Anne M Bowcock

List of Publications by Year
in descending order

173	21,551	13099 68	9589 142
papers	citations	h-index	g-index

186	186	186	23024
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Pathogenesis and therapy of psoriasis. Nature, 2007, 445, 866-873.	27.8	1,543
2	Frequent Mutation of BAP1 in Metastasizing Uveal Melanomas. Science, 2010, 330, 1410-1413.	12.6	1,242
3	Genome-wide scan reveals association of psoriasis with IL-23 and NF-Î²B pathways. Nature Genetics, 2009, 41, 199-204.	21.4	1,229
4	Identification of 15 new psoriasis susceptibility loci highlights the role of innate immunity. Nature Genetics, 2012, 44, 1341-1348.	21.4	848
5	JM2, encoding a fork headâ€“related protein, is mutated in X-linked autoimmunityâ€“allergic disregulation syndrome. Journal of Clinical Investigation, 2000, 106, R75-R81.	8.2	792
6	Identification of a RING protein that can interact in vivo with the BRCA1 gene product. Nature Genetics, 1996, 14, 430-440.	21.4	683
7	A Genome-Wide Association Study of Psoriasis and Psoriatic Arthritis Identifies New Disease Loci. PLoS Genetics, 2008, 4, e1000041.	3.5	572
8	Deletion of the late cornified envelope LCE3B and LCE3C genes as a susceptibility factor for psoriasis. Nature Genetics, 2009, 41, 211-215.	21.4	482
9	AGPAT2 is mutated in congenital generalized lipodystrophy linked to chromosome 9q34. Nature Genetics, 2002, 31, 21-23.	21.4	475
10	Recurrent mutations at codon 625 of the splicing factor SF3B1 in uveal melanoma. Nature Genetics, 2013, 45, 133-135.	21.4	447
11	The immunogenetics of Psoriasis: A comprehensive review. Journal of Autoimmunity, 2015, 64, 66-73.	6.5	447
12	Getting under the skin: the immunogenetics of psoriasis. Nature Reviews Immunology, 2005, 5, 699-711.	22.7	416
13	Nonlesional atopic dermatitis skin is characterized by broad terminal differentiation defects and variable immune abnormalities. Journal of Allergy and Clinical Immunology, 2011, 127, 954-964.e4.	2.9	375
14	PSORS2 Is Due to Mutations in CARD14. American Journal of Human Genetics, 2012, 90, 784-795.	6.2	365
15	The C-terminal (BRCT) Domains of BRCA1 Interact in Vivo with CtIP, a Protein Implicated in the CtBP Pathway of Transcriptional Repression. Journal of Biological Chemistry, 1998, 273, 25388-25392.	3.4	343
16	Oncogenic Mutations inGNAQOccur Early in Uveal Melanoma. , 2008, 49, 5230.		329
17	Genome-wide association analysis identifies three psoriasis susceptibility loci. Nature Genetics, 2010, 42, 1000-1004.	21.4	313
18	Rare and Common Variants in CARD14, Encoding an Epidermal Regulator of NF-kappaB, in Psoriasis. American Journal of Human Genetics, 2012, 90, 796-808.	6.2	306

#	Article	IF	Citations
19	Association analyses identify six new psoriasis susceptibility loci in the Chinese population. Nature Genetics, 2010, 42, 1005-1009.	21.4	287
20	A putative RUNX1 binding site variant between SLC9A3R1 and NAT9 is associated with susceptibility to psoriasis. Nature Genetics, 2003, 35, 349-356.	21.4	284
21	Novel mechanisms of T-cell and dendritic cell activation revealed by profiling of psoriasis on the 63,100-element oligonucleotide array. Physiological Genomics, 2003, 13, 69-78.	2.3	282
22	Mutational and Haplotype Analyses of Families with Familial Partial Lipodystrophy (Dunnigan Variety) Reveal Recurrent Missense Mutations in the Globular C-Terminal Domain of Lamin A/C. American Journal of Human Genetics, 2000, 66, 1192-1198.	6.2	260
23	Psoriasis vulgaris: cutaneous lymphoid tissue supports T-cell activation and â€˜Type 1â€™ inflammatory gene expression. Trends in Immunology, 2004, 25, 295-305.	6.8	255
