Ãlvaro Hermida-Ameijeiras

List of Publications by Year
in descending order

8 papers	36 citations	1937685 4 h-index	1872680 6 g-index
8 all docs	8 docs citations	8 times ranked	24 citing authors

#	Article	IF	Citations
1	LipoDDx: a mobile application for identification of rare lipodystrophy syndromes. Orphanet Journal of Rare Diseases, 2020, 15, 81.	2.7	7
2	Familial partial lipodystrophy syndromes. Presse Medicale, 2021, 50, 104071.	1.9	6
3	Characterization of a Novel Splicing Variant in Acylglycerol Kinase (AGK) Associated with Fatal Sengers Syndrome. International Journal of Molecular Sciences, 2021, 22, 13484.	4.1	6
4	Variable Expressivity and Allelic Heterogeneity in Type 2 Familial Partial Lipodystrophy: The p.(Thr528Met) LMNA Variant. Journal of Clinical Medicine, 2021, 10, 1497.	2.4	5
5	Celiaâ€™s Encephalopathy (BSCL2-Gene-Related): Current Understanding. Journal of Clinical Medicine, 2021, 10, 1435.	2.4	5
6	Rapid Molecular Diagnosis of Genetically Inherited Neuromuscular Disorders Using Next-Generation Sequencing Technologies. Journal of Clinical Medicine, 2022, 11, 2750.	2.4	3
7	Variable Expressivity in Type 2 Familial Partial Lipodystrophy Related to R482 and N466 Variants in the LMNA Gene. Journal of Clinical Medicine, 2021, 10, 1259.	2.4	2
8	Are Differences in Inflammatory Markers between Patients with and without Hypertension-Mediated Organ Damage Influenced by Circadian Blood Pressure Abnormalities?. Journal of Clinical Medicine, 2022, 11, 1252.	2.4	2