

Marie-Louise SyrÅ©n

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

618
citations

759055

12
h-index

839398

18
g-index

19
all docs

19
docs citations

19
times ranked

756
citing authors

#	ARTICLE	IF	CITATIONS
1	Activation/deactivation of renal Na ⁺ ,K ⁺ ATPase: a final common pathway for regulation of natriuresis. <i>FASEB Journal</i> , 1994, 8, 436-439.	0.2	234
2	Patients With Biallelic Mutations in the Chloride Channel Gene CLCNKB: Long-Term Management and Outcome. <i>American Journal of Kidney Diseases</i> , 2007, 49, 91-98.	2.1	59
3	Urinary exosomes in the diagnosis of Gitelman and Bartter syndromes. <i>Nephrology Dialysis Transplantation</i> , 2015, 30, 621-630.	0.4	51
4	Identification of fifteen novel mutations in the SLC12A3 gene encoding the Na-Cl Co-transporter in Italian patients with Gitelman syndrome. <i>Human Mutation</i> , 2002, 20, 78-78.	1.1	49
5	ABCB4 mutations in adult patients with cholestatic liver disease: impact and phenotypic expression. <i>Journal of Gastroenterology</i> , 2016, 51, 271-280.	2.3	45
6	Bioactive Compounds in Edible Oils and Their Role in Oxidative Stress and Inflammation. <i>Frontiers in Physiology</i> , 2021, 12, 659551.	1.3	37
7	Early appearance of hypokalemia in Gitelman syndrome. <i>Pediatric Nephrology</i> , 2010, 25, 2179-2182.	0.9	34
8	Renal phosphate handling in Gitelman syndrome—the results of a case–control study. <i>Pediatric Nephrology</i> , 2013, 28, 65-70.	0.9	22
9	Cardiac arrhythmias due to severe hypokalemia in a patient with classic Bartter disease. <i>Pediatric Nephrology</i> , 2004, 19, 1413-1415.	0.9	18
10	Simultaneous Mutations in the CLCNKB and SLC12A3 Genes in Two Siblings with Phenotypic Heterogeneity in Classic Bartter Syndrome. <i>Pediatric Research</i> , 2005, 58, 1269-1273.	1.1	18
11	The polyunsaturated fatty acid balance in kidney health and disease: A review. <i>Clinical Nutrition</i> , 2018, 37, 1829-1839.	2.3	18
12	N-3 Polyunsaturated Fatty Acids in Menopausal Transition: A Systematic Review of Depressive and Cognitive Disorders with Accompanying Vasomotor Symptoms. <i>International Journal of Molecular Sciences</i> , 2018, 19, 1849.	1.8	12
13	The mutation c.1196_1202dup7bp (p.Ser402X) in the SLC12A3 gene clusters in Italian Gitelman syndrome patients and reflects the presence of a common ancestor. <i>Nephrology Dialysis Transplantation</i> , 2011, 26, 557-561.	0.4	7
14	Whole blood fatty acid profile of young subjects and adherence to the Mediterranean diet: an observational cohort study. <i>Lipids in Health and Disease</i> , 2022, 21, 23.	1.2	4
15	Direct and Indirect Effects of Blood Levels of Omega-3 and Omega-6 Fatty Acids on Reading and Writing (Dis)Abilities. <i>Brain Sciences</i> , 2022, 12, 169.	1.1	3
16	Persistent Abnormalities of Fatty Acids Profile in Children With Idiopathic Nephrotic Syndrome in Stable Remission. <i>Frontiers in Pediatrics</i> , 2020, 8, 633470.	0.9	2
17	Adherence to the Mediterranean Diet Improves Fatty Acids Profile in Pediatric Patients with Idiopathic Nephrotic Syndrome. <i>Nutrients</i> , 2021, 13, 4110.	1.7	2
18	Long-Chain Polyunsaturated Fatty Acids Supplementation and Respiratory Infections. <i>Annals of Nutrition and Metabolism</i> , 2022, 78, 8-15.	1.0	2