24	Mutations in the Gene PRRT2 Cause Paroxysmal Kinesigenic Dyskinesia with Infantile Convulsions. Cell Reports, 2012, 1, 2-12.	6.4	250
25	Genome-wide Association Analysis of Psoriatic Arthritis and Cutaneous Psoriasis Reveals Differences in Their Genetic Architecture. American Journal of Human Genetics, 2015, 97, 816-836.	6.2	245
26	Histone Deacetylase Inhibitors Induce Growth Arrest and Differentiation in Uveal Melanoma. Clinical Cancer Research, 2012, 18, 408-416.	7.0	241
27	Broad defects in epidermal cornification in atopic dermatitis identified through genomic analysis. Journal of Allergy and Clinical Immunology, 2009, 124, 1235-1244.e58.	2.9	231
28	Association Mapping of Disease Loci, by Use of a Pooled DNA Genomic Screen. American Journal of Human Genetics, 1997, 61, 734-747.	6.2	228
29	The genetics of psoriasis, psoriatic arthritis and atopic dermatitis. Human Molecular Genetics, 2004, 13, 43R-55.	2.9	215
30	Deep sequencing of small RNAs from human skin reveals major alterations in the psoriasis miRNAome. Human Molecular Genetics, 2011, 20, 4025-4040.	2.9	213
31	Psoriasis genetics: breaking the barrier. Trends in Genetics, 2010, 26, 415-423.	6.7	203
32	Differential Contributions ofBRCA1andBRCA2to Early-Onset Breast Cancer. New England Journal of Medicine, 1997, 336, 1416-1422.	27.0	197
33	Driver Mutations in Uveal Melanoma. JAMA Ophthalmology, 2016, 134, 728.	2.5	192
34	Linkage of Tunisian autosomal recessive Duchenneâ€“like muscular dystrophy to the pericentromeric region of chromosome 13q. Nature Genetics, 1992, 2, 315-317.	21.4	186
35	Reversal of atopic dermatitis with narrow-band UVB phototherapy and biomarkers for therapeutic response. Journal of Allergy and Clinical Immunology, 2011, 128, 583-593.e4.	2.9	182
36	Fine Mapping Major Histocompatibility Complex Associations in Psoriasis and Its Clinical Subtypes. American Journal of Human Genetics, 2014, 95, 162-172.	6.2	182

#	Article	IF	Citations
37	Completing the map of human genetic variation. Nature, 2007, 447, 161-165.	27.8	178
38	A Gene for Congenital Generalized Lipodystrophy Maps to Human Chromosome 9q34. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 3390-3394.	3.6	167
39	Genome-wide meta-analysis identifies multiple novel associations and ethnic heterogeneity of psoriasis susceptibility. Nature Communications, 2015, 6, 6916.	12.8	154
40	Localization of the gene for familial partial lipodystrophy (Dunnigan variety) to chromosome 1q21â€“22. Nature Genetics, 1998, 18, 292-295.	21.4	151
41	Punctuated evolution of canonical genomic aberrations in uveal melanoma. Nature Communications, 2018, 9, 116.	12.8	144
42	CHD7 Gene Polymorphisms Are Associated with Susceptibility to Idiopathic Scoliosis. American Journal of Human Genetics, 2007, 80, 957-965.	6.2	142
43	Multiple Loci within the Major Histocompatibility Complex Confer Risk of Psoriasis. PLoS Genetics, 2009, 5, e1000606.	3.5	141
44	Multisystem dystrophy syndrome due to novel missense mutations in the amino-terminal head and alpha-helical rod domains of the lamin A/C gene. American Journal of Medicine, 2002, 112, 549-555.	1.5	138
45	A Subset of Methylated CpG Sites Differentiate Psoriatic from Normal Skin. Journal of Investigative Dermatology, 2012, 132, 583-592.	0.7	138
46	Asymmetric Lower-Limb Malformations in Individuals with Homeobox PITX1 Gene Mutation. American Journal of Human Genetics, 2008, 83, 616-622.	6.2	125
47	Localization of Susceptibility to Familial Idiopathic Scoliosis. Spine, 2000, 25, 2372-2380.	2.0	124
48	BAP1 deficiency causes loss of melanocytic cell identity in uveal melanoma. BMC Cancer, 2013, 13, 371.	2.6	123
