List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/595781/publications.pdf Version: 2024-02-01

		8159	4419
211	31,653	76	172
papers	citations	h-index	g-index
221	221	221	40054
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	An integrated map of genetic variation from 1,092 human genomes. Nature, 2012, 491, 56-65.	13.7	7,199
2	Production of monoclonal antibodies to group A erythrocytes, HLA and other human cell surface antigens-new tools for genetic analysis. Cell, 1978, 14, 9-20.	13.5	1,905
3	A serine/threonine kinase gene defective in Peutz–Jeghers syndrome. Nature, 1998, 391, 184-187.	13.7	1,451
4	Localization of the gene for familial adenomatous polyposis on chromosome 5. Nature, 1987, 328, 614-616.	13.7	1,362
5	Common and rare variants in multifactorial susceptibility to common diseases. Nature Genetics, 2008, 40, 695-701.	9.4	1,010
6	Target genes of Â-catenin-T cell-factor/lymphoid-enhancer-factor signaling in human colorectal carcinomas. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 1603-1608.	3.3	764
7	Chromosome 5 allele loss in human colorectal carcinomas. Nature, 1987, 328, 616-619.	13.7	586
8	Monoclonal antibodies to epithelium-specific components of the human milk fat globule membrane: Production and reaction with cells in culture. International Journal of Cancer, 1981, 28, 17-21.	2.3	571
9	Nomenclature for factors of the HLA system, 2004. Tissue Antigens, 2005, 65, 301-369.	1.0	491
10	The Eurasian Heartland: A continental perspective on Y-chromosome diversity. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 10244-10249.	3.3	445
11	Localization of a susceptibility locus for Peutz-Jeghers syndrome to 19p using comparative genomic hybridization and targeted linkage analysis. Nature Genetics, 1997, 15, 87-90.	9.4	444
12	APC mutations in sporadic colorectal tumors: A mutational "hotspot" and interdependence of the "two hits". Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 3352-3357.	3.3	441
13	Â-Catenin mutations in cell lines established from human colorectal cancers. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 10330-10334.	3.3	435
14	The mutation rate and cancer. Proceedings of the National Academy of Sciences of the United States of America, 1996, 93, 14800-14803.	3.3	424
15	The fine-scale genetic structure of the British population. Nature, 2015, 519, 309-314.	13.7	416
16	Cancer stem cells from colorectal cancer-derived cell lines. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 3722-3727.	3.3	392
17	The type of somatic mutation at APC in familial adenomatous polyposis is determined by the site of the germline mutation: a new facet to Knudson's 'two-hit' hypothesis. Nature Medicine, 1999, 5, 1071-1075.	15.2	339
18	Cancer Cell Lines for Drug Discovery and Development. Cancer Research, 2014, 74, 2377-2384.	0.4	324

#	Article	IF	CITATIONS
19	Chromosomal localization of human cellular homologues of two viral oncogenes. Nature, 1982, 299, 747-749.	13.7	317
20	Colorectal Cancer Cell Lines Are Representative Models of the Main Molecular Subtypes of Primary Cancer. Cancer Research, 2014, 74, 3238-3247.	0.4	317
21	Genetic Analysis with Man–Mouse Somatic Cell Hybrids: Linkage between Human Lactate Dehydrogenase B and Peptidase B Genes. Nature, 1970, 227, 248-251.	13.7	308
22	Differentiation antigens expressed by epithelial cells in the lactating breast are also detectable in breast cancers. International Journal of Cancer, 1981, 28, 23-29.	2.3	299
23	Selection, the mutation rate and cancer: Ensuring that the tail does not wag the dog. Nature Medicine, 1999, 5, 11-12.	15.2	289
24	Genetic Analysis with Human—Mouse Somatic Cell Hybrids. Nature, 1969, 223, 358-363.	13.7	282
25	Insulin-like growth factor 1 regulates the location, stability, and transcriptional activity of beta -catenin. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 12103-12108.	3.3	261
26	A Novel Carcinoembryonic Antigen T-Cell Bispecific Antibody (CEA TCB) for the Treatment of Solid Tumors. Clinical Cancer Research, 2016, 22, 3286-3297.	3.2	260
