

Ricardo M Fujita

List of Publications by Year in descending order

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Version: 2024-02-01

59
papers

2,124
citations

218677

26
h-index

233421

45
g-index

61
all docs

61
docs citations

61
times ranked

3091
citing authors

#	ARTICLE	IF	CITATIONS
1	Mannose-binding lectin and its genetic variants. <i>Genes and Immunity</i> , 2006, 7, 85-94.	4.1	395
2	Genome-wide association study identifies five new susceptibility loci for primary angle closure glaucoma. <i>Nature Genetics</i> , 2016, 48, 556-562.	21.4	147
3	Geographic population structure analysis of worldwide human populations infers their biogeographical origins. <i>Nature Communications</i> , 2014, 5, 3513.	12.8	114
4	Y Chromosome Sequences Reveal a Short Beringian Standstill, Rapid Expansion, and early Population structure of Native American Founders. <i>Current Biology</i> , 2019, 29, 149-157.e3.	3.9	94
5	Spectrum of Mutations in the RPGR Gene That Are Identified in 20% of Families with X-Linked Retinitis Pigmentosa. <i>American Journal of Human Genetics</i> , 1997, 61, 1287-1292.	6.2	93
6	Confirmation of linkage of Friedreich ataxia to chromosome 9 and identification of a new closely linked marker. <i>Genomics</i> , 1989, 4, 110-111.	2.9	83
7	Protein-Truncation Mutations in the RP2 Gene in a North American Cohort of Families with X-Linked Retinitis Pigmentosa. <i>American Journal of Human Genetics</i> , 1999, 64, 897-900.	6.2	78
8	Biochemical Characterization and Subcellular Localization of the Mouse Retinitis Pigmentosa GTPase Regulator (mRprgr). <i>Journal of Biological Chemistry</i> , 1998, 273, 19656-19663.	3.4	73
9	A Paleogenomic Reconstruction of the Deep Population History of the Andes. <i>Cell</i> , 2020, 181, 1131-1145.e21.	28.9	69
10	A Novel Locus (RP24) for X-linked Retinitis Pigmentosa Maps to Xq26-27. <i>American Journal of Human Genetics</i> , 1998, 63, 1439-1447.	6.2	65
11	Peruvian horse sickness virus and Yunnan orbivirus, isolated from vertebrates and mosquitoes in Peru and Australia. <i>Virology</i> , 2009, 394, 298-310.	2.4	65
12	Analysis of the RPGR Gene in 11 Pedigrees with the Retinitis Pigmentosa Type 3 Genotype: Paucity of Mutations in the Coding Region but Splice Defects in Two Families. <i>American Journal of Human Genetics</i> , 1997, 61, 571-580.	6.2	61
13	Additional polymorphisms at marker loci D9S5 and D9S15 generate extended haplotypes in linkage disequilibrium with Friedreich ataxia.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1990, 87, 1796-1800.	7.1	59
14	The Current Genomic Landscape of Western South America: Andes, Amazonia, and Pacific Coast. <i>Molecular Biology and Evolution</i> , 2019, 36, 2698-2713.	8.9	59
15	Tracing the genomic ancestry of Peruvians reveals a major legacy of pre-Columbian ancestors. <i>Journal of Human Genetics</i> , 2013, 58, 627-634.	2.3	58
16	Ethics of DNA research on human remains: five globally applicable guidelines. <i>Nature</i> , 2021, 599, 41-46.	27.8	49
17	The Genetic History of Indigenous Populations of the Peruvian and Bolivian Altiplano: The Legacy of the Uros. <i>PLoS ONE</i> , 2013, 8, e73006.	2.5	41
18	Diversity in immunogenomics: the value and the challenge. <i>Nature Methods</i> , 2021, 18, 588-591.	19.0	40