49	Loss of Heterozygosity of Chromosome 3 Detected with Single Nucleotide Polymorphisms Is Superior to Monosomy 3 for Predicting Metastasis in Uveal Melanoma. Clinical Cancer Research, 2007, 13, 2923-2927.	7.0	122
50	THE GENETICS OF PSORIASIS AND AUTOIMMUNITY. Annual Review of Genomics and Human Genetics, 2005, 6, 93-122.	6.2	117
51	Molecular Cloning and Characterization of the HumanCLOCKGene: Expression in the Suprachiasmatic Nuclei. Genomics, 1999, 57, 189-200.	2.9	115
52	A Regulatory Role for 1-Acylglycerol-3-phosphate-O-acyltransferase 2 in Adipocyte Differentiation. Journal of Biological Chemistry, 2006, 281, 11082-11089.	3.4	112
53	New insights into the pathogenesis and genetics of psoriatic arthritis. Nature Clinical Practice Rheumatology, 2009, 5, 83-91.	3.2	112
54	Structure and Characterization of the Human Tissue Inhibitor of Metalloproteinases-2 Gene. Journal of Biological Chemistry, 1996, 271, 25498-25505.	3.4	103

#	Article	IF	Citations
55	Two novel point mutations in the long-range SHH enhancer in three families with triphalangeal thumb and preaxial polydactyly. American Journal of Medical Genetics, Part A, 2007, 143A, 27-32.	1.2	101
56	Genome-Wide Meta-Analysis of Psoriatic Arthritis Identifies Susceptibility Locus at REL. Journal of Investigative Dermatology, 2012, 132, 1133-1140.	0.7	99
57	Phenotypic Heterogeneity in Patients with Familial Partial Lipodystrophy (Dunnigan Variety) Related to the Site of Missense Mutations in Lamin A/C Gene1. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 59-65.	3.6	97
58	Epigenetic reprogramming and aberrant expression of PRAME are associated with increased metastatic risk in Class 1 and Class 2 uveal melanomas. Oncotarget, 2016, 7, 59209-59219.	1.8	94
59	Interferon Î±-Inducible Protein 27 (IFI27) is Upregulated in Psoriatic Skin and Certain Epithelial Cancers. Journal of Investigative Dermatology, 2004, 122, 717-721.	0.7	93
60	Meta-Analysis Confirms the LCE3C_LCE3B Deletion as a Risk Factor for Psoriasis in Several Ethnic Groups and Finds Interaction with HLA-Cw6. Journal of Investigative Dermatology, 2011, 131, 1105-1109.	0.7	89
61	High-Density Genetic Map of the BRCA1 Region of Chromosome 17q12-q21. Genomics, 1993, 17, 618-623.	2.9	87
62	Human Population Expansion and Microsatellite Variation. Molecular Biology and Evolution, 2000, 17, 757-767.	8.9	85
63	Activating Killer Cell Immunoglobulin-Like Receptor Gene KIR2DS1 Is Associated With Psoriatic Arthritis. Human Immunology, 2005, 66, 836-841.	2.4	84
64	A Genetic Risk Score Combining Ten Psoriasis Risk Loci Improves Disease Prediction. PLoS ONE, 2011, 6, e19454.	2.5	84
65	Rapid detection and sequencing of alleles in the 3â€² flanking region of the Interleukin-6 gene. Nucleic Acids Research, 1989, 17, 6855-6864.	14.5	83
66	CARD14-associated papulosquamous eruption: A spectrum including features of psoriasis and pityriasis rubra pilaris. Journal of the American Academy of Dermatology, 2018, 79, 487-494.	1.2	82
67	Direct genomic selection. Nature Methods, 2005, 2, 63-69.	19.0	81
68	Biology of advanced uveal melanoma and next steps for clinical therapeutics. Pigment Cell and Melanoma Research, 2015, 28, 135-147.	3.3	81
69	Novel immunoglobulin superfamily gene cluster, mapping to a region of human chromosome 17q25, linked to psoriasis susceptibility. Human Genetics, 2003, 112, 34-41.	3.8	74
70	Localization of a gene for familial recurrent arthritis. Arthritis and Rheumatism, 2000, 43, 2041-2045.	6.7	70
71	Genetics of psoriasis: the potential impact on new therapies. Journal of the American Academy of Dermatology, 2003, 49, 51-56.	1.2	70