27	Mathematical modeling of cell population dynamics in the colonic crypt and in colorectal cancer. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 4008-4013.	3.3	253
28	Molecular analysis of APC mutations in familial adenomatous polyposis and sporadic colon carcinomas. Lancet, The, 1992, 340, 626-630.	6.3	236
29	Germline Mutations in BMPR1A/ALK3 Cause a Subset of Cases of Juvenile Polyposis Syndrome and of Cowden and Bannayan-Riley-Ruvalcaba Syndromes*. American Journal of Human Genetics, 2001, 69, 704-711.	2.6	236
30	Expression of HLA antigens, β2-microglobulin and enzymes by human amniotic epithelial cells. Nature, 1982, 295, 325-327.	13.7	222
31	Assignment of the structural gene for the third component of human complement to chromosome 19 Proceedings of the National Academy of Sciences of the United States of America, 1982, 79, 5021-5025.	3.3	220
32	Histocompatibility antigens, immune responsiveness and susceptibility to disease. American Journal of Medicine, 1972, 52, 1-8.	0.6	207
33	Analysis of P53 mutations and their expression in 56 colorectal cancer cell lines. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 976-981.	3.3	202
34	The APC variants I1307K and E1317Q are associated with colorectal tumors, but not always with a family history. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 10722-10727.	3.3	194
35	Multiple rare variants in different genes account for multifactorial inherited susceptibility to colorectal adenomas. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 15992-15997.	3.3	181
36	APC mutations are sufficient for the growth of early colorectal adenomas. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 2225-2228.	3.3	169

#	Article	IF	CITATIONS
37	Array Comparative Genomic Hybridization Analysis of Colorectal Cancer Cell Lines and Primary Carcinomas. Cancer Research, 2004, 64, 4817-4825.	0.4	168
38	Failure of programmed cell death and differentiation as causes of tumors: some simple mathematical models Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 11130-11134.	3.3	167
39	Sequence of an HLA-DR α-chain cDNA clone and intron-exon organization of the corresponding gene. Nature, 1982, 299, 750-752.	13.7	162
40	SMAD4 mutations in colorectal cancer probably occur before chromosomal instability, but after divergence of the microsatellite instability pathway. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 9719-9723.	3.3	162
41	Multigene amplification and massively parallel sequencing for cancer mutation discovery. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 9387-9392.	3.3	159
42	Spectral karyotyping suggests additional subsets of colorectal cancers characterized by pattern of chromosome rearrangement. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 2538-2543.	3.3	152
43	DNA mismatch repair genes and colorectal cancer. Gut, 2000, 47, 148-153.	6.1	151
44	Human gene mapping using an X/autosome translocation. Somatic Cell Genetics, 1976, 2, 125-140.	2.7	148
45	Intestinal trefoil factor controls the expression of the adenomatous polyposis coli-catenin and the E-cadherin-catenin complexes in human colon carcinoma cells. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 3122-3127.	3.3	148
46	Genetic steps in colorectal cancer. Nature Genetics, 1994, 6, 217-219.	9.4	147
47	Genetics of colorectal cancer: hereditary aspects and overview of colorectal tumorigenesis. British Medical Bulletin, 2002, 64, 27-43.	2.7	145
48	Mutations in DPC4 (SMAD4) cause juvenile polyposis syndrome, but only account for a minority of cases. Human Molecular Genetics, 1998, 7, 1907-1912.	1.4	142
49	Beta 2-microglobulin gene mutations: a study of established colorectal cell lines and fresh tumors Proceedings of the National Academy of Sciences of the United States of America, 1994, 91, 4751-4755.	3.3	133
50	Hypermethylation of the promoter region of the E-cadherin gene (CDH1) in sporadic and ulcerative colitis associated colorectal cancer. Gut, 2001, 48, 367-371.	6.1	128
