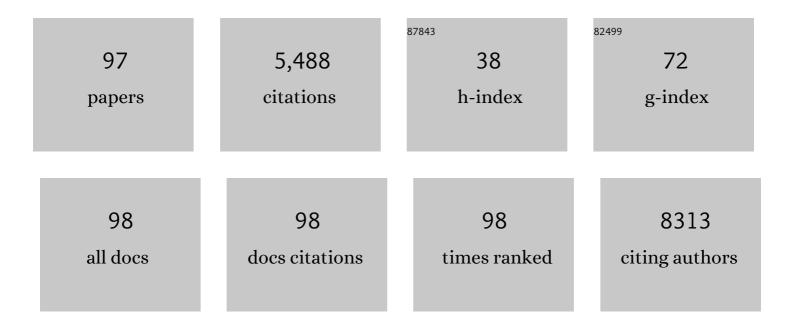
## Anna Pastore

List of Publications by Year in descending order

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ANNA DASTORE

#	Article	IF	CITATIONS
1	Analysis of glutathione: implication in redox and detoxification. Clinica Chimica Acta, 2003, 333, 19-39.	0.5	931
2	COQ2 Nephropathy. Journal of the American Society of Nephrology: JASN, 2007, 18, 2773-2780.	3.0	297
3	Fully automated assay for total homocysteine, cysteine, cysteinylglycine, glutathione, cysteamine, and 2-mercaptopropionylglycine in plasma and urine. Clinical Chemistry, 1998, 44, 825-832.	1.5	242
4	Determination of Blood Total, Reduced, and Oxidized Glutathione in Pediatric Subjects. Clinical Chemistry, 2001, 47, 1467-1469.	1.5	173
5	Extracorporeal dialysis in neonatal hyperammonemia: modalities and prognostic indicators. Pediatric Nephrology, 2001, 16, 862-867.	0.9	167
6	EPI-743 reverses the progression of the pediatric mitochondrial disease—Genetically defined Leigh Syndrome. Molecular Genetics and Metabolism, 2012, 107, 383-388.	0.5	163
7	Clutathione in blood of patients with Friedreich's ataxia. European Journal of Clinical Investigation, 2001, 31, 1007-1011.	1.7	154
8	S-Glutathionylation signaling in cell biology: Progress and prospects. European Journal of Pharmaceutical Sciences, 2012, 46, 279-292.	1.9	152
9	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
10	Actin Glutathionylation Increases in Fibroblasts of Patients with Friedreich's Ataxia. Journal of Biological Chemistry, 2003, 278, 42588-42595.	1.6	142
11	Targeting oxidative stress improves disease outcomes in a rat model of acquired epilepsy. Brain, 2019, 142, e39-e39.	3.7	137
12	A novel disorder involving dyshematopoiesis, inflammation, and HLH due to aberrant CDC42 function. Journal of Experimental Medicine, 2019, 216, 2778-2799.	4.2	132
13	Nitrosylation of Human Glutathione Transferase P1-1 with Dinitrosyl Diglutathionyl Iron Complex in Vitro and in Vivo. Journal of Biological Chemistry, 2005, 280, 42172-42180.	1.6	109
14	Site-directed Mutagenesis of Human Glutathione Transferase P1-1. Journal of Biological Chemistry, 1995, 270, 1243-1248.	1.6	87
15	Activation of the transcription factor EB rescues lysosomal abnormalities in cystinotic kidney cells. Kidney International, 2016, 89, 862-873.	2.6	85
16	Plasma Levels of Homocysteine and Cysteine Increased in Pediatric NAFLD and Strongly Correlated with Severity of Liver Damage. International Journal of Molecular Sciences, 2014, 15, 21202-21214.	1.8	84
17	Protein Glutathionylation in Cardiovascular Diseases. International Journal of Molecular Sciences, 2013, 14, 20845-20876.	1.8	81
18	Treatment of doxorubicin-resistant MCF7/Dx cells with nitric oxide causes histone glutathionylation and reversal of drug resistance. Biochemical Journal, 2011, 440, 175-183.	1.7	77

