

Julia A Newton-Bishop

List of Publications by Year in descending order

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Version: 2024-02-01

191
papers

11,171
citations

28242

55
h-index

34964

98
g-index

202
all docs

202
docs citations

202
times ranked

12275
citing authors

#	ARTICLE	IF	CITATIONS
1	ROR2 promotes epithelial-mesenchymal transition by hyperactivating ERK in melanoma. <i>Journal of Cell Communication and Signaling</i> , 2023, 17, 75-88.	1.8	3
2	Ulcerated melanoma: Systems biology evidence of inflammatory imbalance towards pro-tumorigenicity. <i>Pigment Cell and Melanoma Research</i> , 2022, 35, 252-267.	1.5	4
3	Genes regulated by DNA methylation are involved in distinct phenotypes during melanoma progression and are prognostic factors for patients. <i>Molecular Oncology</i> , 2022, 16, 1913-1930.	2.1	1
4	Cross-cohort gut microbiome associations with immune checkpoint inhibitor response in advanced melanoma. <i>Nature Medicine</i> , 2022, 28, 535-544.	15.2	158
5	Defining novel causal SNPs and linked phenotypes at melanoma-associated loci. <i>Human Molecular Genetics</i> , 2022, 31, 2845-2856.	1.4	3
6	Multiple Primary Melanoma Incidence Trends Over Five Decades: A Nationwide Population-Based Study. <i>Journal of the National Cancer Institute</i> , 2021, 113, 318-328.	3.0	19
7	MX2 mediates establishment of interferon response profile, regulates XAF1, and can sensitize melanoma cells to targeted therapy. <i>Cancer Medicine</i> , 2021, 10, 2840-2854.	1.3	6
8	Birth cohort-specific trends of sun-related behaviors among individuals from an international consortium of melanoma-prone families. <i>BMC Public Health</i> , 2021, 21, 692.	1.2	4
9	Transcriptional signatures underlying dynamic phenotypic switching and novel disease biomarkers in a linear cellular model of melanoma progression. <i>Neoplasia</i> , 2021, 23, 439-455.	2.3	5
10	Inherited duplications of PPP2R3B predispose to nevi and melanoma via a C21orf91-driven proliferative phenotype. <i>Genetics in Medicine</i> , 2021, 23, 1636-1647.	1.1	5
11	Road to Metastasis: The TWEAK Pathway as a Discriminant between Metastasizing and Non-Metastasizing Thick Melanomas. <i>International Journal of Molecular Sciences</i> , 2021, 22, 10568.	1.8	0
12	Cooperation between melanoma cell states promotes metastasis through heterotypic cluster formation. <i>Developmental Cell</i> , 2021, 56, 2808-2825.e10.	3.1	37
13	Tumour gene expression signature in primary melanoma predicts long-term outcomes. <i>Nature Communications</i> , 2021, 12, 1137.	5.8	33
14	ROR2 has a protective role in melanoma by inhibiting Akt activity, cell-cycle progression, and proliferation. <i>Journal of Biomedical Science</i> , 2021, 28, 76.	2.6	8
15	Environmental Exposures Such as Smoking and Low Vitamin D Are Predictive of Poor Outcome in Cutaneous Melanoma rather than Other Deprivation Measures. <i>Journal of Investigative Dermatology</i> , 2020, 140, 327-337.e2.	0.3	14
16	MX 2 is a novel regulator of cell cycle in melanoma cells. <i>Pigment Cell and Melanoma Research</i> , 2020, 33, 446-457.	1.5	11
17	Study of the Female Sex Survival Advantage in Melanoma—A Focus on X-Linked Epigenetic Regulators and Immune Responses in Two Cohorts. <i>Cancers</i> , 2020, 12, 2082.	1.7	16
18	Gene co-expression and histone modification signatures are associated with melanoma progression, epithelial-to-mesenchymal transition, and metastasis. <i>Clinical Epigenetics</i> , 2020, 12, 127.	1.8	9