51	Whole-gene APC deletions cause classical familial adenomatous polyposis, but not attenuated polyposis or "multiple" colorectal adenomas. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 2954-2958.	3.3	127
52	People of the British Isles: preliminary analysis of genotypes and surnames in a UK-control population. European Journal of Human Genetics, 2012, 20, 203-210.	1.4	126
53	Germline APC variants in patients with multiple colorectal adenomas, with evidence for the particular importance of E1317Q. Human Molecular Genetics, 2000, 9, 2215-2221.	1.4	125
54	X-inactivation patch size in human female tissue confounds the assessment of tumor clonality. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 3311-3314.	3.3	121

#	Article	IF	CITATIONS
55	Mutated epithelial cadherin is associated with increased tumorigenicity and loss of adhesion and of responsiveness to the motogenic trefoil factor 2 in colon carcinoma cells. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 2316-2321.	3.3	117
56	Analysis of genetic and phenotypic heterogeneity in juvenile polyposis. Gut, 2000, 46, 656-660.	6.1	117
57	Subcellular Separation and Molecular Nature of Human Histocompatibility Antigens (HL-A). Nature, 1974, 247, 457-461.	13.7	116
58	Dietary fat influences on polyp phenotype in multiple intestinal neoplasia mice. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 3308-3313.	3.3	112
59	Detection of circulating tumour cells in peripheral blood with an automated scanning fluorescence microscope. British Journal of Cancer, 2008, 99, 789-795.	2.9	111
60	How Many Mutations in a Cancer?. American Journal of Pathology, 2002, 160, 755-758.	1.9	110
61	The independent expression of HLA and? 2-microglobulin on human-mouse hybrids. Somatic Cell Genetics, 1976, 2, 483-496.	2.7	102
62	CDX1 is an important molecular mediator of Barrett's metaplasia. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 7565-7570.	3.3	101
63	Mechanisms of inactivation of mismatch repair genes in human colorectal cancer cell lines: The predominant role of hMLH1. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 10296-10301.	3.3	100
64	Hypoxia and lineage specification of cell line-derived colorectal cancer stem cells. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 4382-4387.	3.3	100
65	Functional expression of HLA-DP genes transfected into mouse fibroblasts. Nature, 1985, 313, 61-64.	13.7	98
66	An Ancestral Ashkenazi Haplotype at the HMPS/CRAC1 Locus on 15q13–q14 Is Associated with Hereditary Mixed Polyposis Syndrome. American Journal of Human Genetics, 2003, 72, 1261-1267.	2.6	98
67	Gastrointestinal differentiation marker Cytokeratin 20 is regulated by homeobox gene CDX1. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 1936-1941.	3.3	95
68	Polygenic inheritance, GWAS, polygenic risk scores, and the search for functional variants. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 18924-18933.	3.3	95
69	Isolation and N-terminal amino acid sequence of membrane-bound human HLA-A and HLA-B antigens. Nature, 1976, 261, 200-205.	13.7	93
70	Somatic Mutations in the Peutz-Jegners (LKB1/STKII) Gene in Sporadic Malignant Melanomas. Journal of Investigative Dermatology, 1999, 112, 509-511.	0.3	93
71	Analysis of chromosomal instability in human colorectal adenomas with two mutational hits at APC. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 16910-16915.	3.3	89
72	5-Fluorouracil response in a large panel of colorectal cancer cell lines is associated with mismatch repair deficiency. British Journal of Cancer, 2010, 103, 340-346.	2.9	88

#	Article	IF	CITATIONS
73	Characterization and chromosomal assignment of a human cell surface antigen defined by the monoclonal antibody AUAI. International Journal of Cancer, 1986, 38, 631-636.	2.3	86
74	Nomenclature for factors of the HLA system, 1995. Human Immunology, 1995, 43, 149-164.	1.2	85
75	The peopling of Europe and the cautionary tale of Y chromosome lineage R-M269. Proceedings of the Royal Society B: Biological Sciences, 2012, 279, 884-892.	1.2	84
