

# Cristóbal Colón

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7042512/publications.pdf>

Version: 2024-02-01

10  
papers

187  
citations

1478458

6  
h-index

1474186

9  
g-index

10  
all docs

10  
docs citations

10  
times ranked

348  
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluation and long-term follow-up of infants with inborn errors of metabolism identified in an expanded screening programme. <i>Molecular Genetics and Metabolism</i> , 2011, 104, 470-475.	1.1	77
2	Newborn screening for Fabry disease in the north-west of Spain. <i>European Journal of Pediatrics</i> , 2017, 176, 1075-1081.	2.7	40
3	Reference values of amino acids, acylcarnitines and succinylacetone by tandem mass spectrometry for use in newborn screening in southwest Colombia. , 2017, v48, 113-119.		13
4	A selective screening program for the early detection of mucopolysaccharidosis. <i>Medicine (United States)</i> , 2010, 89, 1075-1081.	1.0	12
5	Proteomic Analysis in Morquio A Cells Treated with Immobilized Enzymatic Replacement Therapy on Nanostructured Lipid Systems. <i>International Journal of Molecular Sciences</i> , 2019, 20, 4610.	4.1	12
6	Characterization of New Proteomic Biomarker Candidates in Mucopolysaccharidosis Type IVA. <i>International Journal of Molecular Sciences</i> , 2021, 22, 226.	4.1	11
7	Plasma Proteomic Analysis in Morquio A Disease. <i>International Journal of Molecular Sciences</i> , 2021, 22, 6165.	4.1	8
8	Congenital Hypothyroidism with Neurological and Respiratory Alterations: A Case Detected Using a Variable Diagnostic Threshold for Tsh. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2011, 3, 208-211.	0.9	7
9	Enzyme-Loaded Gel Core Nanostructured Lipid Carriers to Improve Treatment of Lysosomal Storage Diseases: Formulation and In Vitro Cellular Studies of Elosulfase Alfa-Loaded Systems. <i>Pharmaceutics</i> , 2019, 11, 522.	4.5	5
10	Family study of a novel mutation of mucopolysaccharidosis type VI with a severe phenotype and good response to enzymatic replacement therapy. <i>Medicine (United States)</i> , 2018, 97, e12872.	1.0	2