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19	MC1R variants and cutaneous melanoma risk according to histological type, body site, and Breslow thickness: a pooled analysis from the M-SKIP project. <i>Melanoma Research</i> , 2020, 30, 500-510.	0.6	6
20	Factors Affecting Sentinel Node Metastasis in Thin (T1) Cutaneous Melanomas: Development and External Validation of a Predictive Nomogram. <i>Journal of Clinical Oncology</i> , 2020, 38, 1591-1601.	0.8	50
21	Genome-wide association meta-analyses combining multiple risk phenotypes provide insights into the genetic architecture of cutaneous melanoma susceptibility. <i>Nature Genetics</i> , 2020, 52, 494-504.	9.4	138
22	Gain-of-Function Genetic Alterations of G9a Drive Oncogenesis. <i>Cancer Discovery</i> , 2020, 10, 980-997.	7.7	44
23	Estimating CDKN2A mutation carrier probability among global familial melanoma cases using GenoMELPREDICT. <i>Journal of the American Academy of Dermatology</i> , 2019, 81, 386-394.	0.6	17
24	Transcriptomic Analysis Reveals Prognostic Molecular Signatures of Stage I Melanoma. <i>Clinical Cancer Research</i> , 2019, 25, 7424-7435.	3.2	27
25	High-Resolution Copy Number Patterns From Clinically Relevant FFPE Material. <i>Scientific Reports</i> , 2019, 9, 8908.	1.6	6
26	MC1R variants in childhood and adolescent melanoma: a retrospective pooled analysis of a multicentre cohort. <i>The Lancet Child and Adolescent Health</i> , 2019, 3, 332-342.	2.7	16
27	Genetic and Environmental Determinants of Immune Response to Cutaneous Melanoma. <i>Cancer Research</i> , 2019, 79, 2684-2696.	0.4	45
28	Vitamin D β 2R Signaling Inhibits Wnt/ β 2-Catenin-Mediated Melanoma Progression and Promotes Antitumor Immunity. <i>Cancer Research</i> , 2019, 79, 5986-5998.	0.4	65
29	Application of Circulating Cell-Free Tumor DNA Profiles for Therapeutic Monitoring and Outcome Prediction in Genetically Heterogeneous Metastatic Melanoma. <i>JCO Precision Oncology</i> , 2019, 3, 1-10.	1.5	25
30	Association of the <i>POT1</i> Germline Missense Variant p.I78T With Familial Melanoma. <i>JAMA Dermatology</i> , 2019, 155, 604.	2.0	34
31	ctDNA as a noninvasive monitoring tool in metastatic melanoma. <i>Journal of Clinical Oncology</i> , 2019, 37, 9548-9548.	0.8	0
32	Comparative genomics reveals that loss of lunatic fringe (<i>LFNG</i>) promotes melanoma metastasis. <i>Molecular Oncology</i> , 2018, 12, 239-255.	2.1	20
33	Novel pleiotropic risk loci for melanoma and nevus density implicate multiple biological pathways. <i>Nature Communications</i> , 2018, 9, 4774.	5.8	87
34	Comprehensive Study of the Clinical Phenotype of Germline <i>BAP1</i> Variant-Carrying Families Worldwide. <i>Journal of the National Cancer Institute</i> , 2018, 110, 1328-1341.	3.0	164
35	Somatic cancer genetics in the UK: real-world data from phase I of the Cancer Research UK Stratified Medicine Programme. <i>ESMO Open</i> , 2018, 3, e000408.	2.0	4
36	25-hydroxyvitamin D serum levels in patients with high risk resected melanoma treated in an adjuvant bevacizumab trial. <i>British Journal of Cancer</i> , 2018, 119, 793-800.	2.9	11