76	Detection of human cancer in an animal model using radio-labelled tumour-associated monoclonal antibodies. British Journal of Cancer, 1982, 46, 1-8.	2.9	79
77	On the increase of chromosome mutations under random mating. Theoretical Population Biology, 1976, 9, 260-281.	0.5	74
78	Myofibroblasts are distinguished from activated skin fibroblasts by the expression of AOC3 and other associated markers. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E2162-71.	3.3	73
79	A comparison of the genetic pathways involved in the pathogenesis of three types of colorectal cancer. , 1998, 184, 148-152.		70
80	The HLA system and the analysis of multifactorial genetic disease. Trends in Genetics, 1995, 11, 493-498.	2.9	69
81	Separation of cancer cells from white blood cells by pinched flow fractionation. Lab on A Chip, 2015, 15, 4598-4606.	3.1	66
82	MHC antigens and cancer: implications for T-cell surveillance. Current Opinion in Immunology, 1992, 4, 613-618.	2.4	65
83	Defects in mismatch repair occur afterAPC mutations in the pathogenesis of sporadic colorectal tumours. , 1998, 11, 114-120.		64
84	Genetic Instability Is Not a Requirement for Tumor Development. Cancer Research, 2008, 68, 3558-3561.	0.4	64
85	Genetics of the human face: Identification of large-effect single gene variants. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E676-E685.	3.3	64
86	Cancer genetics: colorectal cancer as a model. Journal of Human Genetics, 2006, 51, 391-396.	1.1	63
87	HIV TESTING ON ALL PREGNANT WOMEN. Lancet, The, 1987, 330, 1277.	6.3	62
88	Rapid isolation of human chromosome-specific DNA probes from a somatic cell hybrid. Genomics, 1990, 7, 257-263.	1.3	62
89	Mitochondrial DNA of Human–Mouse Cell Hybrids. Nature, 1971, 234, 560-562.	13.7	61
90	Loss of CDX1 expression in colorectal carcinoma: Promoter methylation, mutation, and loss of heterozygosity analyses of 37 cell lines. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 574-579.	3.3	61

#	Article	IF	CITATIONS
91	Rare genetic variants and the risk of cancer. Current Opinion in Genetics and Development, 2010, 20, 262-267.	1.5	61
92	Loss of HLA class-I alleles, heavy chains and β2-microglobulin in colorectal cancer. International Journal of Cancer, 1992, 51, 379-385.	2.3	60
93	On the proportion of cancer stem cells in a tumour. Journal of Theoretical Biology, 2010, 266, 708-711.	0.8	59
94	The Evolution of Overdominance: Natural Selection and Heterozygote Advantage. Nature, 1961, 190, 7-12.	13.7	56
95	The E-cadherin gene (CDH1) variants T340A and L599V in gastric and colorectal cancer patients in Korea. Gut, 2000, 47, 262-267.	6.1	56
96	HLA-A locus alleles identified by sequence specific PCR. Lancet, The, 1993, 341, 121-122.	6.3	55
97	Use of SSCP analysis to identify germline mutations in HNPCC families fulfilling the Amsterdam criteria. Human Genetics, 1997, 99, 219-224.	1.8	53
98	An isolated \hat{I}^2 1 exon next to the DR $\hat{I}\pm$ gene in the HLA-D region. Immunogenetics, 1986, 23, 172-180.	1.2	50
99	An update to HLA Nomenclature, 2010. Bone Marrow Transplantation, 2010, 45, 846-848.	1.3	48
100	An immunohistological study of testicular germ cell tumours using two different monoclonal antibodies against placental alkaline phosphatase. British Journal of Cancer, 1984, 49, 11-15.	2.9	47
101	Transforming growth factor stimulation of colorectal cancer cell lines: Type II receptor bypass and changes in adhesion molecule expression. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 3087-3091.	3.3	47
102	Nomenclature for factors of the HLA system, 2002. International Journal of Immunogenetics, 2002, 29, 463-515.	1.2	47
103	Relationship of Centromeric Heterochromatin to Fluorescent Banding Patterns of Metaphase Chromosomes in the Mouse. Nature, 1971, 231, 503-506.	13.7	46
104	Familial adenomatous polyposis (FAP) and its gene, APC. Cytogenetic and Genome Research, 1999, 86, 99-104.	0.6	46
