Juan J TellerÃ-a

List of Publications by Year
in descending order

28	580	567281 15	610901 24
papers	citations	h-index	g-index

32	32	32	859
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	ALOX5 promoter genotype and response to montelukast in moderate persistent asthma. Respiratory Medicine, 2008, 102, 857-861.	2.9	79
2	Spectrum of Mutations in the CFTR Gene in Cystic Fibrosis Patients of Spanish Ancestry. Annals of Human Genetics, 2007, 71, 194-201.	0.8	53
3	Serum levels of CD14 in neonatal sepsis by Gramâ€positive and Gramâ€negative bacteria. Acta Paediatrica, International Journal of Paediatrics, 1996, 85, 728-732.	1.5	49
4	Cytokine gene polymorphisms in retinal detachment patients with and without proliferative vitreoretinopathy: a preliminary study. Acta Ophthalmologica, 2006, 84, 309-313.	0.3	37
5	TNF Î± and LT Î± gene polymorphisms as additional markers of celiac disease susceptibility in a DQ2-positive population. Immunogenetics, 2002, 54, 551-555.	2.4	32
6	Nontraumatic Acute Aortic Emergencies: Part 1, Acute Aortic Syndrome. American Journal of Roentgenology, 2014, 202, 656-665.	2.2	29
7	Antineutrophil cytoplasmic antibodies (ANCA) in idiopathic pulmonary hemosiderosis. Pediatric Allergy and Immunology, 1994, 5, 235-239.	2.6	27
8	PRPH2 (Peripherin/RDS) Mutations Associated with Different Macular Dystrophies in a Spanish Population: A New Mutation. European Journal of Ophthalmology, 2010, 20, 724-732.	1.3	24
9	Tachyphylaxis to Î²2-agonists in Spanish asthmatic patients could be modulated by Î²2-adrenoceptor gene polymorphisms. Respiratory Medicine, 2006, 100, 1072-1078.	2.9	23
10	A nonsense mutation (R1158X) and a splicing mutation (3849 + 4A â†' G) in exon 19 of the cystic fibrosis transmembrane conductance regulator gene. Genomics, 1992, 12, 417-418.	2.9	22
11	TLR2â€"TLR4/CD14 polymorphisms and predisposition to severe invasive infections by Neisseria meningitidis and Streptococcus pneumoniae. Medicina Intensiva, 2014, 38, 356-362.	0.7	21
12	Pathophysiology of Age-Related Macular Degeneration: Implications for Treatment. Ophthalmic Research, 2022, 65, 615-636.	1.9	19
13	CALCA and TRPV1 genes polymorphisms are related to a good outcome in female chronic migraine patients treated with OnabotulinumtoxinA. Journal of Headache and Pain, 2019, 20, 39.	6.0	18
14	Antisilicone Antibodies in Patients with Silicone Implants for Retinal Detachment Surgery. Ophthalmic Research, 2001, 33, 87-90.	1.9	16
15	PRPH2-Related Retinal Diseases: Broadening the Clinical Spectrum and Describing a New Mutation. Genes, 2020, 11, 773.	2.4	16
16	Genotype scores in energy and iron-metabolising genes are higher in elite endurance athletes than in nonathlete controls. Applied Physiology, Nutrition and Metabolism, 2020, 45, 1225-1231.	1.9	16
17	Toll-Like Receptor 2 R753Q Polymorphisms Are Associated With an Increased Risk of Infective Endocarditis. Revista Espanola De Cardiologia (English Ed), 2011, 64, 1056-1059.	0.6	13
18	Polymorphisms in Receptors Involved in Opsonic and Nonopsonic Phagocytosis, and Correlation with Risk of Infection in Oncohematology Patients. Infection and Immunity, 2018, 86, .	2.2	13

#	Article	IF	Citations
19	Lack of association between atopy and RsaI polymorphism within intron 2 of the Pc?RI-? gene in a Spanish population sample. Allergy: European Journal of Allergy and Clinical Immunology, 1998, 53, 1083-1086.	5.7	10
20	Liver-Metabolizing Genes and Their Relationship to the Performance of Elite Spanish Male Endurance Athletes; a Prospective Transversal Study. Sports Medicine - Open, 2019, 5, 50.	3.1	10
21	Genetic Profile in Genes Associated with Cardiorespiratory Fitness in Elite Spanish Male Endurance Athletes. Genes, 2021, 12, 1230.	2.4	9
22	Alpha-1 Antichymotrypsin Levels Are Actively Increased in Normal Colostrum. Journal of Pediatric Gastroenterology and Nutrition, 1998, 26, 376-379.	1.8	6
23	Nontraumatic Acute Aortic Emergencies: Part 2, Pre- and Postsurgical Complications Related to Aortic Aneurysm in the Emergency Clinical Setting. American Journal of Roentgenology, 2014, 202, 666-674.	2.2	5
24	NeuropatÃa auditiva autosÃ³mica dominante y variante DIAPH3 (c.-173C>T). Acta OtorrinolaringolÃ³gica EspaÃ±ola, 2017, 68, 183-185.	0.4	5
25	The ?F508 mutation and RFLP-linked loci in Spanish cystic fibrosis families. Human Genetics, 1991, 87, 516-7.	3.8	1
26	Novel presenilin 1 mutation (p.Thr-Pro116-117Ser-Thr) in a Spanish family with early-onset Alzheimer's disease. Neurobiology of Aging, 2019, 84, 238.e19-238.e24.	3.1	1
27	Relevancia de las nuevas pruebas genÃ©ticas en el diagnÃ³stico de la talla baja con dismorfias. Medicina ClÃnica, 2016, 147, e67-e68.	0.6	0
28	Frequency of Cystic Fibrosis Mutations and Associated Haplotype Distribution in Slovak CF Patients. Advances in Experimental Medicine and Biology, 1991, 290, 383-385.	1.6	0