Jorge Javier Cebolla

List of Publications by Year
in descending order

16 papers	163 citations	1163117 8 h-index	1125743 13 g-index
25 all docs	25 docs citations	25 times ranked	261 citing authors

#	Article	IF	Citations
1	Identification of risk features for complication in Gaucher's disease patients: a machine learning analysis of the Spanish registry of Gaucher disease. Orphanet Journal of Rare Diseases, 2020, 15, 256.	2.7	15
2	Serum protein profile analysis in lysosomal storage disorders patients. Clinica Chimica Acta, 2020, 510, 430-436.	1.1	3
3	LC-MS/MS analysis of plasma glucosylsphingosine as a biomarker for diagnosis and follow-up monitoring in Gaucher disease in the Spanish population. Clinical Chemistry and Laboratory Medicine, 2020, 58, 798-809.	2.3	12
4	New variants in Spanish Niemann–Pick type c disease patients. Molecular Biology Reports, 2020, 47, 2085-2095.	2.3	4
5	Mutations identified in a cohort of Mexican patients with lysosomal acid lipase deficiency. Annals of Hepatology, 2019, 18, 646-650.	1.5	3
6	Evaluation of two approaches to lysosomal acid lipase deficiency patient identification: An observational retrospective study. Atherosclerosis, 2019, 285, 49-54.	0.8	3
7	Muscle-tendon weakness contributes to chronic fatigue syndrome in Gaucher's disease. Journal of Orthopaedic Surgery and Research, 2019, 14, 383.	2.3	2
8	Web-Based Bioinformatics Predictors: Recommendations to Assess Lysosomal Cholesterol Trafficking Diseases-Related Genes. Methods of Information in Medicine, 2019, 58, 050-059.	1.2	1
9	The erythrocyte osmotic resistance test as screening tool for cholesterol-related lysosomal storage diseases. Clinica Chimica Acta, 2018, 480, 161-165.	1.1	7
10	Biomarker combination is necessary for the assessment of Gaucher disease?. Annals of Translational Medicine, 2018, 6, S81-S81.	1.7	14
11	Assessment of plasma chitotriosidase activity, CCL18/PARC concentration and NP-C suspicion index in the diagnosis of Niemann-Pick disease type C: a prospective observational study. Journal of Translational Medicine, 2017, 15, 43.	4.4	19
12	Update on lysosomal acid lipase deficiency: Diagnosis, treatment and patient management. Medicina Clínica (English Edition), 2017, 148, 429.e1-429.e10.	0.2	5
13	Actualización en deficiencia de lipasa ácida lisosomal: diagnóstico, tratamiento y seguimiento de los pacientes. Medicina Clínica, 2017, 148, 429.e1-429.e10.	0.6	18
14	Inhibition of Intermediate-Conductance Calcium-Activated K Channel (KCa3.1) and Fibroblast Mitogenesis by α-Linolenic Acid and Alterations of Channel Expression in the Lysosomal Storage Disorders, Fabry Disease, and Niemann Pick C. Frontiers in Physiology, 2017, 8, 39.	2.8	11
15	Increased glycolipid storage produced by the inheritance of a complex intronic haplotype in the α-galactosidase A (GLA) gene. BMC Genetics, 2015, 16, 109.	2.7	17
16	The Influence of Genetic Variability and Proinflammatory Status on the Development of Bone Disease in Patients with Gaucher Disease. PLoS ONE, 2015, 10, e0126153.	2.5	22