105	Cell growth, global phosphotyrosine elevation, and c-Met phosphorylation through Src family kinases in colorectal cancer cells. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 2358-2362.	3.3	46
106	Genotyping Possible Polymorphic Variants of Human Mismatch Repair Genes in Healthy Korean Individuals and Sporadic Colorectal Cancer Patients. Familial Cancer, 2002, 3, 129-137.	0.9	44
107	Genetic basis of variation in adenoma multiplicity in ApcMin/+ Mom1S mice. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 2868-2873.	3.3	43
108	Mutations in the AXIN1 Gene in Advanced Prostate Cancer. European Urology, 2009, 56, 486-494.	0.9	41

#	Article	IF	CITATIONS
109	Direct and immune mediated antibody targeting of <i>ERBB</i> receptors in a colorectal cancer cell-line panel. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 21046-21051.	3.3	41
110	Stem Cell Differentiation and Lumen Formation in Colorectal Cancer Cell Lines and Primary Tumors. Cancer Research, 2013, 73, 5798-5809.	0.4	41
111	GENETICS OF "4―AND "LA―HUMAN LEUKOCYTE GROUPS*. Annals of the New York Academy of Science 2006, 129, 473-489.	25, 1.8	40
112	CpG island clones from a deletion encompassing the gene for adenomatous polyposis coli Proceedings of the National Academy of Sciences of the United States of America, 1989, 86, 10118-10122.	3.3	39
113	Monoclonal antibody assay of serum placental alkaline phosphatase in the monitoring of testicular tumours. British Journal of Cancer, 1985, 51, 641-644.	2.9	38
114	Stromal uptake and transmission of acid is a pathway for venting cancer cell-generated acid. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E5344-53.	3.3	38
115	Human intraspecific somatic cell hybrids: A genetic and karyotypic analysis of crosses between lymphocytes and D98/AH-2. Somatic Cell Genetics, 1975, 1, 41-64.	2.7	36
116	Allele loss in colorectal cancer at the Cowden disease/Juvenile Polyposis locus on 10q. Cancer Genetics and Cytogenetics, 1997, 97, 64-69.	1.0	36
117	A rapid micro method for counting cells "in situ―using a fluorogenic alkaline phosphatase enzyme assay. In Vitro Cellular & Developmental Biology, 1989, 25, 105-108.	1.0	35
118	A high-frequency polymorphism in exon 6 of the CD45 tyrosine phosphatase gene (PTPRC) resulting in altered isoform expression. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 5997-6002.	3.3	34
119	Disease associations and altered immune function in CD45 138G variant carriers. Human Molecular Genetics, 2004, 13, 2377-2384.	1.4	33
120	Myofibroblast activation in colorectal cancer lymph node metastases. British Journal of Cancer, 2013, 108, 2106-2115.	2.9	33
121	GSTM1 and GSTT1 polymorphisms as modifiers of age at diagnosis of hereditary nonpolyposis colorectal cancer (HNPCC) in a homogeneous cohort of individuals carrying a single predisposing mutation. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2006, 602, 175-181	0.4	30
122	HLA Haplotype Associations with Disease. Tissue Antigens, 1979, 13, 91-102.	1.0	30
123	A pericentric inversion of chromosome six in a patient with Peutz-Jeghers' syndrome and the use of FISH to localise the breakpoints on a genetic map. Human Genetics, 1996, 98, 125-128.	1.8	29
124	MYH biallelic mutation can inactivate the two genetic pathways of colorectal cancer by APC or MLH1 transversions. Familial Cancer, 2010, 9, 589-594.	0.9	29
125	The Irish DNA Atlas: Revealing Fine-Scale Population Structure and History within Ireland. Scientific Reports, 2017, 7, 17199.	1.6	29
126	Expression of a single-chain HLA class I molecule in a human cell line: presentation of exogenous peptide and processed antigen to cytotoxic T lymphocytes Proceedings of the National Academy of Sciences of the United States of America, 1996, 93, 236-240.	3.3	28

#	Article	IF	CITATIONS
127	Microsatellite Instability in Benign Skin Lesions in Hereditary Non-Polyposis Colorectal Cancer Syndrome. Journal of Investigative Dermatology, 1999, 113, 901-905.	0.3	28
128	Nomenclature for factors of the HLA system 1984. Immunogenetics, 1984, 20, 593-601.	1.2	27