ANNA PASTORE

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19	Glutathione metabolism and antioxidant enzymes in patients affected by nonalcoholic steatohepatitis. Clinica Chimica Acta, 2005, 355, 105-111.	0.5	68
20	Long-term outcome of nephropathic cystinosis: a 20-year single-center experience. Pediatric Nephrology, 2010, 25, 2459-2467.	0.9	66
21	A new simple and rapid LC–ESI-MS/MS method for quantification of plasma oxysterols as dimethylaminobutyrate esters. Its successful use for the diagnosis of Niemann–Pick type C disease. Clinica Chimica Acta, 2014, 437, 93-100.	0.5	62
22	Antioxidant enzymes in blood of patients with Friedreich's ataxia. Archives of Disease in Childhood, 2002, 86, 376-379.	1.0	59
23	Nrf2-Inducers Counteract Neurodegeneration in Frataxin-Silenced Motor Neurons: Disclosing New Therapeutic Targets for Friedreich's Ataxia. International Journal of Molecular Sciences, 2017, 18, 2173.	1.8	58
24	Glutathione metabolism and antioxidant enzymes in children with down syndrome. Journal of Pediatrics, 2003, 142, 583-585.	0.9	56
25	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
26	Common mutation in methylenetetrahydrofolate reductase. Correlation with homocysteine and other risk factors for vascular disease. Atherosclerosis, 1998, 139, 377-383.	0.4	49
27	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
28	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
29	Endo-Lysosomal Dysfunction in Human Proximal Tubular Epithelial Cells Deficient for Lysosomal Cystine Transporter Cystinosin. PLoS ONE, 2015, 10, e0120998.	1.1	47
30	Glutathione metabolism in cobalamin deficiency type C (cblC). Journal of Inherited Metabolic Disease, 2014, 37, 125-129.	1.7	46
31	Altered mTOR signalling in nephropathic cystinosis. Journal of Inherited Metabolic Disease, 2016, 39, 457-464.	1.7	45
32	Cystinuria in children and young adults: success of monitoring free-cystine urine levels. Pediatric Nephrology, 2007, 22, 1869-1873.	0.9	44
33	Tissue Factor and Homocysteine Levels in Ischemic Heart Disease Are Associated with Angiographically Documented Clinical Recurrences after Coronary Angioplasty. Thrombosis and Haemostasis, 2000, 83, 826-832.	1.8	43
34	Impaired Activity of the Î <sup>3</sup> -Glutamyl Cycle in Nephropathic Cystinosis Fibroblasts. Pediatric Research, 2006, 59, 332-335.	1.1	43
35	Effects of levosimendan on mitochondrial function in patients withÂseptic shock: A randomized trial. Biochimie, 2014, 102, 166-173.	1.3	41
36	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

**ANNA PASTORE** 

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37	Glutathione imbalance in patients with X-linked adrenoleukodystrophy. Molecular Genetics and Metabolism, 2013, 109, 366-370.	0.5	39
38	Homocysteine, cysteine, folate and vitamin B12 status in type 2 diabetic patients with chronic kidney disease. Journal of Nephrology, 2015, 28, 571-576.	0.9	39
39	Myosin as a potential redox-sensor: an inÂvitro study. Journal of Muscle Research and Cell Motility, 2008, 29, 119-126.	0.9	37
40	Brown–Vialetto–van Laere and Fazio–Londe overlap syndromes: A clinical, biochemical and genetic study. Neuromuscular Disorders, 2012, 22, 1075-1082.	0.3	36
41	Cystinosis (ctns) zebrafish mutant shows pronephric glomerular and tubular dysfunction. Scientific Reports, 2017, 7, 42583.	1.6	36
42	Erythrocyte glutathione transferase: a potential new biomarker in chronic kidney diseases which correlates with plasma homocysteine. Amino Acids, 2012, 43, 347-354.	1.2	35
43	Conformational States of Human Placental Glutathione Transferase as Probed by Limited Proteolysis. Biochemical and Biophysical Research Communications, 1993, 194, 804-810.	1.0	34
44	Determination of superoxide dismutase and glutathione peroxidase activities in blood of healthy pediatric subjects. Clinica Chimica Acta, 2002, 322, 117-120.	0.5	34
45	The proteome of cblC defect: in vivo elucidation of altered cellular pathways in humans. Journal of Inherited Metabolic Disease, 2015, 38, 969-979.	1.7	34
46	Effects of Folic Acid Before and After Vitamin B12 on Plasma Homocysteine Concentrations in Hemodialysis Patients with Known MTHFR Genotypes. Clinical Chemistry, 2006, 52, 145-148.	1.5	33
47	Lack of association between carotid intima-media thickness and methylenetetrahydrofolate reductase gene polymorphism or serum homocysteine in non—insulin-dependent diabetes mellitus. Metabolism: Clinical and Experimental, 2000, 49, 718-723.	1.5	32
48	Erythrocyte glutathione transferase activity: a possible early biomarker for blood toxicity in uremic diabetic patients. Acta Diabetologica, 2014, 51, 219-224.	1.2	32
49	The fine-tuning of TRAF2–GSTP1-1 interaction: effect of ligand binding and in situ detection of the complex. Cell Death and Disease, 2014, 5, e1015-e1015.	2.7	31
50	Effect of protein glutathionylation on neuronal cytoskeleton: a potential link to neurodegeneration. Neuroscience, 2011, 192, 285-294.	1.1	29
51	Determination of glutathionyl-hemoglobin in human erythrocytes by cation-exchange high-performance liquid chromatography. Analytical Biochemistry, 2003, 312, 85-90.	1.1	28
52	Role of GST P1-1 in mediating the effect of etoposide on human neuroblastoma cell line Sh-Sy5y. Journal of Cellular Biochemistry, 2002, 86, 340-347.	1.2	27
53	Frataxin silencing alters microtubule stability in motor neurons: implications for Friedreich's ataxia. Human Molecular Genetics, 2016, 25, 4288-4301.	1.4	27
54	GSSG-mediated Complex I defect in isolated cardiac mitochondria. International Journal of Molecular Medicine, 2010, 26, 95-9.	1.8	26

