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	Pathophysiology of Age-Related Macular Degeneration: Implications for Treatment. Ophthalmic Research, 2022, 65, 615-636.	1.9	19
2	Genetic Profile in Genes Associated with Cardiorespiratory Fitness in Elite Spanish Male Endurance Athletes. Genes, 2021, 12, 1230.	2.4	9
3	PRPH2-Related Retinal Diseases: Broadening the Clinical Spectrum and Describing a New Mutation. Genes, 2020, 11, 773.	2.4	16
4	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
5	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
6	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
7	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
8	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
9	NeuropatÃa auditiva autosÃ³mica dominante y variante DIAPH3 (c.-173C>T). Acta OtorrinolaringolÃ³gica EspaÃ±ola, 2017, 68, 183-185.	0.4	5
10	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
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	Nontraumatic Acute Aortic Emergencies: Part 1, Acute Aortic Syndrome. American Journal of Roentgenology, 2014, 202, 656-665.	2.2	29
13	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
14	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
15	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
16	ALOX5 promoter genotype and response to montelukast in moderate persistent asthma. Respiratory Medicine, 2008, 102, 857-861.	2.9	79
17	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
18	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

#	Article	IF	Citations
19	Cytokine gene polymorphisms in retinal detachment patients with and without proliferative vitreoretinopathy: a preliminary study. Acta Ophthalmologica, 2006, 84, 309-313.	0.3	37
20	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
21	Antisilicone Antibodies in Patients with Silicone Implants for Retinal Detachment Surgery. Ophthalmic Research, 2001, 33, 87-90.	1.9	16
22	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
23	Alpha-1 Antichymotrypsin Levels Are Actively Increased in Normal Colostrum. Journal of Pediatric Gastroenterology and Nutrition, 1998, 26, 376-379.	1.8	6
24	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
25	Antineutrophil cytoplasmic antibodies (ANCA) in idiopathic pulmonary hemosiderosis. Pediatric Allergy and Immunology, 1994, 5, 235-239.	2.6	27
26	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
27	The ?F508 mutation and RFLP-linked loci in Spanish cystic fibrosis families. Human Genetics, 1991, 87, 516-7.	3.8	1
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