129	New vector for transfer of yeast artificial chromosomes to mammalian cells. Somatic Cell and Molecular Genetics, 1993, 19, 161-169.	0.7	27
130	Genetic Characterization of Human Populations: From ABO to a Genetic Map of the British People. Genetics, 2015, 199, 267-279.	1.2	27
131	Introduction of amyc reporter tag to improve the quality of mutation detection using the protein truncation test. , 1997, 9, 172-176.		26
132	Targeted killing of colorectal cancer cell lines by a humanised IgG1 monoclonal antibody that binds to membrane-bound carcinoembryonic antigen. British Journal of Cancer, 2008, 98, 1217-1225.	2.9	26
133	Humanised IgC1 antibody variants targeting membrane-bound carcinoembryonic antigen by antibody-dependent cellular cytotoxicity and phagocytosis. British Journal of Cancer, 2009, 101, 1758-1768.	2.9	26
134	Comprehensive assessment of variation at the transforming growth factor Î ² type 1 receptor locus and colorectal cancer predisposition. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 7858-7862.	3.3	26
135	A sensitive micro-immunoassay using β-galactosidase/ anti-β-galactosidase complexes. Journal of Immunological Methods, 1987, 97, 19-27.	0.6	25
136	Distribution of carcinoembryonic antigen and biologic behavior in colorectal carcinoma. Diseases of the Colon and Rectum, 1999, 42, 640-648.	0.7	25
137	CDX2 mutations do not account for juvenile polyposis or Peutz–Jeghers syndrome and occur infrequently in sporadic colorectal cancers. British Journal of Cancer, 2001, 84, 1314-1316.	2.9	25
138	Effects of Maternal Age on the Incidence of Congenital Abnormalities in Mouse and Man. Nature, 1961, 190, 1134-1135.	13.7	24
139	Linkage disequilibrium and age of HLA region SNPs in relation to classic HLA gene alleles within Europe. European Journal of Human Genetics, 2010, 18, 924-932.	1.4	24
140	HLA today. Human Immunology, 1986, 17, 490-503.	1.2	23
141	HLA: what's in a name?. Tissue Antigens, 1997, 49, 293-296.	1.0	23
142	Dsh Homolog DVL3 Mediates Resistance to IGFIR Inhibition by Regulating IGF-RAS Signaling. Cancer Research, 2014, 74, 5866-5877.	0.4	23
143	Characterization and mapping of microdissected genomic clones from the adenomatous polyposis coli (APC) region. Genomics, 1991, 11, 247-251.	1.3	22
144	Altered CD45 expression in C77G carriers influences immune function and outcome of hepatitis C infection. Journal of Medical Genetics, 2006, 43, 678-684.	1.5	20

#	Article	IF	CITATIONS
145	Cytostatic drug treatment causes seeding of gene promoter methylation. European Journal of Cancer, 2007, 43, 947-954.	1.3	19
146	Public Understanding of Science: The BA, the Royal Society and COPUS. Notes and Records of the Royal Society, 2010, 64, .	0.1	19
147	Fine Mapping of Probes in the Adenomatous Polyposis Coli Region of Chromosome 5 by In Situ Hybridization. Genes Chromosomes and Cancer, 1991, 3, 382-389.	1.5	17
148	A somatic cell hybrid panel for regional mapping of human chromosome 18. Genomics, 1992, 14, 431-436.	1.3	17
149	Replication error deficient and proficient colorectal cancer gene expression differences caused by 3′UTR polyT sequence deletions. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 21058-21063.	3.3	17
150	IMMUNOLOGY AND THE FETUS. Lancet, The, 1978, 311, 326-327.	6.3	15
151	T-cell immune responses to cancer— A new look. Human Immunology, 1991, 30, 259-261.	1.2	15
152	PLAP -CAR T nbsp cells mediate high specific cytotoxicity against colon cancer cells. Frontiers in Bioscience - Landmark, 2020, 25, 1765-1786.	3.0	15
153	PCR-based detection of two Mspl Polymorphic sites at D18S8. Nucleic Acids Research, 1991, 19, 6983-6983.	6.5	14
154	Distribution and Quantity of Leukocyte Antigens in the Formed Elements of the Blood. Transfusion, 1966, 6, 193-204.	0.8	13
155	Bristol Cancer Help Centre. Lancet, The, 1990, 336, 1185-1188.	6.3	13
156	Immune responses in advanced colorectal cancer following repeated intradermal vaccination with the anti-CEA murine monoclonal antibody, PR1A3: results of a phase I study. International Journal of Colorectal Disease, 2005, 20, 403-414.	1.0	13