ANNA PASTORE

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55	Modulation of CTNS gene expression by intracellular thiols. Free Radical Biology and Medicine, 2010, 48, 865-872.	1.3	26
56	Simultaneous determination of inulin and p-aminohippuric acid in plasma and urine by reversed-phase high-performance liquid chromatography. Biomedical Applications, 2001, 751, 187-191.	1.7	25
57	Serum homocysteine, methylenetetrahydrofolate reductase gene polymorphism and cardiovascular disease in heterozygous familial hypercholesterolemia. Atherosclerosis, 2005, 179, 333-338.	0.4	24
58	Pyroglutamic aciduria and nephropathic cystinosis. Journal of Inherited Metabolic Disease, 1999, 22, 224-226.	1.7	23
59	All glutathione forms are depleted in blood of obese and type 1 diabetic children. Pediatric Diabetes, 2012, 13, 272-277.	1.2	23
60	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
61	Characterization of a new trabectedinâ€resistant myxoid liposarcoma cell line that shows collateral sensitivity to methylating agents. International Journal of Cancer, 2012, 131, 59-69.	2.3	22
62	Creatine metabolism in urea cycle defects. Journal of Inherited Metabolic Disease, 2012, 35, 647-653.	1.7	22
63	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
64	Susceptibility of isolated myofibrils to in vitro glutathionylation: Potential relevance to muscle functions. Cytoskeleton, 2010, 67, 81-89.	1.0	20
65	Glutathione <scp>S</scp> â€transferase <scp>P</scp> 1â€1 as a target for mesothelioma treatment. Cancer Science, 2013, 104, 223-230.	1.7	20
66	High performance liquid chromatographic determination of plasma free and total tazobactam and piperacillin. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2009, 877, 86-88.	1.2	18
67	Glutathionylation of p65NF-κB correlates with proliferating/apoptotic hepatoma cells exposed to pro- and anti-oxidants. International Journal of Molecular Medicine, 2009, 24, 319-26.	1.8	18
68	Simultaneous determination of ubiquinol and ubiquinone in skeletal muscle of pediatric patients. Analytical Biochemistry, 2005, 342, 352-355.	1.1	15
69	Long-term renal function in heart transplant children on cyclosporine treatment. Pediatric Nephrology, 2006, 21, 561-565.	0.9	15
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	Intrinsic Bone Defects in Cystinotic Mice. American Journal of Pathology, 2019, 189, 1053-1064.	1.9	14
72	Selective Adsorption of Homocysteine Using an HFR-ON LINE Technique. Artificial Organs, 2004, 28, 592-595.	1.0	11

