Ferran Casals

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

71	9,531	25	78
papers	citations	h-index	g-index
78	11,020 ext. citations	8.8	5.41
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
71	A forensic population database in El Salvador: 58 STRs and 94 SNPs. <i>Forensic Science International: Genetics</i> , 2021 , 57, 102646	4.3	O
70	Assessment of the gene mosaicism burden in blood and its implications for immune disorders. <i>Scientific Reports</i> , 2021 , 11, 12940	4.9	0
69	Genome-wide postnatal changes in immunity following fetal inflammatory response. <i>FEBS Journal</i> , 2021 , 288, 2311-2331	5.7	
68	Second GHEP-ISFG exercise for DVI: "DNA-led" victims Udentification in a simulated air crash. <i>Forensic Science International: Genetics</i> , 2021 , 53, 102527	4.3	1
67	Historical human remains identification through maternal and paternal genetic signatures in a founder population with extensive genealogical record. <i>American Journal of Physical Anthropology</i> , 2020 , 171, 645-658	2.5	2
66	FHLdb: A Comprehensive Database on the Molecular Basis of Familial Hemophagocytic Lymphohistiocytosis. <i>Frontiers in Immunology</i> , 2020 , 11, 107	8.4	3
65	The first GHEP-ISFG collaborative exercise on forensic applications of massively parallel sequencing. <i>Forensic Science International: Genetics</i> , 2020 , 49, 102391	4.3	2
64	Severe Autoinflammatory Manifestations and Antibody Deficiency Due to Novel Hypermorphic PLCG2 Mutations. <i>Journal of Clinical Immunology</i> , 2020 , 40, 987-1000	5.7	12
63	Genetic diagnosis of autoinflammatory disease patients using clinical exome sequencing. <i>European Journal of Medical Genetics</i> , 2020 , 63, 103920	2.6	9
62	Flow Sorting Enrichment and Nanopore Sequencing of Chromosome 1 From a Chinese Individual. <i>Frontiers in Genetics</i> , 2019 , 10, 1315	4.5	1
61	Genetic identification of Spanish civil war victims. The state of the art in Catalonia (Northeastern Spain). <i>Forensic Science International: Genetics Supplement Series</i> , 2019 , 7, 419-421	0.5	3
60	The BIOMEPOC Project: Personalized Biomarkers and Clinical Profiles in Chronic Obstructive Pulmonary Disease. <i>Archivos De Bronconeumologia</i> , 2019 , 55, 93-99	0.7	11
59	Unexpected relevant role of gene mosaicism in patients with primary immunodeficiency diseases. Journal of Allergy and Clinical Immunology, 2019 , 143, 359-368	11.5	29
58	The impact of endogenous content, replicates and pooling on genome capture from faecal samples. <i>Molecular Ecology Resources</i> , 2018 , 18, 319-333	8.4	24
57	Evaluating the Genetics of Common Variable Immunodeficiency: Monogenetic Model and Beyond. <i>Frontiers in Immunology</i> , 2018 , 9, 636	8.4	81
56	Genomes reveal marked differences in the adaptive evolution between orangutan species. <i>Genome Biology</i> , 2018 , 19, 193	18.3	5
55	Reply to Wo evidence for unknown archaic ancestry in South AsiaU <i>Nature Genetics</i> , 2018 , 50, 1637-163	9 36.3	2

(2013-2018)

