## Antonio Salas

## List of Publications by Year in descending order

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318 18,108 62 119 g-index

337 337 337 337 15880

337 337 15880 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Ancient human genomes suggest three ancestral populations for present-day Europeans. Nature, 2014, 513, 409-413.	27.8	1,179
2	Distinctive Paleo-Indian Migration Routes from Beringia Marked by Two Rare mtDNA Haplogroups. Current Biology, 2009, 19, 1-8.	3.9	738
3	Reconstructing Native American population history. Nature, 2012, 488, 370-374.	27.8	699
4	HaploGrep 2: mitochondrial haplogroup classification in the era of high-throughput sequencing. Nucleic Acids Research, 2016, 44, W58-W63.	14.5	688
5	Correcting for Purifying Selection: An Improved Human Mitochondrial Molecular Clock. American Journal of Human Genetics, 2009, 84, 740-759.	6.2	643
6	A multiplex assay with 52 single nucleotide polymorphisms for human identification. Electrophoresis, 2006, 27, 1713-1724.	2.4	462
7	The Making of the African mtDNA Landscape. American Journal of Human Genetics, 2002, 71, 1082-1111.	6.2	451
8	Updating the East Asian mtDNA phylogeny: a prerequisite for the identification of pathogenic mutations. Human Molecular Genetics, 2006, 15, 2076-2086.	2.9	346
9	Inferring ancestral origin using a single multiplex assay of ancestry-informative marker SNPs. Forensic Science International: Genetics, 2007, 1, 273-280.	3.1	332
10	Genome-wide association study identifies variants in the CFH region associated with host susceptibility to meningococcal disease. Nature Genetics, 2010, 42, 772-776.	21.4	275
11	Diagnostic Test Accuracy of a 2-Transcript Host RNA Signature for Discriminating Bacterial vs Viral Infection in Febrile Children. JAMA - Journal of the American Medical Association, 2016, 316, 835.	7.4	263
12	The Fingerprint of Phantom Mutations in Mitochondrial DNA Data. American Journal of Human Genetics, 2002, 71, 1150-1160.	6.2	249
13	DNA Commission of the International Society for Forensic Genetics: Revised and extended guidelines for mitochondrial DNA typing. Forensic Science International: Genetics, 2014, 13, 134-142.	3.1	243
14	Early human dispersals within the Americas. Science, 2018, 362, .	12.6	230
15	The Phylogeny of the Four Pan-American MtDNA Haplogroups: Implications for Evolutionary and Disease Studies. PLoS ONE, 2008, 3, e1764.	2.5	227
16	Drug Consumption and the Risk of Microscopic Colitis. American Journal of Gastroenterology, 2007, 102, 324-330.	0.4	216
17	A global analysis of Y-chromosomal haplotype diversity for 23 STR loci. Forensic Science International: Genetics, 2014, 12, 12-23.	3.1	214
18	The African Diaspora: Mitochondrial DNA and the Atlantic Slave Trade. American Journal of Human Genetics, 2004, 74, 454-465.	6.2	213