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37	Amelanotic melanoma. <i>BMJ: British Medical Journal</i> , 2018, 360, k826.	2.4	9
38	MC1R variants as melanoma risk factors independent of at-risk phenotypic characteristics: a pooled analysis from the M-SKIP project. <i>Cancer Management and Research</i> , 2018, Volume 10, 1143-1154.	0.9	57
39	Assessing the Incremental Contribution of Common Genomic Variants to Melanoma Risk Prediction in Two Population-Based Studies. <i>Journal of Investigative Dermatology</i> , 2018, 138, 2617-2624.	0.3	52
40	Î²-Catenin-mediated immune evasion pathway frequently operates in primary cutaneous melanomas. <i>Journal of Clinical Investigation</i> , 2018, 128, 2048-2063.	3.9	71
41	Abstract 5204:MYC expression and smoking as drivers of failure of immune response to melanoma. , 2018, , .		0
42	Abstract 5205: Primary melanoma expression of the vitamin D receptor (VDR) is protective for melanoma survival and is associated with increased tumor immune response, decreased Wnt/B-catenin signaling and tumor proliferation. , 2018, , .		0
43	Abstract 228: Association analysis across different populations identifies 26 new cutaneous melanoma risk loci. , 2018, , .		0
44	Abstract 234: Understanding melanoma susceptibility through GWAS of risk phenotypes. , 2018, , .		0
45	Abstract 5372: The mutational landscape of primary cutaneous melanoma. , 2018, , .		0
46	A population-based analysis of germline <i>BAP1</i> mutations in melanoma. <i>Human Molecular Genetics</i> , 2017, 26, ddw403.	1.4	31
47	Functional characterization of a multi-cancer risk locus on chr5p15.33 reveals regulation of TERT by ZNF148. <i>Nature Communications</i> , 2017, 8, 15034.	5.8	40
48	Germline <i>CDKN2A</i> / <i>P16INK4A</i> mutations contribute to genetic determinism of sarcoma. <i>Journal of Medical Genetics</i> , 2017, 54, 607-612.	1.5	19
49	Rare Variant, Gene-Based Association Study of Hereditary Melanoma Using Whole-Exome Sequencing. <i>Journal of the National Cancer Institute</i> , 2017, 109, .	3.0	32
50	Tests to assist in the staging of cutaneous melanoma: a generic protocol. <i>The Cochrane Library</i> , 2017, , .	1.5	1
51	Germline Variation at <i>CDKN2A</i> and Associations with Nevus Phenotypes among Members of Melanoma Families. <i>Journal of Investigative Dermatology</i> , 2017, 137, 2606-2612.	0.3	18
52	A common intronic variant of <i>PARP1</i> confers melanoma risk and mediates melanocyte growth via regulation of <i>MITF</i> . <i>Nature Genetics</i> , 2017, 49, 1326-1335.	9.4	51
53	Identification of a gene signature for discriminating metastatic from primary melanoma using a molecular interaction network approach. <i>Scientific Reports</i> , 2017, 7, 17314.	1.6	32
54	Which symptoms are linked to a delayed presentation among melanoma patients? A retrospective study. <i>BMC Cancer</i> , 2017, 17, 5.	1.1	5

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55	Genomic expression differences between cutaneous cells from red hair color individuals and black hair color individuals based on bioinformatic analysis. <i>Oncotarget</i> , 2017, 8, 11589-11599.	0.8	5
56	Abstract 4006: Immune cell profiles and β -catenin signaling in melanoma. , 2017, , .		1
57	Abstract 3717: New therapies for the treatment of BRAF/NRAS wild type melanoma. , 2017, , .		0
58	Abstract 1741: Whole-transcriptome characterisation of NRAS and BRAF mutated primary melanomas associated with immune cell infiltration signatures and differential survival benefit. , 2017, , .		0
59	Psychosocial, clinical and demographic features related to worry in patients with melanoma. <i>Melanoma Research</i> , 2016, 26, 497-504.	0.6	8
60	Diagnosis and treatment of melanoma. European consensus-based interdisciplinary guideline "Update 2016. <i>European Journal of Cancer</i> , 2016, 63, 201-217.	1.3	330
61	Vitamin D, vitamin A, the primary melanoma transcriptome and survival. <i>British Journal of Dermatology</i> , 2016, 175, 30-34.	1.4	11
62	Histopathology of melanocytic lesions in a family with an inherited <i>BAP1</i> mutation. <i>Journal of Cutaneous Pathology</i> , 2016, 43, 287-289.	0.7	10
63	Association of Melanocortin-1 Receptor Variants with Pigmentary Traits in Humans: A Pooled Analysis from the M-Skip Project. <i>Journal of Investigative Dermatology</i> , 2016, 136, 1914-1917.	0.3	16
64	The <i>CDKN2A/p16</i> ^{INK4a} UTR sequence and translational regulation: impact of novel variants predisposing to melanoma. <i>Pigment Cell and Melanoma Research</i> , 2016, 29, 210-221.	1.5	9
65	Consensus of Melanoma Gene Expression Subtypes Converges on Biological Entities. <i>Journal of Investigative Dermatology</i> , 2016, 136, 2502-2505.	0.3	23
66	Wide versus narrow excision margins for high-risk, primary cutaneous melanomas: long-term follow-up of survival in a randomised trial. <i>Lancet Oncology</i> , The, 2016, 17, 184-192.	5.1	116
67	Phenotypic and Histopathological Tumor Characteristics According to CDKN2A Mutation Status among Affected Members of Melanoma Families. <i>Journal of Investigative Dermatology</i> , 2016, 136, 1066-1069.	0.3	13
68	Loss-of-Function Mutations in the Cell-Cycle Control Gene <i>CDKN2A</i> Impact on Glucose Homeostasis in Humans. <i>Diabetes</i> , 2016, 65, 527-533.	0.3	38
69	Germline TERT promoter mutations are rare in familial melanoma. <i>Familial Cancer</i> , 2016, 15, 139-144.	0.9	51
70	<i>MC1R</i> variants increased the risk of sporadic cutaneous melanoma in darker pigmented Caucasians: A pooled analysis from the M-Skip project. <i>International Journal of Cancer</i> , 2015, 136, 618-631.	2.3	92
71	25-Hydroxyvitamin D ₂ /D ₃ levels and factors associated with systemic inflammation and melanoma survival in the Leeds Melanoma Cohort. <i>International Journal of Cancer</i> , 2015, 136, 2890-2899.	2.3	61
72	Authors' reply to: High naevus counts confer a favourable prognosis in patients with melanoma. <i>International Journal of Cancer</i> , 2015, 137, 3008-3009.	2.3	0