54	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4-insufficient subjects. <i>Journal of Allergy and Clinical Immunology</i> , 2018 , 142, 1932-1946	11.5	204
53	Y-chromosomal sequences of diverse Indian populations and the ancestry of the Andamanese. <i>Human Genetics</i> , 2017 , 136, 499-510	6.3	11
52	Length and repeat-sequence variation in 58 STRs and 94 SNPs in two Spanish populations. <i>Forensic Science International: Genetics</i> , 2017 , 30, 66-70	4.3	17
51	Morphometric, Behavioral, and Genomic Evidence for a New Orangutan Species. <i>Current Biology</i> , 2017 , 27, 3487-3498.e10	6.3	116
50	Detection of genomic rearrangements from targeted resequencing data in Parkinson'd disease patients. <i>Movement Disorders</i> , 2017 , 32, 165-169	7	10
49	Genomic analysis of Andamanese provides insights into ancient human migration into Asia and adaptation. <i>Nature Genetics</i> , 2016 , 48, 1066-70	36.3	88
48	Chimpanzee genomic diversity reveals ancient admixture with bonobos. <i>Science</i> , 2016 , 354, 477-481	33.3	139
47	Natural Selection in the Great Apes. <i>Molecular Biology and Evolution</i> , 2016 , 33, 3268-3283	8.3	44
46	Genetic Load of Loss-of-Function Polymorphic Variants in Great Apes. <i>Genome Biology and Evolution</i> , 2016 , 8, 871-7	3.9	15
45	An assessment of a massively parallel sequencing approach for the identification of individuals from mass graves of the Spanish Civil War (1936-1939). <i>Electrophoresis</i> , 2016 , 37, 2841-2847	3.6	16
44	Extreme selective sweeps independently targeted the X chromosomes of the great apes. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 6413-8	11.5	52
43	Mendelian genes for Parkinson's disease contribute to the sporadic forms of the disease. <i>Human Molecular Genetics</i> , 2015 , 24, 2023-34	5.6	20
42	Population and genomic lessons from genetic analysis of two Indian populations. <i>Human Genetics</i> , 2014 , 133, 1273-87	6.3	23
41	Selective constraint, background selection, and mutation accumulation variability within and between human populations. <i>BMC Genomics</i> , 2013 , 14, 495	4.5	13
40	Great ape genetic diversity and population history. <i>Nature</i> , 2013 , 499, 471-5	50.4	574
39	Exome sequencing identifies mutations in the gene TTC7A in French-Canadian cases with hereditary multiple intestinal atresia. <i>Journal of Medical Genetics</i> , 2013 , 50, 324-9	5.8	93
38	Rare allelic forms of PRDM9 associated with childhood leukemogenesis. <i>Genome Research</i> , 2013 , 23, 419-30	9.7	37
37	Whole-exome sequencing reveals a rapid change in the frequency of rare functional variants in a founding population of humans. <i>PLoS Genetics</i> , 2013 , 9, e1003815	6	52

36	Long-standing balancing selection in the THBS4 gene: influence on sex-specific brain expression and gray matter volumes in Alzheimer disease. <i>Human Mutation</i> , 2013 , 34, 743-53	4.7	6
35	Next-generation sequencing approaches for genetic mapping of complex diseases. <i>Journal of Neuroimmunology</i> , 2012 , 248, 10-22	3.5	15
34	Genetics. Human genetic variation, shared and private. <i>Science</i> , 2012 , 337, 39-40	33.3	27
33	Similarity in recombination rate estimates highly correlates with genetic differentiation in humans. <i>PLoS ONE</i> , 2011 , 6, e17913	3.7	13
32	Variation in genome-wide mutation rates within and between human families. <i>Nature Genetics</i> , 2011 , 43, 712-4	36.3	404
31	Genetic adaptation of the antibacterial human innate immunity network. <i>BMC Evolutionary Biology</i> , 2011 , 11, 202	3	20
30	Recent human evolution has shaped geographical differences in susceptibility to disease. <i>BMC Genomics</i> , 2011 , 12, 55	4.5	22
29	A population genetic approach to mapping neurological disorder genes using deep resequencing. <i>PLoS Genetics</i> , 2011 , 7, e1001318	6	72
28	A targeted association study of immunity genes and networks suggests novel associations with placental malaria infection. <i>PLoS ONE</i> , 2011 , 6, e24996	3.7	6
27	A map of human genome variation from population-scale sequencing. <i>Nature</i> , 2010 , 467, 1061-73	50.4	6142
27	A map of human genome variation from population-scale sequencing. <i>Nature</i> , 2010 , 467, 1061-73 Serotype-specific pneumococcal disease may be influenced by mannose-binding lectin deficiency. <i>European Respiratory Journal</i> , 2010 , 36, 856-63	50.4	6142
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26	Serotype-specific pneumococcal disease may be influenced by mannose-binding lectin deficiency. European Respiratory Journal, 2010, 36, 856-63 Direct measure of the de novo mutation rate in autism and schizophrenia cohorts. American Journal	13.6	11
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26 25 24	Serotype-specific pneumococcal disease may be influenced by mannose-binding lectin deficiency. European Respiratory Journal, 2010, 36, 856-63 Direct measure of the de novo mutation rate in autism and schizophrenia cohorts. American Journal of Human Genetics, 2010, 87, 316-24 Human pseudogenes of the ABO family show a complex evolutionary dynamics and loss of function. Glycobiology, 2009, 19, 583-91 Interrogating 11 fast-evolving genes for signatures of recent positive selection in worldwide	13.6 11 5.8	11 181 12
26 25 24 23	Serotype-specific pneumococcal disease may be influenced by mannose-binding lectin deficiency. European Respiratory Journal, 2010, 36, 856-63 Direct measure of the de novo mutation rate in autism and schizophrenia cohorts. American Journal of Human Genetics, 2010, 87, 316-24 Human pseudogenes of the ABO family show a complex evolutionary dynamics and loss of function. Glycobiology, 2009, 19, 583-91 Interrogating 11 fast-evolving genes for signatures of recent positive selection in worldwide human populations. Molecular Biology and Evolution, 2009, 26, 2285-97 A variant in the gene FUT9 is associated with susceptibility to placental malaria infection. Human	13.6 11 5.8 8.3 5.6	11 181 12 20
26 25 24 23 22	Serotype-specific pneumococcal disease may be influenced by mannose-binding lectin deficiency. European Respiratory Journal, 2010, 36, 856-63 Direct measure of the de novo mutation rate in autism and schizophrenia cohorts. American Journal of Human Genetics, 2010, 87, 316-24 Human pseudogenes of the ABO family show a complex evolutionary dynamics and loss of function. Glycobiology, 2009, 19, 583-91 Interrogating 11 fast-evolving genes for signatures of recent positive selection in worldwide human populations. Molecular Biology and Evolution, 2009, 26, 2285-97 A variant in the gene FUT9 is associated with susceptibility to placental malaria infection. Human Molecular Genetics, 2009, 18, 3136-44	13.6 11 5.8 8.3 5.6	11 181 12 20