#	Article	IF	CITATIONS
19	Development of a Panel of Genome-Wide Ancestry Informative Markers to Study Admixture Throughout the Americas. PLoS Genetics, 2012, 8, e1002554.	3.5	212
20	A Critical Reassessment of the Role of Mitochondria in Tumorigenesis. PLoS Medicine, 2005, 2, e296.	8.4	188
21	Collagenous and lymphocytic colitis: evaluation of clinical and histological features, response to treatment, and long-term follow-up. American Journal of Gastroenterology, 2003, 98, 340-347.	0.4	174
22	Rapid coastal spread of First Americans: Novel insights from South America's Southern Cone mitochondrial genomes. Genome Research, 2012, 22, 811-820.	5.5	167
23	Incidence of collagenous and lymphocytic colitis: a 5-year population-based study. American Journal of Gastroenterology, 1999, 94, 418-423.	0.4	164
24	Typing of mitochondrial DNA coding region SNPs of forensic and anthropological interest using SNaPshot minisequencing. Forensic Science International, 2004, 140, 251-257.	2.2	161
25	Maternal traces of deep common ancestry and asymmetric gene flow between Pygmy hunter–gatherers and Bantu-speaking farmers. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 1596-1601.	7.1	157
26	The initial peopling of the Americas: A growing number of founding mitochondrial genomes from Beringia. Genome Research, 2010, 20, 1174-1179.	5.5	147
27	Inferring the Demographic History of African Farmers and Pygmy Hunter–Gatherers Using a Multilocus Resequencing Data Set. PLoS Genetics, 2009, 5, e1000448.	3.5	142
28	mtDNA analysis of the Galician population: a genetic edge of European variation. European Journal of Human Genetics, 1998, 6, 365-375.	2.8	141
29	A practical guide to mitochondrial DNA error prevention in clinical, forensic, and population genetics. Biochemical and Biophysical Research Communications, 2005, 335, 891-899.	2.1	138
30	Participation of thromboxane and other eicosanoid synthesis in the course of experimental inflammatory colitis. Gastroenterology, 1990, 98, 269-277.	1.3	137
31	The genetic legacy of western Bantu migrations. Human Genetics, 2005, 117, 366-375.	3.8	131
32	Systematic Evaluation of the Causes of Chronic Watery Diarrhea With Functional Characteristics. American Journal of Gastroenterology, 2007, 102, 2520-2528.	0.4	121
33	Efficacy of anti-TNF therapies in refractory severe microscopic colitis. Journal of Crohn's and Colitis, 2011, 5, 612-618.	1.3	120
34	Genetic association study of exfoliation syndrome identifies a protective rare variant at LOXL1 and five new susceptibility loci. Nature Genetics, 2017, 49, 993-1004.	21.4	114
35	Phylogeographic investigations: The role of trees in forensic genetics. Forensic Science International, 2007, 168, 1-13.	2.2	110
36	Ancestry Analysis in the 11-M Madrid Bomb Attack Investigation. PLoS ONE, 2009, 4, e6583.	2.5	110

3

#	Article	IF	Citations
37	Genome-wide ancestry of 17th-century enslaved Africans from the Caribbean. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 3669-3673.	7.1	110
38	Role of intestinal microflora in chronic inflammation and ulceration of the rat colon Gut, 1994, 35, 1090-1097.	12.1	109
39	Charting the Ancestry of African Americans. American Journal of Human Genetics, 2005, 77, 676-680.	6.2	109
40	Pseudomitochondrial genome haunts disease studies. Journal of Medical Genetics, 2008, 45, 769-772.	3.2	106
41	Spectrum of gluten-sensitive enteropathy in first-degree relatives of patients with coeliac disease: clinical relevance of lymphocytic enteritis. Gut, 2006, 55, 1739-1745.	12.1	104
42	Genetic origin, admixture, and asymmetry in maternal and paternal human lineages in Cuba. BMC Evolutionary Biology, 2008, 8, 213.	3.2	101
43	Resolving relationship tests that show ambiguous STR results using autosomal SNPs as supplementary markers. Forensic Science International: Genetics, 2008, 2, 198-204.	3.1	100
44	Artificial recombination in forensic mtDNA population databases. International Journal of Legal Medicine, 2004, 118, 267-273.	2.2	97
45	SPSmart: adapting population based SNP genotype databases for fast and comprehensive web access. BMC Bioinformatics, 2008, 9, 428.	2.6	95
46	<p>Role of Monocytes/Macrophages in Covid-19 Pathogenesis: Implications for Therapy</p> . Infection and Drug Resistance, 2020, Volume 13, 2485-2493.	2.7	93
47	Evaluating HapMap SNP data transferability in a large-scale genotyping project involving 175 cancer-associated genes. Human Genetics, 2006, 118, 669-679.	3.8	92
48	Mapping genome variation of SARS-CoV-2 worldwide highlights the impact of COVID-19 super-spreaders. Genome Research, 2020, 30, 1434-1448.	5.5	91
49	A reappraisal of complete mtDNA variation in East Asian families with hearing impairment. Human Genetics, 2006, 119, 505-515.	3.8	87
50	Exaggerated status of "novel―and "pathogenic―mtDNA sequence variants due to inadequate database searches. Human Mutation, 2009, 30, 191-196.	2.5	79
51	Coding region mitochondrial DNA SNPs: Targeting East Asian and Native American haplogroups. Forensic Science International: Genetics, 2007, 1, 44-55.	3.1	78
52	Heteroplasmy in mtDNA and the weight of evidence in forensic mtDNA analysis: a case report. International Journal of Legal Medicine, 2001, 114, 186-190.	2.2	75
53	New Population and Phylogenetic Features of the Internal Variation within Mitochondrial DNA Macro-Haplogroup R0. PLoS ONE, 2009, 4, e5112.	2.5	75
54	Low "penetrance―of phylogenetic knowledge in mitochondrial disease studies. Biochemical and Biophysical Research Communications, 2005, 333, 122-130.	2.1	74

