

# Álvvaro Hermida-Ameijeiras

## List of Publications by Year in descending order

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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	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
2	Rapid Molecular Diagnosis of Genetically Inherited Neuromuscular Disorders Using Next-Generation Sequencing Technologies. Journal of Clinical Medicine, 2022, 11, 2750.	2.4	3
3	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
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	Celiacâ€™s Encephalopathy (BSCL2-Gene-Related): Current Understanding. Journal of Clinical Medicine, 2021, 10, 1435.	2.4	5
6	Familial partial lipodystrophy syndromes. Presse Medicale, 2021, 50, 104071.	1.9	6
7	Characterization of a Novel Splicing Variant in Acylglycerol Kinase (ACK) Associated with Fatal Sengers Syndrome. International Journal of Molecular Sciences, 2021, 22, 13484.	4.1	6
8	LipoDDx: a mobile application for identification of rare lipodystrophy syndromes. Orphanet Journal of Rare Diseases, 2020, 15, 81.	2.7	7