18	Balancing selection is the main force shaping the evolution of innate immunity genes. <i>Journal of Immunology</i> , 2008 , 181, 1315-22	5.3	150
17	SNP analysis to results (SNPator): a web-based environment oriented to statistical genomics analyses upon SNP data. <i>Bioinformatics</i> , 2008 , 24, 1643-4	7.2	57
16	Is there selection for the pace of successive inactivation of the arpAT gene in primates?. <i>Journal of Molecular Evolution</i> , 2008 , 67, 23-8	3.1	2
15	Evolutionary analysis of genes of two pathways involved in placental malaria infection. <i>Human Genetics</i> , 2008 , 123, 343-57	6.3	6
14	Signatures of selection in the human olfactory receptor OR5I1 gene. <i>Molecular Biology and Evolution</i> , 2008 , 25, 144-54	8.3	24
13	Extreme individual marker F(ST) values do not imply population-specific selection in humans: the NRG1 example. <i>Human Genetics</i> , 2007 , 121, 759-62	6.3	20
12	Testing chromosomal phylogenies and inversion breakpoint reuse in Drosophila. <i>Genetics</i> , 2007 , 175, 167-77	4	22
11	Abundance and chromosomal distribution of six Drosophila buzzatii transposons: BuT1, BuT2, BuT3, BuT4, BuT5, and BuT6. <i>Chromosoma</i> , 2006 , 115, 403-12	2.8	15
10	A BAC-based physical map of the Drosophila buzzatii genome. <i>Genome Research</i> , 2005 , 15, 885-92	9.7	20
9	Molecular characterization and chromosomal distribution of Galileo, Kepler and Newton, three foldback transposable elements of the Drosophila buzzatii species complex. <i>Genetics</i> , 2005 , 169, 2047	-5 9	23
8	Duplicative and conservative transpositions of larval serum protein 1 genes in the genus Drosophila. <i>Genetics</i> , 2004 , 168, 253-64	4	12
7	The foldback-like transposon Galileo is involved in the generation of two different natural chromosomal inversions of Drosophila buzzatii. <i>Molecular Biology and Evolution</i> , 2003 , 20, 674-85	8.3	64
6	A new split of the Hox gene complex in Drosophila: relocation and evolution of the gene labial. <i>Molecular Biology and Evolution</i> , 2003 , 20, 2042-54	8.3	36
5	Low occurrence of gene transposition events during the evolution of the genus Drosophila. <i>Evolution; International Journal of Organic Evolution</i> , 2003 , 57, 1325-35	3.8	39
4	LOW OCCURRENCE OF GENE TRANSPOSITION EVENTS DURING THE EVOLUTION OF THE GENUS DROSOPHILA. <i>Evolution; International Journal of Organic Evolution</i> , 2003 , 57, 1325	3.8	10
3	How malleable is the eukaryotic genome? Extreme rate of chromosomal rearrangement in the genus Drosophila. <i>Genome Research</i> , 2001 , 11, 230-9	9.7	129
2	Further confirmation for unknown archaic ancestry in Andaman and South Asia		2
1	Next-generation sequencing for rare diseases231-242		