72	Risk Factors for Diabetes in Familial Partial Lipodystrophy, Dunnigan Variety. Diabetes Care, 2003, 26, 1350-1355.	8.6	68

#	Article	IF	Citations
73	AKT Inhibitors Promote Cell Death in Cervical Cancer through Disruption of mTOR Signaling and Glucose Uptake. PLoS ONE, 2014, 9, e92948.	2.5	68
74	Further Genetic Evidence for Three Psoriasis-Risk Genes: ADAM33, CDKAL1, and PTPN22. Journal of Investigative Dermatology, 2009, 129, 629-634.	0.7	67
75	Genetic Linkage Localizes an Adolescent Idiopathic Scoliosis and Pectus Excavatum Gene to Chromosome 18 q. Spine, 2009, 34, E94-E100.	2.0	66
76	Psoriasis Patients Are Enriched for Genetic Variants That Protect against HIV-1 Disease. PLoS Genetics, 2012, 8, e1002514.	3.5	66
77	Carriers of Rare Missense Variants in IFIH1 Are Protected from Psoriasis. Journal of Investigative Dermatology, 2010, 130, 2768-2772.	0.7	65
78	SNTG1, the gene encoding ?1-syntrophin: a candidate gene for idiopathic scoliosis. Human Genetics, 2004, 115, 81-89.	3.8	64
79	Association of Cardiovascular and Metabolic Disease Genes with Psoriasis. Journal of Investigative Dermatology, 2013, 133, 836-839.	0.7	62
80	Psoriasis mutations disrupt CARD14 autoinhibition promoting BCL10-MALT1-dependent NF-Î¹B activation. Biochemical Journal, 2016, 473, 1759-1768.	3.7	62
81	Localization of PSORS1 to a haplotype block harboring HLA-C and distinct from corneodesmosin and HCR. Human Genetics, 2005, 118, 466-476.	3.8	61
82	Disruption of Sodium Bicarbonate Transporter SLC4A10 in a Patient With Complex Partial Epilepsy and Mental Retardation. Archives of Neurology, 2008, 65, 550.	4.5	61
83	A transmission disequilibrium test for general pedigrees that is robust to the presence of random genotyping errors and any number of untyped parents. European Journal of Human Genetics, 2004, 12, 752-761.	2.8	57
84	A Gene for Congenital Generalized Lipodystrophy Maps to Human Chromosome 9q34. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 3390-3394.	3.6	56
85	Microsatellite Polymorphism Linkage Map of Human Chromosome 13q. Genomics, 1993, 15, 376-386.	2.9	55
86	The CEPH Consortium Linkage Map of Human Chromosome 13. Genomics, 1993, 16, 486-496.	2.9	55
87	A linkage map of human chromosome 15 with an average resolution of 2 cM and containing 55 polymorphic microsatellites. Human Molecular Genetics, 1993, 2, 2019-2030.	2.9	54
88	CARD14 Expression in Dermal Endothelial Cells in Psoriasis. PLoS ONE, 2014, 9, e111255.	2.5	52
89	Isolation of Two Novel WNT Genes, WNT14 and WNT15, One of Which (WNT15) Is Closely Linked to WNT3 on Human Chromosome 17q21. Genomics, 1997, 46, 450-458.	2.9	50
90	Characterization ofEZH1,a Human Homolog ofDrosophila Enhancer of zestenearBRCA1. Genomics, 1996, 37, 161-171.	2.9	49

#	Article	IF	Citations
91	HOXD10 M319K mutation in a family with isolated congenital vertical talus. Journal of Orthopaedic Research, 2006, 24, 448-453.	2.3	48
92	Peptidoglycan recognition proteins Pglyrp3 and Pglyrp4 are encoded from the epidermal differentiation complex and are candidate genes for the Psors4 locus on chromosome 1q21. Human Genetics, 2006, 119, 113-125.	3.8	45
93	The study of variation in the human genome. Genomics, 1991, 11, 491-498.	2.9	43
94	Noncanonical microRNAs and endogenous siRNAs in normal and psoriatic human skin. Human Molecular Genetics, 2013, 22, 737-748.	2.9	43
95	The human cationic amino acid transporter (ATRC1): Physical and genetic mapping to 13q12â€“q14. Genomics, 1992, 12, 430-434.	2.9	42
96	Guilt by association. Nature, 2007, 447, 645-646.	27.8	40