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19	Variation in optineurin (OPTN) allele frequencies between and within populations. <i>Molecular Vision</i> , 2007, 13, 151-63.	1.1	40
20	A new subhaplogroup of native American Y-Chromosomes from the Andes. <i>American Journal of Physical Anthropology</i> , 2011, 146, 553-559.	2.1	38
21	Enclaves of genetic diversity resisted Inca impacts on population history. <i>Scientific Reports</i> , 2017, 7, 17411.	3.3	32
22	The gene encoding the human spasmodic protein (SML1hSP) is in 21q 22.3, physically linked to the homologous breast cancer marker gene BCE1pS2. <i>Genomics</i> , 1992, 13, 1328-1330.	2.9	30
23	Frequencies of Blood Group Systems MNS, Diego, and Duffy and Clinical Phases of Carrion's Disease in Amazonas, Peru. <i>Interdisciplinary Perspectives on Infectious Diseases</i> , 2014, 2014, 1-8.	1.4	30
24	The Genetic History of Peruvian Quechua Mestizas and Chankas: Uniparental DNA Patterns among Autochthonous Amazonian and Andean Populations. <i>Annals of Human Genetics</i> , 2016, 80, 88-101.	0.8	29
25	New native South American Y chromosome lineages. <i>Journal of Human Genetics</i> , 2016, 61, 593-603.	2.3	28
26	Physical mapping of two loci (D9S5 and D9S15) tightly linked to Friedreich ataxia locus (FRDA) and identification of nearby CpG islands by pulse-field gel electrophoresis. <i>Genomics</i> , 1991, 10, 915-920.	2.9	27
27	Population genetics of the jumbo squid <i>Dosidicus gigas</i> (Cephalopoda: Ommastrephidae) in the northern Humboldt Current system based on mitochondrial and microsatellite DNA markers. <i>Fisheries Research</i> , 2016, 175, 1-9.	1.7	20
28	A 530kb YAC contig tightly linked to the Friedreich ataxia locus contains five CpG clusters and a new highly polymorphic microsatellite. <i>Human Genetics</i> , 1992, 89, 531-538.	3.8	19
29	Letter to the Editor: A Carboxy-terminal Truncation of 99 Amino Acids Resulting from a Novel Mutation (Arg555*stop) in the CHM Gene leads to Choroideremia. <i>Experimental Eye Research</i> , 1997, 64, 487-490.	2.6	16
30	Mutational analysis of BRCA1 and BRCA2 genes in Peruvian families with hereditary breast and ovarian cancer. <i>Molecular Genetics & Genomic Medicine</i> , 2017, 5, 481-494.	1.2	14
31	Complement Receptor 1 availability on red blood cell surface modulates <i>Plasmodium vivax</i> invasion of human reticulocytes. <i>Scientific Reports</i> , 2019, 9, 8943.	3.3	14
32	From colorectal cancer pattern to the characterization of individuals at risk: Picture for genetic research in Latin America. <i>International Journal of Cancer</i> , 2019, 145, 318-326.	5.1	14
33	Genetics and genomics in Peru: Clinical and research perspective. <i>Molecular Genetics & Genomic Medicine</i> , 2018, 6, 873-886.	1.2	12
34	Five novel RPGR mutations in families with X-linked retinitis pigmentosa. <i>Human Mutation</i> , 2001, 17, 151-151.	2.5	11
35	Genetic ancestry of families of putative Inka descent. <i>Molecular Genetics and Genomics</i> , 2018, 293, 873-881.	2.1	11
36	Novel and known MYOC exon 3 mutations in an admixed Peruvian primary open-angle glaucoma population. <i>Molecular Vision</i> , 2012, 18, 2067-75.	1.1	11