157	DNA mismatch repair in lymphoblastoid cells from hereditary non-polyposis colorectal cancer (HNPCC) patients is normal under conditions of rapid cell division and increased mutational load. Mutation Research DNA Repair, 1997, 383, 177-182.	3.8	12
158	Alkaline-mediated differential interaction (AMDI): A simple automatable single-nucleotide polymorphism assay. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 2694-2697.	3.3	12
159	The human chromosome content in human × rodent somatic cell hybrids analyzed by a screening technique using Alu PCR. Genomics, 1991, 10, 186-192.	1.3	11
160	RA Fisher, statistician and geneticist extraordinary: a personal view. International Journal of Epidemiology, 2003, 32, 938-942.	0.9	11
161	Carcino-embryonic antigen may function as a chemo-attractant in colorectal-carcinoma cell lines. , 1999, 82, 880-885.		10
162	Antibody targeting studies in a transgenic murine model of spontaneous colorectal tumors. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 10256-10260.	3.3	10

#	Article	IF	CITATIONS
163	Geographical distribution and disease associations of the CD45 exon 6 138G variant. Immunogenetics, 2006, 58, 235-239.	1.2	10
164	Growth factor requirements of human colorectal tumour cells: Relations to cellular differentiation. European Journal of Cancer & Clinical Oncology, 1991, 27, 1680-1684.	0.9	9
165	Detection of a 4-bp Insertion (CACA) Functional Polymorphism at Nucleotide 241 of the Cellular Adhesion Regulatory Molecule CMAR (Formerly CAR). Genomics, 1994, 19, 181-182.	1.3	9
166	Enhancement of colorectal tumor targeting using a novel biparatopic monoclonal antibody against carcinoembryonic antigen in experimental radioimmunoguided surgery. International Journal of Cancer, 2002, 97, 542-547.	2.3	9
167	CD45 variant alleles: possibly increased frequency of a novel exon 4 CD45 polymorphism in HIV seropositive Ugandans. Immunogenetics, 2004, 56, 107-110.	1.2	9
168	Cyclin D1 rare variants in UK multiple adenoma and early-onset colorectal cancer patients. Journal of Human Genetics, 2011, 56, 58-63.	1.1	9
169	Role of rare variants in undetermined multiple adenomatous polyposis and early-onset colorectal cancer. Journal of Human Genetics, 2012, 57, 709-716.	1.1	9
170	Subdividing Y-chromosome haplogroup R1a1 reveals Norse Viking dispersal lineages in Britain. European Journal of Human Genetics, 2021, 29, 512-523.	1.4	9
171	Histocompatibility testing international. Nature, 1975, 256, 696-697.	13.7	8
172	A new look at tumour immunology. European Journal of Cancer, 1992, 28, 1761-1762.	1.3	8
173	High-throughput class I HLA genotyping using fluorescence resonance energy transfer (FRET) probes and sequence-specific primer-polymerase chain reaction (SSP-PCR). Tissue Antigens, 1999, 54, 603-614.	1.0	8
174	Preclinical Application of Radioimmunoguided Surgery Using Anti-Carcinoembryonic Antigen Biparatopic Antibody in the Colon Cancer. European Surgical Research, 2005, 37, 36-44.	0.6	8
175	Ruggero Ceppellini: A Perspective on His Contributions to Genetics and Immunology. Frontiers in Immunology, 2019, 10, 1280.	2.2	8
176	Radioimmunodetection of cancer. European Journal of Cancer & Clinical Oncology, 1982, 18, 1221-1223.	0.9	6
177	PTPRC (CD45) variation and disease association studied using single nucleotide polymorphism tagging. Tissue Antigens, 2008, 71, 458-463.	1.0	6
178	The outstanding scientist, R.A. Fisher: his views on eugenics and race. Heredity, 2021, 126, 565-576.	1.2	6
179	A novel xenonucleic acid-mediated molecular clamping technology for early colorectal cancer screening. PLoS ONE, 2021, 16, e0244332.	1.1	6
180	Regional mapping of 22 microclones around the adenomatous polyposis coli (APC) locus on chromosome 5q. Human Genetics, 1991, 88, 112-114.	1.8	5

#	Article	IF	CITATIONS
181	A Long-Range Restriction Map of Human Chromosome 5q21-q23. Genomics, 1993, 17, 15-24.	1.3	5
