JoÃ£o Lavinha

List of Publications by Year
in descending order

13	368	1163117 8	1199594 12
papers	citations	h-index	g-index

13	13	13	574
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Zinc Deficiency Interacts with Intestinal/Urogenital Parasites in the Pathway to Anemia in Preschool Children, Bengoâ€“Angola. Nutrients, 2022, 14, 1392.	4.1	3
2	Effectiveness of Nutrition and WASH/malaria educational community-based interventions in reducing anemia in children from Angola. Scientific Reports, 2021, 11, 5603.	3.3	3
3	Iron deficiency anaemia among 6-to-36-month children from northern Angola. BMC Pediatrics, 2020, 20, 298.	1.7	14
4	Efficacy of Nutrition and WASH/Malaria Educational Community-Based Interventions in Reducing Anemia in Preschool Children from Bengo, Angola: Study Protocol of a Randomized Controlled Trial. International Journal of Environmental Research and Public Health, 2019, 16, 466.	2.6	7
5	Early modification of sickle cell disease clinical course by UDP-glucuronosyltransferase 1A1 gene promoter polymorphism. Journal of Human Genetics, 2008, 53, 524-528.	2.3	12
6	Comment on â€˜Nonsense-mediated mRNA decay modulates clinical outcome of genetic diseaseâ€™. European Journal of Human Genetics, 2007, 15, 533-534.	2.8	4
7	Nonsense Mutations in Close Proximity to the Initiation Codon Fail to Trigger Full Nonsense-mediated mRNA Decay. Journal of Biological Chemistry, 2004, 279, 32170-32180.	3.4	116
8	Cystic Fibrosis F508del Patients Have Apically Localized CFTR in a Reduced Number of Airway Cells. Laboratory Investigation, 2000, 80, 857-868.	3.7	93
9	Cystic fibrosis patients with the 3272-26A?G mutation have mild disease, leaky alternative mRNA splicing, and CFTR protein at the cell membrane. Human Mutation, 1999, 14, 133-144.	2.5	59
10	Missense mutation R1066C in the second transmembrane domain of CFTR causes a severe cystic fibrosis phenotype: Study of 19 heterozygous and 2 homozygous patients. Human Mutation, 1997, 10, 387-392.	2.5	12
11	Complex cystic fibrosis allele R334W-R1158X results in reduced levels of correctly processed mRNA in a pancreatic sufficient patient. , 1996, 8, 134-139.		25
12	Characterization of a splicing mutation in the factor VIII gene at the RNA level. Human Genetics, 1995, 95, 109-11.	3.8	9
13	Single-strand conformation polymorphism (SSCP) analysis of the molecular pathology of hemophilia B. Human Mutation, 1993, 2, 355-361.	2.5	11