

Sonia Fargue

List of Publications by Year in descending order

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Version: 2024-02-01

38
papers

1,689
citations

331259

21
h-index

377514

34
g-index

41
all docs

41
docs citations

41
times ranked

1308
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary hyperoxaluria Type 1: indications for screening and guidance for diagnosis and treatment. <i>Nephrology Dialysis Transplantation</i> , 2012, 27, 1729-1736.	0.4	266
2	Primary hyperoxaluria type 1: still challenging!. <i>Pediatric Nephrology</i> , 2006, 21, 1075-1081.	0.9	135
3	Disease recurrence in paediatric renal transplantation. <i>Pediatric Nephrology</i> , 2009, 24, 2097-2108.	0.9	135
4	Genotype-phenotype correlation in primary hyperoxaluria type 1: the p.Gly170Arg AGXT mutation is associated with a better outcome. <i>Kidney International</i> , 2010, 77, 443-449.	2.6	117
5	Multiple mechanisms of action of pyridoxine in primary hyperoxaluria type 1. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2013, 1832, 1776-1783.	1.8	85
6	Screening for NPHS2 Mutations May Help Predict FSGS Recurrence after Transplantation. <i>Journal of the American Society of Nephrology: JASN</i> , 2011, 22, 579-585.	3.0	82
7	Primary Hyperoxaluria. <i>International Journal of Nephrology</i> , 2011, 2011, 1-11.	0.7	76
8	Four of the Most Common Mutations in Primary Hyperoxaluria Type 1 Unmask the Cryptic Mitochondrial Targeting Sequence of Alanine:glyoxylate Aminotransferase Encoded by the Polymorphic Minor Allele. <i>Journal of Biological Chemistry</i> , 2013, 288, 2475-2484.	1.6	76
9	Effect of conservative treatment on the renal outcome of children with primary hyperoxaluria type 1. <i>Kidney International</i> , 2009, 76, 767-773.	2.6	57
10	Pharmacologic rescue of an enzyme-trafficking defect in primary hyperoxaluria 1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 14406-14411.	3.3	56
11	Survey of First-Year Medical Students to Assess Their Knowledge and Attitudes Toward Organ Transplantation and Donation. <i>Transplantation Proceedings</i> , 2009, 41, 634-638.	0.3	52
12	End Points for Clinical Trials in Primary Hyperoxaluria. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2020, 15, 1056-1065.	2.2	51
13	Pyridoxamine and pyridoxal are more effective than pyridoxine in rescuing folding-defective variants of human alanine:glyoxylate aminotransferase causing primary hyperoxaluria type I. <i>Human Molecular Genetics</i> , 2015, 24, 5500-5511.	1.4	50
14	Primary hyperoxaluria type 1: strategy for organ transplantation. <i>Current Opinion in Organ Transplantation</i> , 2010, 15, 590-593.	0.8	47
15	Hydroxyproline Metabolism and Oxalate Synthesis in Primary Hyperoxaluria. <i>Journal of the American Society of Nephrology: JASN</i> , 2018, 29, 1615-1623.	3.0	44
16	Contribution of Dietary Oxalate and Oxalate Precursors to Urinary Oxalate Excretion. <i>Nutrients</i> , 2021, 13, 62.	1.7	39
17	Tacrolimus nephrotoxicity: beware of the association of diarrhea, drug interaction and pharmacogenetics. <i>Pediatric Nephrology</i> , 2010, 25, 965-969.	0.9	37
18	Metabolism of 13C5-hydroxyproline in mouse models of Primary Hyperoxaluria and its inhibition by RNAi therapeutics targeting liver glyoxylate oxidase and hydroxyproline dehydrogenase. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2016, 1862, 233-239.	1.8	33

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19	Reduction in urinary oxalate excretion in mouse models of Primary Hyperoxaluria by RNA interference inhibition of liver lactate dehydrogenase activity. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2019, 1865, 2203-2209.	1.8	33
20	Bone metabolism in oxalosis: a single-center study using new imaging techniques and biomarkers. <i>Pediatric Nephrology</i> , 2010, 25, 1081-1089.	0.9	31
21	The N-terminal extension is essential for the formation of the active dimeric structure of liver peroxisomal alanine:glyoxylate aminotransferase. <i>International Journal of Biochemistry and Cell Biology</i> , 2012, 44, 536-546.	1.2	29
22	High throughput cell-based assay for identification of glycolate oxidase inhibitors as a potential treatment for Primary Hyperoxaluria Type 1. <i>Scientific Reports</i> , 2016, 6, 34060.	1.6	20
23	Development of a Phenotypic High-Content Assay to Identify Pharmacoperone Drugs for the Treatment of Primary Hyperoxaluria Type 1 by High-Throughput Screening. <i>Assay and Drug Development Technologies</i> , 2015, 13, 16-24.	0.6	19
24	Effects of alanine:glyoxylate aminotransferase variants and pyridoxine sensitivity on oxalate metabolism in a cell-based cytotoxicity assay. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2016, 1862, 1055-1062.	1.8	19
25	The effects of the inactivation of Hydroxyproline dehydrogenase on urinary oxalate and glycolate excretion in mouse models of primary hyperoxaluria. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020, 1866, 165633.	1.8	13
26	Review: New treatment strategies for proliferative lupus nephritis: keep children in mind!. <i>Lupus</i> , 2007, 16, 684-691.	0.8	12
27	Future treatments for hyperoxaluria. <i>Current Opinion in Urology</i> , 2020, 30, 171-176.	0.9	12
28	Primary hyperoxaluria type 1: pathophysiology and genetics. <i>CKJ: Clinical Kidney Journal</i> , 2022, 15, i4-i8.	1.4	12
29	Hyperuricemia after liver transplantation in children. <i>Pediatric Transplantation</i> , 2008, 12, 847-853.	0.5	10
30	A mutation creating an out-of-frame alternative translation initiation site in the <i>GRHPR</i> 5'UTR causing primary hyperoxaluria type 1. <i>Clinical Genetics</i> , 2015, 88, 494-498.	1.0	8
31	Factors influencing clinical outcome in patients with primary hyperoxaluria type 1. <i>Kidney International</i> , 2014, 86, 1074-1076.	2.6	7
32	Effect of alanine supplementation on oxalate synthesis. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2021, 1867, 165981.	1.8	6
33	Generation of a GLO-2 deficient mouse reveals its effects on liver carbonyl and glutathione levels. <i>Biochemistry and Biophysics Reports</i> , 2021, 28, 101138.	0.7	3
34	Primary Hyperoxaluria. , 2009, , 1069-1079.		1
35	Metabolism of Glycolate to Oxalate in Kidney Proximal Tubule Cells. <i>FASEB Journal</i> , 2019, 33, 863.7.	0.2	1
36	MP34-04 INHIBITION OF GLYCOLATE OXIDASE REDUCES URINARY OXALATE EXCRETION IN A MOUSE MODEL OF PRIMARY HYPEROXALURIA TYPE 1. <i>Journal of Urology</i> , 2015, 193, .	0.2	0

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37	MP67-02 THE EFFECT OF ALANINE ON CELL VIABILITY AND AGT ACTIVITY IN TRANSFORMED CHINESE HAMSTER OVARY CELLS. <i>Journal of Urology</i> , 2016, 195, .	0.2	0
38	MP67-03 MITOCHONDRIAL IMPLICATIONS OF GLYCOLATE METABOLISM IN PRIMARY HYPEROXALURIA. <i>Journal of Urology</i> , 2016, 195, .	0.2	0