97	Conservation of function and primary structure in the BRCA1-associated RING domain (BARD1) protein. Oncogene, 1998, 17, 2143-2148.	5.9	36
98	Management of resectable colorectal lung metastases. Clinical and Experimental Metastasis, 2016, 33, 285-296.	3.3	36
99	Deletion of the activating <scp>NKG</scp>2C receptor and a functional polymorphism in its ligand <scp>HLA</scp>â€Ł in psoriasis susceptibility. Experimental Dermatology, 2013, 22, 679-681.	2.9	31
100	Psoriasis Bench to Bedside. Archives of Dermatology, 2009, 145, 462-4.	1.4	29
101	The Tetratricopeptide Repeat Domain 7 Gene is Mutated in Flaky Skin Mice: A Model for Psoriasis, Autoimmunity, and Anemia. Experimental Biology and Medicine, 2005, 230, 659-667.	2.4	26
102	A contiguous linkage map of chromosome 13q with 39 distinct loci separated on average by 5.1 centimorgans. Genomics, 1991, 11, 517-529.	2.9	25
103	Fine mapping of eight psoriasis susceptibility loci. European Journal of Human Genetics, 2015, 23, 844-853.	2.8	25
104	Genomic structure, mapping, and expression analysis of the mammalian Lunatic, Manic, and Radical fringe genes. Mammalian Genome, 1999, 10, 535-541.	2.2	24
105	Familial and Somatic BAP1 Mutations Inactivate ASXL1/2-Mediated Allosteric Regulation of BAP1 Deubiquitinase by Targeting Multiple Independent Domains. Cancer Research, 2018, 78, 1200-1213.	0.9	24
106	Protective Effect of Human Endogenous Retrovirus K dUTPase Variants on Psoriasis Susceptibility. Journal of Investigative Dermatology, 2012, 132, 1833-1840.	0.7	22
107	Direct selection of expressed sequences within a 1-Mb region flanking BRCA1 on human chromosome 17q21. Genomics, 1995, 25, 248-255.	2.9	21
108	DNA copy number changes as diagnostic tools for lung cancer. Thorax, 2014, 69, 496-497.	5.6	21

#	Article	IF	Citations
109	Predicting genotypes at loci for autosomal recessive disorders using linked genetic markers: application to Wilson's disease. Human Genetics, 1988, 79, 109-117.	3.8	20
110	Variable hand and foot abnormalities in family with congenital vertical talus and CDMP-1 gene mutation. Journal of Orthopaedic Research, 2005, 23, 1490-1494.	2.3	19
111	Absence of HOXD10 Mutations in Idiopathic Clubfoot and Sporadic Vertical Talus. Clinical Orthopaedics and Related Research, 2007, 462, 27-31.	1.5	19
112	Primary Ciliary Dyskinesia-Causing Mutations in Amish and Mennonite Communities. Journal of Pediatrics, 2013, 163, 383-387.	1.8	19
113	Loss of heterozygosity for chromosome 22 DNA sequences in human meningioma. Cancer Genetics and Cytogenetics, 1991, 53, 271-277.	1.0	18
114	Molecular cloning of BRCA1: a gene for early onset familial breast and ovarian cancer. Breast Cancer Research and Treatment, 1993, 28, 121-135.	2.5	18
115	In vitro transformation of cell lines from human salivary gland tumors. , 1999, 81, 793-798.		18
116	Psoriasis Genetics: The Way Forward. Journal of Investigative Dermatology, 2004, 122, xv-xvii.	0.7	18
117	Understanding the Pathogenesis of Psoriasis, Psoriatic Arthritis, and Autoimmunity via a Fusion of Molecular Genetics and Immunology. Immunologic Research, 2005, 32, 045-056.	2.9	18
118	The tumor genetics of acral melanoma: What should a dermatologist know?. JAAD International, 2020, 1, 135-147.	2.2	18
119	Polymorphism and mapping of the IGF1 gene, and absence of association with stature among African Pygmies. Human Genetics, 1990, 85, 349-54.	3.8	17
120	Evolution of Modern Humans: Evidence From Nuclear DNA Polymorphisms. , 1993, , 69-83.		17
121	Evidence for an additional locus for split hand/foot malformation in chromosome region 8q21.11â€“q22.3. American Journal of Medical Genetics, Part A, 2006, 140A, 1744-1748.	1.2	17