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37	Recurrent Myocilin Asn480Lys Glaucoma Causative Mutation Arises De Novo in a Family of Andean Descent. <i>Journal of Glaucoma</i> , 2008, 17, 67-72.	1.6	10
38	Patterns of mitochondrial and microsatellite DNA markers describe historical and contemporary dynamics of the Humboldt squid <i>Dosidicus gigas</i> in the Eastern Pacific Ocean. <i>Reviews in Fish Biology and Fisheries</i> , 2020, 30, 519-533.	4.9	10
39	Prenatal diagnosis of Friedreich ataxia. <i>Lancet, The</i> , 1990, 335, 1102.	13.7	9
40	Precision medicine for locally advanced breast cancer: frontiers and challenges in Latin America. <i>Ecancermedicalsecience</i> , 2019, 13, 896.	1.1	8
41	Extreme High Prevalence of a Defective Mannose-Binding Lectin (MBL2) Genotype in Native South American West Andean Populations. <i>PLoS ONE</i> , 2014, 9, e108943.	2.5	6
42	Mapping the Friedreich ataxia locus (FRDA) by linkage disequilibrium analysis with highly polymorphic microsatellites. <i>Biomedicine and Pharmacotherapy</i> , 1994, 48, 219-224.	5.6	5
43	Tracing the genetic history of the Ca^{2+} channel from Ecuador and Peru using uniparental DNA markers. <i>BMC Genomics</i> , 2020, 21, 413.	2.8	5
44	MLPA followed by targeted NGS to detect mutations in the dystrophin gene of Peruvian patients suspected of DMD/DMB. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1759.	1.2	5
45	Clinical expression of X-linked retinitis pigmentosa in a Swedish family with the RP2 genotype. <i>Ophthalmic Genetics</i> , 1998, 19, 187-196.	1.2	4
46	PUM1 and RNase P genes as potential cell-free DNA markers in breast cancer. <i>Journal of Clinical Laboratory Analysis</i> , 2021, 35, e23720.	2.1	4
47	Angiotensin-Converting Enzyme (ACE) genetic variation and longevity in Peruvian older people: a cross-sectional study. <i>Annals of Human Biology</i> , 2020, 47, 309-312.	1.0	3
48	Variabilidad genética de la respuesta inflamatoria. I. Polimorfismo -511 C/T en el gen $IL1\beta$ en diferentes subpoblaciones peruanas. <i>Anales De La Facultad De Medicina</i> , 2012, 73, 221.	0.1	3
49	Mutational profile of KIT and PDGFRA genes in gastrointestinal stromal tumors in Peruvian samples. <i>Revista Espanola De Enfermedades Digestivas</i> , 2015, 107, 72-8.	0.3	3
50	Dinucleotide repeat polymorphism at the DXS977 locus. <i>Human Molecular Genetics</i> , 1994, 3, 1030-1030.	2.9	2
51	Enfermedades Raras o Huérfanas, en Perú más huérfanas que raras. <i>Horizonte Médico</i> , 2018, 18, 4-5.	0.2	2
52	Failure to obtain hybridomas between human macrophages and human tumoral U-937 cells is probably due to parental macrophages. <i>In Vitro Cellular & Developmental Biology</i> , 1993, 29, 362-370.	1.0	1
53	Dinucleotide polymorphism at the DXS1178 locus is tightly linked to PCK1 at Xq13. <i>Human Genetics</i> , 1995, 95, 467-8.	3.8	1
54	Novel mutation in ENG gene causing Hereditary Hemorrhagic Telangiectasia in a Peruvian family. <i>Genetics and Molecular Biology</i> , 2020, 43, e20190126.	1.3	1

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55	[11]Isolation of candidate genes for inherited diseases: Application to X-linked retinal degenerations. Methods in Molecular Genetics, 1996, 8, 207-228.	0.6	0
56	Three novel polymorphic microsatellite markers for the glaucoma locus GLC1B by datamining tetranucleotide repeats on chromosome 2p12-q12. Genetics and Molecular Biology, 2009, 32, 720-722.	1.3	0
57	Population Variation of the Human Genome. , 2021, , 329-350.		0
58	La COVID-19, la ciencia y la pseudociencia, un yin-yang peligroso. Horizonte MÃ©dico, 2021, 21, e1535.	0.2	0
59	Rapid YAC End Sequencing by Alu-Vector PCR and Biotinylated Primers. , 1996, , 227-237.		0