Gabriella de Medeiros Abreu

List of Publications by Year
in descending order

13	153	1163117 8	1199594 12
papers	citations	h-index	g-index

13	13	13	312
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of Variants Responsible for Monogenic Forms of Diabetes in Brazil. Frontiers in Endocrinology, 2022, 13, 827325.	3.5	2
2	A Rare Potential Pathogenic Variant in the BDNF Gene is Found in a Brazilian Patient with Severe Childhood-Onset Obesity. Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy, 2021, Volume 14, 11-22.	2.4	7
3	Identification of the First PAX4-MODY Family Reported in Brazil . Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy, 2020, Volume 13, 2623-2631.	2.4	10
4	Identification of a Rare and Potential Pathogenic MC4R Variant in a Brazilian Patient With Adulthood-Onset Severe Obesity. Frontiers in Genetics, 2020, 11, 608840.	2.3	2
5	Identification of a novel large deletion and other copy number variations in the CFTR gene in patients with Cystic Fibrosis from a multiethnic population. Molecular Genetics & Genomic Medicine, 2019, 7, e00645.	1.2	8
6	The association of the fat mass and obesity-associated gene (FTO) rs9939609 polymorphism and the severe obesity in a Brazilian population . Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy, 2019, Volume 12, 667-684.	2.4	26
7	Identification of the MC4R start lost mutation in a morbidly obese Brazilian patient . Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy, 2019, Volume 12, 257-266.	2.4	11
8	The first case of NEUROD1â€MODY reported in Latin America. Molecular Genetics & Genomic Medicine, 2019, 7, e989.	1.2	11
9	MODY probability calculator for GCK and HNF1A screening in a multiethnic background population. Archives of Endocrinology and Metabolism, 2019, 64, 17-23.	0.6	7
10	Clinical profiles associated with LRRK2 and GBA mutations in Brazilians with Parkinson's disease. Journal of the Neurological Sciences, 2017, 381, 160-164.	0.6	27
11	Autosomal dominant Parkinsonâ€™s disease: Incidence of mutations in LRRK2, SNCA, VPS35 and GBA genes in Brazil. Neuroscience Letters, 2016, 635, 67-70.	2.1	13
12	Association of LRRK2 and GBA mutations in a Brazilian family with Parkinson's disease. Parkinsonism and Related Disorders, 2015, 21, 825-826.	2.2	11
13	Parkinson disease: Î±-synuclein mutational screening and new clinical insight into the p.E46K mutation. Parkinsonism and Related Disorders, 2015, 21, 586-589.	2.2	18