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55	Insights into the western Bantu dispersal: mtDNA lineage analysis in Angola. Human Genetics, 2004, 115, 439-47.	3.8	70
56	Dissection of mitochondrial superhaplogroup H using coding region SNPs. Electrophoresis, 2006, 27, 2541-2550.	2.4	70
57	Life-threatening infections in children in Europe (the EUCLIDS Project): a prospective cohort study. The Lancet Child and Adolescent Health, 2018, 2, 404-414.	5.6	69
58	More evidence for non-maternal inheritance of mitochondrial DNA?. Journal of Medical Genetics, 2005, 42, 957-960.	3.2	67
59	Viral Co-Infections in Pediatric Patients Hospitalized with Lower Tract Acute Respiratory Infections. PLoS ONE, 2015, 10, e0136526.	2.5	67
60	Case report: Identification of skeletal remains using short-amplicon marker analysis of severely degraded DNA extracted from a decomposed and charred femur. Forensic Science International: Genetics, 2008, 2, 212-218.	3.1	66
61	Evaluating the Ability of Treeâ€Based Methods and Logistic Regression for the Detection of SNPâ€SNP Interaction. Annals of Human Genetics, 2009, 73, 360-369.	0.8	66
62	Surface hydrophobicity of the rat colonic mucosa is a defensive barrier against macromolecules and toxins. Gut, 2000, 46, 515-521.	12.1	64
63	Problems in FBI mtDNA Database. Science, 2004, 305, 1402b-1404b.	12.6	64
64	Origins and genetic legacies of the Caribbean Taino. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 2341-2346.	7.1	64
65	Sequence variation of a hypervariable short tandem repeat at the D1S1656 locus. International Journal of Legal Medicine, 1998, 111, 244-247.	2.2	63
66	mtDNA Data Mining in GenBank Needs Surveying. American Journal of Human Genetics, 2009, 85, 929-933.	6.2	63
67	The impact of modern migrations on present-day multi-ethnic Argentina as recorded on the mitochondrial DNA genome. BMC Genetics, 2011, 12, 77.	2.7	63
68	Gender bias in the multiethnic genetic composition of central Argentina. Journal of Human Genetics, 2008, 53, 662-674.	2.3	62
69	Mitochondrial Haplogroup U5b3: A Distant Echo of the Epipaleolithic in Italy and the Legacy of the Early Sardinians. American Journal of Human Genetics, 2009, 84, 814-821.	6.2	62
70	What is a â€~novel' mtDNA mutation – and does â€~novelty' really matter?. Journal of Human Genetics 51, 1073-1082.	, 2006, 2.3	61
71	A Bidirectional Corridor in the Sahel-Sudan Belt and the Distinctive Features of the Chad Basin Populations: A History Revealed by the Mitochondrial DNA Genome. Annals of Human Genetics, 2007, 71, 433-452.	0.8	61
72	Current Next Generation Sequencing technology may not meet forensic standards. Forensic Science International: Genetics, 2012, 6, 143-145.	3.1	60