122	Contribution of genetic factors for melanoma susceptibility in sporadic US melanoma patients. Experimental Dermatology, 2009, 18, 485-487.	2.9	17
123	The haemoglobin H disease mental retardation syndrome: molecular studies on the South African case. British Journal of Haematology, 1984, 56, 69-78.	2.5	16
124	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. Genomics, 1990, 7, 110-114.	2.9	16
125	Infrequency of BRCA2 alterations in head and neck squamous cell carcinoma. Oncogene, 1997, 14, 2189-2193.	5.9	16
126	Investigation of the Chromosome 17q25 PSORS2 Locus in Atopic Dermatitis. Journal of Investigative Dermatology, 2006, 126, 603-606.	0.7	16

#	Article	IF	Citations
127	Association of Melanocortin-1 Receptor Variants with Pigmentary Traits in Humans: AÂPooled Analysis from the M-Skip Project. Journal of Investigative Dermatology, 2016, 136, 1914-1917.	0.7	16
128	Integrative Copy Number Analysis of Uveal Melanoma Reveals Novel Candidate Genes Involved in Tumorigenesis Including a Tumor Suppressor Role for PHF10/BAF45a. Clinical Cancer Research, 2019, 25, 5156-5166.	7.0	16
129	CARD14E138A signalling in keratinocytes induces TNF-dependent skin and systemic inflammation. ELife, 2020, 9, .	6.0	16
130	A YAC-, P1-, and cosmid-based physical Map of the BRCA1 region on chromosome 17q21. Genomics, 1995, 25, 264-273.	2.9	15
131	CARD14â€associated papulosquamous eruption (CAPE) in pediatric patients: Three additional cases and review of the literature. Pediatric Dermatology, 2021, 38, 1237-1242.	0.9	15
132	Influence of CrohnÊ¼s Disease Risk Alleles and Smoking on Disease Location. Diseases of the Colon and Rectum, 2011, 54, 1020-1025.	1.3	15
133	The Molecular Revolution in Cutaneous Biology: The Era of Genome-Wide Association Studies and Statistical, BigÂData, and Computational Topics. Journal of Investigative Dermatology, 2017, 137, e113-e118.	0.7	14
134	Molecular profiling of colorectal pulmonary metastases and primary tumours: implications for targeted treatment. Oncotarget, 2017, 8, 64999-65008.	1.8	14
135	Primary Ciliary Dyskinesia in Amish Communities. Journal of Pediatrics, 2010, 156, 1023-1025.	1.8	13
136	Targeting Skin: Vitiligo and Autoimmunity. Journal of Investigative Dermatology, 2012, 132, 13-15.	0.7	13
137	Gain of function p.E138A alteration in Card14 leads to psoriasiform skin inflammation and implicates genetic modifiers in disease severity. Experimental and Molecular Pathology, 2019, 110, 104286.	2.1	13
138	Genetic Locus for Psoriasis Identified. Annals of Medicine, 1995, 27, 183-186.	3.8	12
139	JAK3 Maps to Human Chromosome 19p12 within a Cluster of Proto-oncogenes and Transcription Factors. Genomics, 1997, 43, 109-111.	2.9	12
140	The NGF and kallikrein genes of mouse, the African rat Mastomys natalensis and man: their distribution and mode of expression in the salivary gland. Molecular Brain Research, 1988, 3, 165-172.	2.3	11
141	Global expression and CpG methylation analysis of primary endothelial cells before and after TNFa stimulation reveals gene modules enriched in inflammatory and infectious diseases and associated DMRs. PLoS ONE, 2020, 15, e0230884.	2.5	11
142	Breast Cancer Genes. Breast Journal, 1997, 3, 1-6.	1.0	9
143	Genome-Wide Association Studies and Infectious Disease. Critical Reviews in Immunology, 2010, 30, 305-309.	0.5	9
144	Runx Transcription Factors Repress Human and Murine c-Myc Expression in a DNA-Binding and C-Terminally Dependent Manner. PLoS ONE, 2013, 8, e69083.	2.5	9