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73	High nevus counts confer a favorable prognosis in melanoma patients. <i>International Journal of Cancer</i> , 2015, 137, 1691-1698.	2.3	37
74	Development and Validation of a Melanoma Risk Score Based on Pooled Data from 16 Case-Control Studies. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015, 24, 817-824.	1.1	25
75	Histologic features of melanoma associated with CDKN2A genotype. <i>Journal of the American Academy of Dermatology</i> , 2015, 72, 496-507.e7.	0.6	19
76	MGMT promoter methylation is associated with temozolomide response and prolonged progression-free survival in disseminated cutaneous melanoma. <i>International Journal of Cancer</i> , 2015, 136, 2844-2853.	2.3	45
77	The clinicopathological and gene expression patterns associated with ulceration of primary melanoma. <i>Pigment Cell and Melanoma Research</i> , 2015, 28, 94-104.	1.5	26
78	Nonsense Mutations in the Shelterin Complex Genes ACD and TERF2IP in Familial Melanoma. <i>Journal of the National Cancer Institute</i> , 2015, 107, .	3.0	134
79	Genome-wide meta-analysis identifies five new susceptibility loci for cutaneous malignant melanoma. <i>Nature Genetics</i> , 2015, 47, 987-995.	9.4	218
80	Fine mapping of genetic susceptibility loci for melanoma reveals a mixture of single variant and multiple variant regions. <i>International Journal of Cancer</i> , 2015, 136, 1351-1360.	2.3	30
81	Regressing Eruptive Disseminated Spitz Nevi. <i>Pediatric Dermatology</i> , 2015, 32, e181-3.	0.5	9
82	Melanoma: summary of NICE guidance. <i>BMJ, The</i> , 2015, 351, h3708.	3.0	23
83	Prognostic Significance of Promoter Hypermethylation and Diminished Gene Expression of SYNPO2 in Melanoma. <i>Journal of Investigative Dermatology</i> , 2015, 135, 2328-2331.	0.3	13
84	Independent replication of a melanoma subtype gene signature and evaluation of its prognostic value and biological correlates in a population cohort. <i>Oncotarget</i> , 2015, 6, 11683-11693.	0.8	44
85	Identification of a melanoma susceptibility locus and somatic mutation in <i>TET2</i> . <i>Carcinogenesis</i> , 2014, 35, 2097-2101.	1.3	41
86	Somatic BRAF and NRAS Mutations in Familial Melanomas with Known Germline CDKN2A Status: A GenoMEL Study. <i>Journal of Investigative Dermatology</i> , 2014, 134, 287-290.	0.3	18
87	Inherited variation in the PARP1 gene and survival from melanoma. <i>International Journal of Cancer</i> , 2014, 135, 1625-1633.	2.3	24
88	An inherited variant in the gene coding for vitamin D-binding protein and survival from cutaneous melanoma: a BIOGENO-MEL study. <i>Pigment Cell and Melanoma Research</i> , 2014, 27, 234-243.	1.5	25
89	Prevalence and predictors of germline CDKN2A mutations for melanoma cases from Australia, Spain and the United Kingdom. <i>Hereditary Cancer in Clinical Practice</i> , 2014, 12, 20.	0.6	45
90	The Effect on Melanoma Risk of Genes Previously Associated With Telomere Length. <i>Journal of the National Cancer Institute</i> , 2014, 106, .	3.0	109