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73	Results of a collaborative study of the EDNAP group regarding mitochondrial DNA heteroplasmy and segregation in hair shafts. Forensic Science International, 2004, 140, 1-11.	2.2	59
74	Predisposing HLA-DQ2 and HLA-DQ8 haplotypes of coeliac disease and associated enteropathy in microscopic colitis. European Journal of Gastroenterology and Hepatology, 2005, 17, 1333-1338.	1.6	59
75	The peopling of South America and the trans-Andean gene flow of the first settlers. Genome Research, 2018, 28, 767-779.	5.5	59
76	Increased Serum Levels of sCD14 and sCD163 Indicate a Preponderant Role for Monocytes in COVID-19 Immunopathology. Frontiers in Immunology, 2020, 11, 560381.	4.8	59
77	ERCC4 Associated with Breast Cancer Risk: A Two-Stage Case-Control Study Using High-throughput Genotyping. Cancer Research, 2006, 66, 9420-9427.	0.9	58
78	Haplogrouping mitochondrial DNA sequences in Legal Medicine/Forensic Genetics. International Journal of Legal Medicine, 2012, 126, 901-916.	2.2	58
79	Ethical-legal problems of DNA databases in criminal investigation. Journal of Medical Ethics, 2000, 26, 266-271.	1.8	57
80	Arrival of Paleo-Indians to the Southern Cone of South America: New Clues from Mitogenomes. PLoS ONE, 2012, 7, e51311.	2.5	57
81	Reconstructing ancient mitochondrial DNA links between Africa and Europe. Genome Research, 2012, 22, 821-826.	5.5	57
82	Cuba: Exploring the History of Admixture and the Genetic Basis of Pigmentation Using Autosomal and Uniparental Markers. PLoS Genetics, 2014, 10, e1004488.	3.5	57
83	Incrimination of anaerobic bacteria in the induction of experimental colitis. American Journal of Physiology - Renal Physiology, 1997, 272, G10-G15.	3.4	56
84	Subepithelial myofibroblasts and tenascin expression in microscopic colitis. Histopathology, 2003, 43, 48-54.	2.9	54
85	Impact of Current Smoking on the Clinical Course of Microscopic Colitis. Inflammatory Bowel Diseases, 2013, 19, 1470-1476.	1.9	54
86	The Paleo-Indian Entry into South America According to Mitogenomes. Molecular Biology and Evolution, 2018, 35, 299-311.	8.9	54
87	Evolution of the incidence of collagenous colitis and lymphocytic colitis in Terrassa, Spain: A population-based study. Inflammatory Bowel Diseases, 2011, 17, 1015-1020.	1.9	53
88	Shipwrecks and founder effects: Divergent demographic histories reflected in Caribbean mtDNA. American Journal of Physical Anthropology, 2005, 128, 855-860.	2.1	52
89	SNPs as Supplements in Simple Kinship Analysis or as Core Markers in Distant Pairwise Relationship Tests: When Do SNPs Add Value or Replace Well-Established and Powerful STR Tests?. Transfusion Medicine and Hemotherapy, 2012, 39, 202-210.	1.6	52
90	Mutation spectra of ABCC8 gene in Spanish patients with hyperinsulinism of infancy (HI). Human Mutation, 2006, 27, 214-214.	2.5	51

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91	Genome-wide Ancestry and Demographic History of African-Descendant Maroon Communities from French Guiana and Suriname. American Journal of Human Genetics, 2017, 101, 725-736.	6.2	50
92	Analysis of the CODIS autosomal STR loci in four main Colombian regions. Forensic Science International, 2003, 137, 67-73.	2.2	49
93	Pharmacogenetics of OATP Transporters Reveals That SLCO1B1 c.388A>G Variant Is Determinant of Increased Atorvastatin Response. International Journal of Molecular Sciences, 2011, 12, 5815-5827.	4.1	49
94	Linking the sub-Saharan and West Eurasian gene pools: maternal and paternal heritage of the Tuareg nomads from the African Sahel. European Journal of Human Genetics, 2010, 18, 915-923.	2.8	47
95	A western route of prehistoric human migration from Africa into the Iberian Peninsula. Proceedings of the Royal Society B: Biological Sciences, 2019, 286, 20182288.	2.6	47
96	Distilling Artificial Recombinants from Large Sets of Complete mtDNA Genomes. PLoS ONE, 2008, 3, e3016.	2.5	46
97	Does Viral Co-Infection Influence the Severity of Acute Respiratory Infection in Children?. PLoS ONE, 2016, 11, e0152481.	2.5	46
98	Results of the 1999–2000 collaborative exercise and proficiency testing program on mitochondrial DNA of the GEP-ISFG: an inter-laboratory study of the observed variability in the heteroplasmy level of hair from the same donor. Forensic Science International, 2002, 125, 1-7.	2.2	45
99	The mtDNA ancestry of admixed Colombian populations. American Journal of Human Biology, 2008, 20, 584-591.	1.6	44
100	Contamination and sample mix-up can best explain some patterns of mtDNA instabilities in buccal cells and oral squamous cell carcinoma. BMC Cancer, 2009, 9, 113.	2.6	44
101	Contamination detection in sequencing studies using the mitochondrial phylogeny. Genome Research, 2021, 31, 309-316.	5.5	44
102	Human genome-wide screen of haplotype-like blocks of reduced diversity. Gene, 2005, 349, 219-225.	2.2	43
103	High penetrance of sequencing errors and interpretative shortcomings in mtDNA sequence analysis of LHON patients. Biochemical and Biophysical Research Communications, 2007, 352, 283-291.	2.1	42
104	Is Mitochondrial DNA Variation Associated with Sporadic Breast Cancer Risk?. Cancer Research, 2008, 68, 623-625.	0.9	42
105	ldentification of novel risk loci and causal insights for sporadic Creutzfeldt-Jakob disease: a genome-wide association study. Lancet Neurology, The, 2020, 19, 840-848.	10.2	42
106	The Genetic Legacy of the Pre-Colonial Period in Contemporary Bolivians. PLoS ONE, 2013, 8, e58980.	2.5	42
107	mtDNA mutations in tumors of the central nervous system reflect the neutral evolution of mtDNA in populations. Oncogene, 2004, 23, 1314-1320.	5.9	41
108	Development and Validation of a New Clinical Scale for Infants with Acute Respiratory Infection: The ReSVinet Scale. PLoS ONE, 2016, 11, e0157665.	2.5	41

