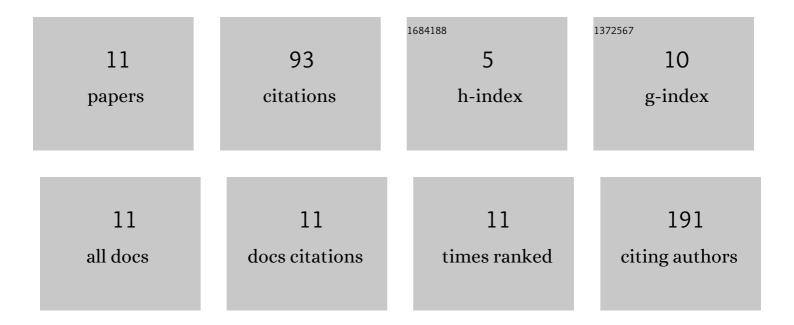
Luciana Cardoso Bonadia

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6387395/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	CFTR genotype and clinical outcomes of adult patients carried as cystic fibrosis disease. Gene, 2014, 540, 183-190.	2.2	37
2	Brain Structural Signature of <scp><i>RFC1</i></scp> â€Related Disorder. Movement Disorders, 2021, 36, 2634-2641.	3.9	19
3	Intermediate-length CAG repeat in ATXN2 is associated with increased risk for amyotrophic lateral sclerosis in Brazilian patients. Neurobiology of Aging, 2018, 69, 292.e15-292.e18.	3.1	10
4	<p>Clinical and Molecular Investigation of Familial Multiple Lipomatosis: Variants in the HMGA2 Gene</p> . Clinical, Cosmetic and Investigational Dermatology, 2020, Volume 13, 1-10.	1.8	6
5	Diagnostic Yield of Whole Exome Sequencing for Adults with Ataxia: a Brazilian Perspective. Cerebellum, 2022, 21, 49-54.	2.5	6
6	Association Between C1236T (rs1128503) Variant in ABCB1 Gene and Breast Cancer Recurrence. Clinical Cancer Drugs, 2018, 5, 60-64.	0.3	4
7	Burkholderia cepacia complex in cystic fibrosis in a Brazilian reference center. Medical Microbiology and Immunology, 2017, 206, 447-461.	4.8	3
8	Frequency and Genetic Profile of Compound Heterozygous Friedreich's Ataxia Patients—the Brazilian Experience. Cerebellum, 2019, 18, 1143-1146.	2.5	2
9	CAG repeatsâ€^≥â€^34 in Ataxin-1 gene are associated with amyotrophic lateral sclerosis in a Brazilian cohort. Journal of the Neurological Sciences, 2020, 414, 116842.	0.6	2
10	Slowly progressive behavioral frontotemporal dementia syndrome in a family coâ€segregating the C9orf 72 expansion and a Synaptophysin mutation. Alzheimer's and Dementia, 2021, , .	0.8	2
11	DRPLA: An unusual disease or an underestimated cause of ataxia in Brazil?. Parkinsonism and Related Disorders, 2021, 92, 67-71.	2.2	2