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91	POT1 loss-of-function variants predispose to familial melanoma. <i>Nature Genetics</i> , 2014, 46, 478-481.	9.4	319
92	Primary Melanoma Tumors from CDKN2A Mutation Carriers Do Not Belong to a Distinct Molecular Subclass. <i>Journal of Investigative Dermatology</i> , 2014, 134, 3000-3003.	0.3	8
93	Targeting human apurinic/aprimidinic endonuclease 1 (APE1) in phosphatase and tensin homolog (PTEN) deficient melanoma cells for personalized therapy. <i>Oncotarget</i> , 2014, 5, 3273-3286.	0.8	47
94	Abstract 20: POT1 mutations predispose to familial melanoma. , 2014, , .		0
95	MC1R genotype as a predictor of early-onset melanoma, compared with self-reported and physician-measured traditional risk factors: an Australian case-control-family study. <i>BMC Cancer</i> , 2013, 13, 406.	1.1	30
96	Do vitamin A serum levels moderate outcome or the protective effect of vitamin D on outcome from malignant melanoma?. <i>Clinical Nutrition</i> , 2013, 32, 1012-1016.	2.3	7
97	A variant in FTO shows association with melanoma risk not due to BMI. <i>Nature Genetics</i> , 2013, 45, 428-432.	9.4	111
98	Association between putative functional variants in the <i>PSMB9</i> gene and risk of melanoma – reanalysis of published melanoma genome-wide association studies. <i>Pigment Cell and Melanoma Research</i> , 2013, 26, 392-401.	1.5	5
99	Melanoma prone families with <i>CDK4</i> germline mutation: phenotypic profile and associations with <i>MC1R</i> variants. <i>Journal of Medical Genetics</i> , 2013, 50, 264-270.	1.5	112
100	Association between functional polymorphisms in genes involved in the MAPK signaling pathways and cutaneous melanoma risk. <i>Carcinogenesis</i> , 2013, 34, 885-892.	1.3	10
101	Erythema Nodosum-like Panniculitis in Patients With Melanoma Treated With Vemurafenib. <i>Journal of Clinical Oncology</i> , 2013, 31, e320-e321.	0.8	29
102	Evaluation of <i>PAX3</i> genetic variants and nevus number. <i>Pigment Cell and Melanoma Research</i> , 2013, 26, 666-676.	1.5	7
103	Response to P. Autier and M. Boniol regarding our article-Relationship between sunbed use and melanoma risk in a large case-control study in the United Kingdom. <i>International Journal of Cancer</i> , 2013, 132, 1960-1961.	2.3	1
104	Vitamin D and melanoma. <i>Dermato-Endocrinology</i> , 2013, 5, 121-129.	1.9	28
105	Skin Examination Behavior. <i>Archives of Dermatology</i> , 2012, 148, 1142.	1.7	36
106	Objective assessment of blood and lymphatic vessel invasion and association with macrophage infiltration in cutaneous melanoma. <i>Modern Pathology</i> , 2012, 25, 493-504.	2.9	105
107	Clinicopathologic Features of V600E and V600K Melanoma – Letter. <i>Clinical Cancer Research</i> , 2012, 18, 6792-6792.	3.2	14
108	Authors' reply to Bayley and Cave. <i>BMJ</i> , The, 2012, 345, e5417-e5417.	3.0	0