182	Tumor burden and clonality in multiple intestinal neoplasia mouse/normal mouse aggregation chimeras. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 12553-12558.	3.3	5
183	A mutated HLA-A*0101 allele in the colorectal cell line HCA-7. Tissue Antigens, 2005, 66, 231-237.	1.0	5
184	Somatic selection of poorly differentiating variant stem cell clones could be a key to human ageing. Journal of Theoretical Biology, 2020, 489, 110153.	0.8	5
185	Population stratification as an explanation of IQ and ABO association (reply). Nature, 1975, 254, 363-364.	13.7	4
186	The Role of TGF?s in Controlling Cell Adhesion and Differentiation of Colon Carcinoma Cells. Annals of the New York Academy of Sciences, 1990, 593, 360-362.	1.8	4
187	Genomic and cDNA sequence analysis of the cell matrix adhesion regulator gene. Proceedings of the National Academy of Sciences of the United States of America, 1997, 94, 14578-14583.	3.3	4
188	Sam Karlin: A personal appreciation. Theoretical Population Biology, 2009, 75, 230-232.	0.5	4
189	Commentary: Connections between genetics and statistics: a commentary on Fisher's 1951 Bateson lecture'Statistical Methods in Genetics'. International Journal of Epidemiology, 2010, 39, 340-344.	0.9	4
190	Joshua Lederberg. 23 May 1925 — 2 February 2008. Biographical Memoirs of Fellows of the Royal Society, 2011, 57, 229-251.	0.1	3
191	A Mathematician's Odyssey. Annual Review of Genomics and Human Genetics, 2015, 16, 1-29.	2.5	3
192	Molecular analysis of the HLA-D region genes. Human Immunology, 1983, 8, 105-112.	1.2	2
193	Isolation of a polymorphic DNA sequence (λEMBL3.2S7, D8S8) from chromosome 8. Nucleic Acids Research, 1986, 14, 9224-9224.	6.5	2
194	A new RFLP for L1.4 (D5S4) an anonymous genomic clone localised to chromosome 5. Nucleic Acids Research, 1987, 15, 6762-6762.	6.5	2
195	Where Will Genome Analysis Lead Us Forty Years On?. Annals of the New York Academy of Sciences, 1995, 758, 414-426.	1.8	2
196	Isolation of a polymorphic DNA sequence (λEMBL3.121, D14S12) from chromosome 14. Nucleic Acids Research, 1986, 14, 9225-9225.	6.5	1
197	UNDIFFERENTIATED CELLS, METAPLASIA, c-myc EXPRESSION, AND GASTROINTESTINAL CARCINOGENESIS. Lancet, The, 1989, 333, 566-567.	6.3	1
198	A fluorescence based cell adhesion assay using terasaki plates. In Vitro Cellular and Developmental Biology - Animal, 1995, 31, 81-83.	0.7	1

#	Article	IF	CITATIONS
199	Genetic testing and insurance. Nature, 1996, 380, 384-385.	13.7	1
200	Reply: In vitro and in vivo anticancer efficacy of unconjugated humanised anti-CEA monoclonal antibodies. British Journal of Cancer, 2008, 99, 839-840.	2.9	1
201	Jon Van Rood. International Journal of Immunogenetics, 2017, 44, 271-273.	0.8	1
202	Development and validation of ColoScape: A new colorectal cancer mutation detection assay Journal of Clinical Oncology, 2018, 36, e24189-e24189.	0.8	1
203	Molecular and Genetic Organization: The Future. Novartis Foundation Symposium, 1979, , 395-411.	1.2	1
204	Isolation of a DNA sequence (λEMBL3.123, D4S64) from chromosome 4 showing a BgIII polymorphism. Nucleic Acids Research, 1987, 15, 1342-1342.	6.5	0
205	Isolation of a polymorphic DNA sequence (λEMBL3.303, D2S14) from chromosome 2. Nucleic Acids Research, 1987, 15, 1341-1341.	6.5	0
206	The influence of charitable foundations on medical research policy. Journal of Medical Engineering and Technology, 1994, 18, 138-142.	0.8	0
207	Clinical features and molecular analysis of a family with multiple colon tumours and reduced plasminogen activator activity. International Journal of Colorectal Disease, 1997, 12, 1-3.	1.0	0
208	A Haldane perspective from a Fisher student. Journal of Genetics, 2017, 96, 743-746.	0.4	0
209	Connecting gene expression subtypes of colorectal cancer (CRC) with cell lines and drug resistance Journal of Clinical Oncology, 2013, 31, e14544-e14544.	0.8	0
210	A novel xenonucleic acid mediated molecular clamping technology for early colorectal cancer diagnostics Journal of Clinical Oncology, 2020, 38, e16106-e16106.	0.8	0
211	Blood Lines of the British People. , 2022, , 84-117.		0