#	Article	IF	Citations
145	Canonical and Interior Circular RNAs Function as Competing Endogenous RNAs in Psoriatic Skin. International Journal of Molecular Sciences, 2021, 22, 5182.	4.1	8
146	Zeroing in on tolerance. Nature Medicine, 2001, 7, 279-281.	30.7	7
147	Additional evidence of a locus for complex febrile and afebrile seizures on chromosome 12q22-23.3. Neurogenetics, 2007, 8, 61-63.	1.4	7
148	A somatic cell hybrid map of human chromosome 13. Genomics, 1993, 18, 486-495.	2.9	6
149	Detection of a polymorphism within the pepsinogen C gene with PCR: Construction of a linkage map around PGC from 6p11â€“6p21.3. Genomics, 1992, 14, 398-402.	2.9	5
150	An SspI RFLP at the D13S25 locus identified by the anonymous single copy probe H2â€“42. Nucleic Acids Research, 1990, 18, 7194-7194.	14.5	4
151	Dinucleotide repeat polymorphism at the D1S167 locus. Human Molecular Genetics, 1992, 1, 138-138.	2.9	4
152	Use of pharmacogenomics in psoriasis. Clinical Investigation, 2011, 1, 399-411.	0.0	4
153	Circulating tumor DNA is readily detectable among Ghanaian breast cancer patients supporting non-invasive cancer genomic studies in Africa. Npj Precision Oncology, 2021, 5, 83.	5.4	4
154	The anonymous probe pR1â€“4 which identifies the locus D13S59 detects a BanII RFLP. Nucleic Acids Research, 1989, 17, 8396-8396.	14.5	3
155	The Gene for Pancreatic Polypeptide (PPY) and the Anonymous Marker D17S78 Are within 45 kb of Each Other on Chromosome 17q21. Genomics, 1994, 21, 458-460.	2.9	3
156	A Highly Informative Polymorphism of the Pepsinogen C Gene Detected by Polymerase Chain Reaction. Advances in Experimental Medicine and Biology, 1991, 306, 95-99.	1.6	3
157	The CA repeat marker D17S791 islocated within 40 kb of the WNT3 gene on chromosome 17q. Genomics, 1993, 18, 728-729.	2.9	2
158	W1122 Serology Testing, Genotype, and Other Risk Factors for Repeat Ileocolic Resection in Crohn's Disease: A Time-to-Event Analysis. Gastroenterology, 2008, 134, A-637-A-638.	1.3	2
159	Clinicopathological features and associations in a series of South African acral melanomas. Pigment Cell and Melanoma Research, 2021, 34, 1120-1122.	3.3	2
160	The anonymous probe pF5A identifying the locus D13S61 detects RFLPs with XmnI and BanII. Nucleic Acids Research, 1989, 17, 8397-8397.	14.5	1
161	The Î± chain of human propionyl CoA carboxylase (PCCA) (mapped to chromosome 13) detects an RFLP with XmnI. Nucleic Acids Research, 1989, 17, 8400-8400.	14.5	1
162	Dinucleotide repeat polymorphism at the D6S223 locus. Human Molecular Genetics, 1992, 1, 66-66.	2.9	1

#	Article	IF	Citations
163	Kinetic Characterization of ASXL1/2-Mediated Allosteric Regulation of the BAP1 Deubiquitinase. Molecular Cancer Research, 2021, 19, 1099-1112.	3.4	1
164	Hereditary Breast Cancer Genes. , 1999, , 199-224.		1
165	The anonymous DNA probe p7-26 identifying the locus [D7S17] reveals an XmnI polymorphism. Nucleic Acids Research, 1989, 17, 1787-1787.	14.5	0
166	Association mapping with a pooled DNA genomic screen of the non-HLA genes in IDDM, RA and MS. Human Immunology, 1996, 47, 152.	2.4	0
167	Markers and Methods for Reconstructing Modern Human History. DNA Sequence, 1998, 8, 329-342.	0.7	0
168	A Breath of Fresh Air: Opening up the Lung Cancer Genome. Cancer Research, 2019, 79, 4808-4810.	0.9	0
169	Abstract 81: Studying Ghanian Cancer Genomes Using Cell-free DNA. , 2021, , .		0
170	The Immunogenetics of Inflammatory Skin Disease. , 2004, , 55-73.		0
171	CARD14-Mediated Psoriasis and Pityriasis Rubra Piliaris (PRP). , 2018, , 1-4.		0
172	CARD14-Mediated Psoriasis and Pityriasis Rubra Piliaris (PRP). , 2020, , 92-95.		0
173	Hereditary Breast Cancer Genes. , 0, , 199-224.		0