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109	Rotavirus and autoimmunity. Journal of Infection, 2020, 81, 183-189.	3.3	41
110	Mitochondrial DNA error prophylaxis: assessing the causes of errors in the GEP'02–03 proficiency testing trial. Forensic Science International, 2005, 148, 191-198.	2.2	40
111	Linguistic and maternal genetic diversity are not correlated in Native Mexicans. Human Genetics, 2009, 126, 521-531.	3.8	40
112	Intestinal Intraepithelial Lymphocyte Cytometric Pattern Is More Accurate than Subepithelial Deposits of Anti-Tissue Transglutaminase IgA for the Diagnosis of Celiac Disease in Lymphocytic Enteritis. PLoS ONE, 2014, 9, e101249.	2.5	40
113	Impact of Rotavirus Vaccination on Childhood Hospitalization for Seizures. Pediatric Infectious Disease Journal, 2015, 34, 769-773.	2.0	40
114	Mapping human dispersals into the Horn of Africa from Arabian Ice Age refugia using mitogenomes. Scientific Reports, 2016, 6, 25472.	3.3	40
115	The 1998–1999 collaborative exercises and proficiency testing program on DNA typing of the Spanish and Portuguese Working Group of the International Society for Forensic Genetics (GEP-ISFG). Forensic Science International, 2000, 114, 21-30.	2.2	39
116	Mitochondrial DNA Haplogroup Background Affects LHON, but Not Suspected LHON, in Chinese Patients. PLoS ONE, 2011, 6, e27750.	2.5	39
117	Micro-geographical differentiation in Northern Iberia revealed by Y-chromosomal DNA analysis. Gene, 2004, 329, 17-25.	2.2	38
118	No evidence of association between common European mitochondrial DNA variants in Alzheimer, Parkinson, and migraine in the Spanish population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 54-65.	1.7	37
119	The 2000–2001 GEP–ISFG Collaborative Exercise on mtDNA: assessing the cause of unsuccessful mtDNA PCR amplification of hair shaft samples. Forensic Science International, 2003, 134, 46-53.	2.2	36
120	Timing and deciphering mitochondrial DNA macro-haplogroup RO variability in Central Europe and Middle East. BMC Evolutionary Biology, 2008, 8, 191.	3.2	36
121	Uniparental Markers of Contemporary Italian Population Reveals Details on Its Pre-Roman Heritage. PLoS ONE, 2012, 7, e50794.	2.5	36
122	The saga of the many studies wrongly associating mitochondrial DNA with breast cancer. BMC Cancer, 2014, 14, 659.	2.6	36
123	Bacteremia in Children Hospitalized with Respiratory Syncytial Virus Infection. PLoS ONE, 2016, 11, e0146599.	2.5	36
124	Impact of mass screening for gluten-sensitive enteropathy in working population. World Journal of Gastroenterology, 2009, 15, 1331.	3.3	35
125	Inferring the Most Likely Geographical Origin of mtDNA Sequence Profiles. Annals of Human Genetics, 2004, 68, 461-471.	0.8	34
126	ENGINES: exploring single nucleotide variation in entire human genomes. BMC Bioinformatics, 2011, 12, 105.	2.6	34

