Paula SÃ¡nchez-Pintos

List of Publications by Year
in descending order

12 papers	342 citations	1162889 8 h-index	1199470 12 g-index
12 all docs	12 docs citations	12 times ranked	433 citing authors

#	Article	IF	Citations
1	Rapid Molecular Diagnosis of Genetically Inherited Neuromuscular Disorders Using Next-Generation Sequencing Technologies. Journal of Clinical Medicine, 2022, 11, 2750.	1.0	3
2	Transferrin Isoforms, Old but New Biomarkers in Hereditary Fructose Intolerance. Journal of Clinical Medicine, 2021, 10, 2932.	1.0	4
3	Evaluation of Body Composition, Physical Activity, and Food Intake in Patients with Inborn Errors of Intermediary Metabolism. Nutrients, 2021, 13, 2111.	1.7	6
4	Treatment adherence in tyrosinemia type 1 patients. Orphanet Journal of Rare Diseases, 2021, 16, 256.	1.2	12
5	Bone Status in Patients with Phenylketonuria: A Systematic Review. Nutrients, 2020, 12, 2154.	1.7	23
6	Clinical Utility of LCT Genotyping in Children with Suspected Functional Gastrointestinal Disorder. Nutrients, 2020, 12, 3017.	1.7	4
7	Automated therapy preparation of isoleucine formulations using 3D printing for the treatment of MSUD: First single-centre, prospective, crossover study in patients. International Journal of Pharmaceutics, 2019, 567, 118497.	2.6	171
8	Acylcarnitine profile in neonatal hypoxic-ischemic encephalopathy. Medicine (United States), 2019, 98, e15221.	0.4	21
9	Evolution of tyrosinemia type 1 disease in patients treated with nitisinone in Spain. Medicine (United) Tj ETQq1 1 0.784314 rgBT /Ove	0.4	17
10	Carbohydrate status in patients with phenylketonuria. Orphanet Journal of Rare Diseases, 2018, 13, 103.	1.2	36
11	Similarities between acylcarnitine profiles in large for gestational age newborns and obesity. Scientific Reports, 2017, 7, 16267.	1.6	19
12	Lipid profile status and other related factors in patients with Hyperphenylalaninaemia. Orphanet Journal of Rare Diseases, 2016, 11, 123.	1.2	26