Alejandro GarcÃ-a CastaÃ±o

List of Publications by Year
in descending order

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