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127	Phylogeography of SARS-CoV-2 pandemic in Spain: a story of multiple introductions, micro-geographic stratification, founder effects, and super-spreaders. Zoological Research, 2020, 41, 605-620.	2.1	34
128	Natural resistance to Meningococcal Disease related to CFH loci: Meta-analysis of genome-wide association studies. Scientific Reports, 2016, 6, 35842.	3.3	33
129	Median network analysis of defectively sequenced entire mitochondrial genomes from early and contemporary disease studies. Journal of Human Genetics, 2009, 54, 174-181.	2.3	32
130	The Etruscan timeline: a recent Anatolian connection. European Journal of Human Genetics, 2009, 17, 693-696.	2.8	32
131	Genomic insights on the ethno-history of the Maya and the  Ladinos' from Guatemala. BMC Genomics, 2015, 16, 131.	2.8	32
132	Rotavirus infection beyond the gut. Infection and Drug Resistance, 2019, Volume 12, 55-64.	2.7	32
133	Investigating the Role of Mitochondrial Haplogroups in Genetic Predisposition to Meningococcal Disease. PLoS ONE, 2009, 4, e8347.	2.5	32
134	The complete mitogenome of a 500-year-old Inca child mummy. Scientific Reports, 2015, 5, 16462.	3.3	31
135	Superspreading in the emergence of COVID-19 variants. Trends in Genetics, 2021, 37, 1069-1080.	6.7	31
136	Insights into Iberian population origins through the construction of highly informative Yâ€chromosome haplotypes using biallelic markers, STRs, and the MSY1 minisatellite. American Journal of Physical Anthropology, 2003, 122, 147-161.	2.1	30
137	Diagnostic value of duodenal antitissue transglutaminase antibodies in glutenâ€sensitive enteropathy. Alimentary Pharmacology and Therapeutics, 2008, 27, 820-829.	3.7	30
138	Mitochondrial DNA as a Risk Factor for False Positives in Case-Control Association Studies. Journal of Genetics and Genomics, 2015, 42, 169-172.	3.9	30
139	Archaeogenomic distinctiveness of the Isthmo-Colombian area. Cell, 2021, 184, 1706-1723.e24.	28.9	30
140	Rapid and enhanced detection of mitochondrial DNA variation using single-strand conformation analysis of superposed restriction enzyme fragments from polymerase chain reaction-amplified products. Electrophoresis, 1997, 18, 52-54.	2.4	29
141	Y chromosome microsatellite genetic variation in two Native American populations from Argentina: Population stratification and mutation data. Forensic Science International: Genetics, 2008, 2, 274-280.	3.1	29
142	Estimating Haplotype Frequency and Coverage of Databases. PLoS ONE, 2008, 3, e3988.	2.5	29
143	A melting pot of multicontinental mtDNA lineages in admixed Venezuelans. American Journal of Physical Anthropology, 2012, 147, 78-87.	2.1	29
144	Evaluating the accuracy of AIM panels at quantifying genome ancestry. BMC Genomics, 2014, 15, 543.	2.8	29

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145	Whole Exome Sequencing reveals new candidate genes in host genomic susceptibility to Respiratory Syncytial Virus Disease. Scientific Reports, 2017, 7, 15888.	3.3	29
146	mtDNA hypervariable region II (HVII) sequences in human evolution studies. European Journal of Human Genetics, 2000, 8, 964-974.	2.8	27
147	SNaPshot Typing of Mitochondrial DNA Coding Region Variants. , 2005, 297, 197-208.		27
148	A qPCR expression assay of IFI44L gene differentiates viral from bacterial infections in febrile children. Scientific Reports, 2019, 9, 11780.	3.3	27
149	Sequence variation of a hypervariable short tandem repeat at the D12S391 locus. Gene, 1996, 182, 151-153.	2.2	26
150	Applications of MALDIâ€TOF MS to largeâ€scale human mtDNA populationâ€based studies. Electrophoresis, 2009, 30, 3665-3673.	2.4	26
151	New Insights into the Lake Chad Basin Population Structure Revealed by High-Throughput Genotyping of Mitochondrial DNA Coding SNPs. PLoS ONE, 2011, 6, e18682.	2.5	26
152	Interdisciplinary approach to the demography of Jamaica. BMC Evolutionary Biology, 2012, 12, 24.	3.2	26
153	Ancestry analysis reveals a predominant Native American component with moderate European admixture in Bolivians. Forensic Science International: Genetics, 2013, 7, 537-542.	3.1	26
154	The Mitochondrial Genome Is a "Genetic Sanctuary―during the Oncogenic Process. PLoS ONE, 2011, 6, e23327.	2.5	26
155	Testing for genetic structure in different urban Argentinian populations. Forensic Science International, 2007, 165, 35-40.	2.2	25
156	â€~Distorted' mitochondrial DNA sequences in schizophrenic patients. European Journal of Human Genetics, 2007, 15, 400-402.	2.8	25
157	D9S1120, a simple STR with a common Native American-specific allele: Forensic optimization, locus characterization and allele frequency studies. Forensic Science International: Genetics, 2008, 3, 7-13.	3.1	25
158	No evidence that major mtDNA European haplogroups confer risk to schizophrenia. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 414-421.	1.7	25
159	Results of the 2003–2004 GEP-ISFG collaborative study on mitochondrial DNA: Focus on the mtDNA profile of a mixed semen-saliva stain. Forensic Science International, 2006, 160, 157-167.	2.2	24
160	Analysis of body fluid mixtures by mtDNA sequencing: An inter-laboratory study of the GEP-ISFG working group. Forensic Science International, 2007, 168, 42-56.	2,2	24
161	A cautionary note on switching mitochondrial DNA reference sequences in forensic genetics. Forensic Science International: Genetics, 2012, 6, e182-e184.	3.1	24
162	Indian Signatures in the Westernmost Edge of the European Romani Diaspora: New Insight from Mitogenomes. PLoS ONE, 2013, 8, e75397.	2.5	24