Anna Pastore

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73	Drug Repurposing in Rare Diseases: An Integrative Study of Drug Screening and Transcriptomic Analysis in Nephropathic Cystinosis. International Journal of Molecular Sciences, 2021, 22, 12829.	1.8	11
74	Optimizing the dose of hydroxocobalamin in cobalamin C (cblC) defect. Molecular Genetics and Metabolism, 2013, 109, 329-330.	0.5	10
75	Detection of iron deficiency in children with Down syndrome. Genetics in Medicine, 2020, 22, 317-325.	1.1	10
76	Serum homocysteine, MTHFR gene polymorphism, and carotid intimal-medial thickness in NIDDM subjects. Journal of Thrombosis and Thrombolysis, 1999, 8, 207-212.	1.0	9
77	Evaluation of carbohydrate-cysteamine thiazolidines as pro-drugs for the treatment of cystinosis. Carbohydrate Research, 2017, 439, 9-15.	1.1	9
78	Cystinosin-LKG rescues cystine accumulation and decreases apoptosis rate in cystinotic proximal tubular epithelial cells. Pediatric Research, 2017, 81, 113-119.	1.1	9
79	Protein glutathionylation in cellular compartments: A constitutive redox signal. Redox Report, 2012, 17, 63-71.	1.4	8
80	Semiautomated Method for Determination of Cystine Concentration in Polymorphonuclear Leukocytes. Clinical Chemistry, 2000, 46, 560-576.	1.5	8
81	Influence of dialysis techniques and alternate vitamin supplementation on homocysteine levels in patients with known MTHFR genotypes. Clinical and Experimental Nephrology, 2015, 19, 140-145.	0.7	7
82	Reverseâ€phase highâ€performance liquid chromatography for the simultaneous determination of sildenafil and <i>Nâ€</i> desmethyl sildenafil in plasma of children. Biomedical Chromatography, 2016, 30, 2070-2073.	0.8	7
83	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
84	Gender-related effects on urine l-cystine metastability. Amino Acids, 2014, 46, 415-427.	1.2	6
85	Systemic Redox Biomarkers in Neurodegenerative Diseases. Current Drug Metabolism, 2015, 16, 46-70.	0.7	6
86	Changes in Total Homocysteine and Glutathione Levels After Laparoscopic Sleeve Gastrectomy in Children with Metabolic-Associated Fatty Liver Disease. Obesity Surgery, 2021, , 1.	1.1	6
87	Pitfalls in the quantitative imaging of glutathione in living cells. Nature Communications, 2018, 9, 1588.	5.8	5
88	Benefits and Toxicity of Disulfiram in Preclinical Models of Nephropathic Cystinosis. Cells, 2021, 10, 3294.	1.8	5
89	Transcriptional and Posttranscriptional Regulation of the CTNS Gene. Pediatric Research, 2011, 70, 130-135.	1.1	4
90	Purification and characterization of a novel alpha-class glutathione transferase from human liver. International Journal of Biochemistry and Cell Biology, 1998, 30, 1235-1243.	1.2	2

Anna Pastore

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91	Renal hemodynamic effect of tacrolimus in renal transplanted children. Pediatric Nephrology, 2001, 16, 773-776.	0.9	2
92	Studying nonobstructive azoospermia in cystinosis: histologicÂexamination of testes andÂepididymis and sperm analysis inÂa Ctnsâ''/â^' mouse model. Fertility and Sterility, 2012, 98, 162-165.	0.5	2
93	Pediatric reference intervals for muscle coenzyme Q10. Biomarkers, 2012, 17, 764-766.	0.9	2
94	Drastic Reduction of Piperacillin-Tazobactam Concentrations in an in-vitro Model of Continuous Venovenous Hemofiltration: Proposal of An Innovative Modality of Administration to Maintain them at Constant Concentration. Cardiovascular and Hematological Agents in Medicinal Chemistry, 2014, 11, 187-193.	0.4	1
95	Cerebrospinal Fluid Levels of AFP and hCG: Validation of the Analytical Method and Application in the Diagnosis of Central Nervous System Germ Cell Tumors. Diagnostics, 2021, 11, 1980.	1.3	1
96	Clutathione Status in MMACHC Patients. Free Radical Biology and Medicine, 2012, 53, S69-S70.	1.3	0
97	Response to Zhang et al Genetics in Medicine, 2020, 22, 662-662.	1.1	0