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109	Nick Hayward. Pigment Cell and Melanoma Research, 2012, 25, 116-116.	1.5	0
110	Identification of differentially expressed genes in matched formalin-fixed paraffin-embedded primary and metastatic melanoma tumor pairs. Pigment Cell and Melanoma Research, 2012, 25, 284-286.	1.5	5
111	Inherited variants in the <i>MC1R</i> gene and survival from cutaneous melanoma: a BioGenoMEL study. Pigment Cell and Melanoma Research, 2012, 25, 384-394.	1.5	61
112	Perceptions of genetic research and testing among members of families with an increased risk of malignant melanoma. European Journal of Cancer, 2012, 48, 3052-3062.	1.3	17
113	Diagnosis and treatment of melanoma. European consensus-based interdisciplinary guideline – Update 2012. European Journal of Cancer, 2012, 48, 2375-2390.	1.3	407
114	EORTC Melanoma Group achievements. European Journal of Cancer, Supplement, 2012, 10, 112-119.	2.2	0
115	Melanocortin-1 receptor, skin cancer and phenotypic characteristics (M-SKIP) project: study design and methods for pooling results of genetic epidemiological studies. BMC Medical Research Methodology, 2012, 12, 116.	1.4	12
116	Identifying melanomas in primary care: can we do better?. BMJ, The, 2012, 345, e4244-e4244.	3.0	6
117	Relationship between sunbed use and melanoma risk in a large case-control study in the United Kingdom. International Journal of Cancer, 2012, 130, 3011-3013.	2.3	17
118	Genome-wide association study identifies three new melanoma susceptibility loci. Nature Genetics, 2011, 43, 1108-1113.	9.4	230
119	Melanoma and vitamin D. Molecular Oncology, 2011, 5, 197-214.	2.1	58
120	Relationship between sun exposure and melanoma risk for tumours in different body sites in a large case-control study in a temperate climate. European Journal of Cancer, 2011, 47, 732-741.	1.3	90
121	Infliximab in the treatment of a child with cutaneous granulomas associated with ataxia telangiectasia. Journal of the American Academy of Dermatology, 2011, 65, 676-677.	0.6	28
122	Pathway-Based Analysis of a Melanoma Genome-Wide Association Study: Analysis of Genes Related to Tumour-Immunosuppression. PLoS ONE, 2011, 6, e29451.	1.1	18
123	The determinants of periorbital skin ageing in participants of a melanoma case-control study in the U.K.. British Journal of Dermatology, 2011, 165, 1011-1021.	1.4	17
124	The determinants of serum vitamin D levels in participants in a melanoma case-control study living in a temperate climate. Cancer Causes and Control, 2011, 22, 1471-1482.	0.8	32
125	Melanoma risk for CDKN2A mutation carriers who are relatives of population-based case carriers in Australia and the UK. Journal of Medical Genetics, 2011, 48, 266-272.	1.5	41
126	Prognosis in Patients With Sentinel Node-Positive Melanoma Is Accurately Defined by the Combined Rotterdam Tumor Load and Dewar Topography Criteria. Journal of Clinical Oncology, 2011, 29, 2206-2214.	0.8	195

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127	A novel recurrent mutation in MITF predisposes to familial and sporadic melanoma. <i>Nature</i> , 2011, 480, 99-103.	13.7	413
128	Genome-wide association study identifies a new melanoma susceptibility locus at 1q21.3. <i>Nature Genetics</i> , 2011, 43, 1114-1118.	9.4	140
129	Melanoma susceptibility genes. <i>Melanoma Research</i> , 2010, 20, 161-162.	0.6	5
130	Melanoma risk factors, perceived threat and intentional tanning: an international online survey. <i>European Journal of Cancer Prevention</i> , 2010, 19, 216-226.	0.6	47
131	IRF4 Variants Have Age-Specific Effects on Nevus Count and Predispose to Melanoma. <i>American Journal of Human Genetics</i> , 2010, 87, 6-16.	2.6	114
132	Deletion at chromosome arm 9p in relation to <i>BRAF</i> / <i>NRAS</i> mutations and prognostic significance for primary melanoma. <i>Genes Chromosomes and Cancer</i> , 2010, 49, 425-438.	1.5	46
133	Predictors of Sun Protection Behaviors and Severe Sunburn in an International Online Study. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2010, 19, 2199-2210.	1.1	106
134	Patterns of Expression of DNA Repair Genes and Relapse From Melanoma. <i>Clinical Cancer Research</i> , 2010, 16, 5211-5221.	3.2	53
135	Melanocytic Nevi, Nevus Genes, and Melanoma Risk in a Large Case-Control Study in the United Kingdom. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2010, 19, 2043-2054.	1.1	102
136	Reply to P.E. Hutchinson et al. <i>Journal of Clinical Oncology</i> , 2010, 28, e494-e495.	0.8	0
137	Diagnosis and treatment of melanoma: European consensus-based interdisciplinary guideline. <i>European Journal of Cancer</i> , 2010, 46, 270-283.	1.3	284
138	Laryngo-Onycho-Cutaneous Syndrome. <i>Ophthalmology</i> , 2010, 117, 1056-1056.e2.	2.5	7
139	Melanoma sentinel node biopsy and prediction models for relapse and overall survival. <i>British Journal of Cancer</i> , 2010, 103, 1229-1236.	2.9	54
140	Sun exposure and melanoma risk at different latitudes: a pooled analysis of 5700 cases and 7216 controls. <i>International Journal of Epidemiology</i> , 2009, 38, 814-830.	0.9	219
141	Gene Expression Profiling of Paraffin-Embedded Primary Melanoma Using the DASL Assay Identifies Increased Osteopontin Expression as Predictive of Reduced Relapse-Free Survival. <i>Clinical Cancer Research</i> , 2009, 15, 6939-6946.	3.2	93
142	Serum 25-Hydroxyvitamin D ₃ Levels Are Associated With Breslow Thickness at Presentation and Survival From Melanoma. <i>Journal of Clinical Oncology</i> , 2009, 27, 5439-5444.	0.8	263
143	Cutaneous Melanoma During Pregnancy: Is the Controversy Over?. <i>Journal of Clinical Oncology</i> , 2009, 27, e11-e12.	0.8	22
144	A pooled analysis of melanocytic nevus phenotype and the risk of cutaneous melanoma at different latitudes. <i>International Journal of Cancer</i> , 2009, 124, 420-428.	2.3	84

