Marie-Louise SyrÃ©n

List of Publications by Year
in descending order

18	618	759055 12	839398 18
papers	citations	h-index	g-index

19	19	19	756
all docs	docs citations	times ranked	citing authors

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