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163	The relationship between surname frequency and Y chromosome variation in Spain. European Journal of Human Genetics, 2016, 24, 120-128.	2.8	24
164	Mitochondrial Echoes of First Settlement and Genetic Continuity in El Salvador. PLoS ONE, 2009, 4, e6882.	2.5	23
165	Extraordinary claims require extraordinary evidence in asserted mtDNA biparental inheritance. Forensic Science International: Genetics, 2020, 47, 102274.	3.1	23
166	A Genome-Wide Study of Modern-Day Tuscans: Revisiting Herodotus's Theory on the Origin of the Etruscans. PLoS ONE, 2014, 9, e105920.	2.5	23
167	A multi-tissue study of immune gene expression profiling highlights the key role of the nasal epithelium in COVID-19 severity. Environmental Research, 2022, 210, 112890.	7.5	23
168	High Mitochondrial DNA Stability in B-Cell Chronic Lymphocytic Leukemia. PLoS ONE, 2009, 4, e7902.	2.5	22
169	Testing the performance of mtSNP minisequencing in forensic samples. Forensic Science International: Genetics, 2009, 3, 261-264.	3.1	22
170	Male lineages in South American native groups: Evidence of M19 traveling south. American Journal of Physical Anthropology, 2011, 146, 188-196.	2.1	22
171	Analysis of a claimed distant relationship in a deficient pedigree using high density SNP data. Forensic Science International: Genetics, 2012, 6, 350-353.	3.1	22
172	Nonbinary single-nucleotide polymorphism markers. International Congress Series, 2004, 1261, 27-29.	0.2	21
173	The search of â€~novel' mtDNA mutations in hypertrophic cardiomyopathy: MITOMAPping as a risk factor. International Journal of Cardiology, 2008, 126, 439-442.	1.7	21
174	2006 GEP-ISFG collaborative exercise on mtDNA: reflections about interpretation, artefacts, and DNA mixtures. Forensic Science International: Genetics, 2008, 2, 126-133.	3.1	21
175	Mild enteropathy as a cause of iron-deficiency anaemia of previously unknown origin. Digestive and Liver Disease, 2011, 43, 448-453.	0.9	20
176	Reassessing the role of mitochondrial DNA mutations in autism spectrum disorder. BMC Medical Genetics, 2011, 12, 50.	2.1	20
177	Prevalence and Natural History of Microscopic Colitis: A Population-Based Study With Long-term Clinical Follow-up in Terrassa, Spain. Journal of Crohn's and Colitis, 2016, 10, 805-811.	1.3	20
178	Revealing latitudinal patterns of mitochondrial DNA diversity in Chileans. Forensic Science International: Genetics, 2016, 20, 81-88.	3.1	20
179	A 2-transcript host cell signature distinguishes viral from bacterial diarrhea and it is influenced by the severity of symptoms. Scientific Reports, 2018, 8, 8043.	3.3	20
180	Y-chromosome STR haplotypes from a Western Mediterranean population sample. Forensic Science International, 2001, 119, 254-257.	2.2	19

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