

Alejandro GarcÃ-a CastaÃ±o

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7053653/publications.pdf>

Version: 2024-02-01

7
papers

120
citations

1684188

5
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

242
citing authors

#	ARTICLE	IF	CITATIONS
1	Claudin-16 Deficiency Impairs Tight Junction Function in Ameloblasts, Leading to Abnormal Enamel Formation. <i>Journal of Bone and Mineral Research</i> , 2016, 31, 498-513.	2.8	50
2	Poor phenotype-genotype association in a large series of patients with Type III Bartter syndrome. <i>PLoS ONE</i> , 2017, 12, e0173581.	2.5	27
3	Genetics of Type III Bartter Syndrome in Spain, Proposed Diagnostic Algorithm. <i>PLoS ONE</i> , 2013, 8, e74673.	2.5	16
4	Novel variant in the <i>CNNM2</i> gene associated with dominant hypomagnesemia. <i>PLoS ONE</i> , 2020, 15, e0239965.	2.5	10
5	Forty-One Individuals With Mutations in the <i>AVP-NP2</i> Gene Associated With Familial Neurohypophyseal Diabetes Insipidus. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 1112-1118.	3.6	9
6	Novel compound heterozygous mutations of <i>CLDN16</i> in a patient with familial hypomagnesemia with hypercalciuria and nephrocalcinosis. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1475.	1.2	7
7	Response to Letter to the Editor: “Forty-One Individuals with Mutations in the <i>AVP-NP2</i> Gene Associated with Familial Neurohypophyseal Diabetes Insipidus. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e2687-e2688.	3.6	1