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145	Overseas Sun Exposure, Nevus Counts, and Premature Skin Aging in Young English Women: A Population-Based Survey. <i>Journal of Investigative Dermatology</i> , 2009, 129, 50-59.	0.3	40
146	Genome-wide association study identifies variants at 9p21 and 22q13 associated with development of cutaneous nevi. <i>Nature Genetics</i> , 2009, 41, 915-919.	9.4	204
147	Genome-wide association study identifies three loci associated with melanoma risk. <i>Nature Genetics</i> , 2009, 41, 920-925.	9.4	422
148	Vitamin D receptor gene polymorphisms, serum 25-hydroxyvitamin D levels, and melanoma: UK case-control comparisons and a meta-analysis of published VDR data. <i>European Journal of Cancer</i> , 2009, 45, 3271-3281.	1.3	127
149	Selection criteria for genetic assessment of patients with familial melanoma. <i>Journal of the American Academy of Dermatology</i> , 2009, 61, 677.e1-677.e14.	0.6	154
150	The emergence of networks in human genome epidemiology: challenges and opportunities. , 2009, , 120-134.		0
151	Trends in prognostic factors and survival from cutaneous melanoma in Yorkshire, UK and New South Wales, Australia between 1993 and 2003. <i>International Journal of Cancer</i> , 2008, 123, 861-866.	2.3	43
152	Spontaneous involution of congenital melanocytic nevi of the scalp. <i>Journal of the American Academy of Dermatology</i> , 2008, 58, 508-511.	0.6	58
153	Environmental risk factors for relapse of melanoma. <i>European Journal of Cancer</i> , 2008, 44, 1717-1725.	1.3	18
154	Frequent p16-Independent Inactivation of p14ARF in Human Melanoma. <i>Journal of the National Cancer Institute</i> , 2008, 100, 784-795.	3.0	94
155	CDKN2A and CDK4 variants in Latvian melanoma patients: analysis of a clinic-based population. <i>Melanoma Research</i> , 2007, 17, 185-191.	0.6	34
156	The Emergence of Networks in Human Genome Epidemiology. <i>Epidemiology</i> , 2007, 18, 1-8.	1.2	102
157	Sun-Protective Behaviors in Families at Increased Risk of Melanoma. <i>Journal of Investigative Dermatology</i> , 2007, 127, 1343-1350.	0.3	14
158	Genetics: What Advice for Patients Who Present With a Family History of Melanoma?. <i>Seminars in Oncology</i> , 2007, 34, 452-459.	0.8	21
159	A road map for efficient and reliable human genome epidemiology. <i>Nature Genetics</i> , 2006, 38, 3-5.	9.4	244
160	A mutation hotspot at the p14ARF splice site. <i>Oncogene</i> , 2005, 24, 4604-4608.	2.6	70
161	The Effect of Sun Exposure in Determining Nevus Density in UK Adolescent Twins. <i>Journal of Investigative Dermatology</i> , 2005, 124, 56-62.	0.3	62
162	Intronic sequence variants of the CDKN2A gene in melanoma pedigrees. <i>Genes Chromosomes and Cancer</i> , 2005, 43, 128-136.	1.5	32

#	ARTICLE	IF	CITATIONS
163	Prevalence of 9p21 deletions in UK melanoma families. <i>Genes Chromosomes and Cancer</i> , 2005, 44, 292-300.	1.5	36
164	A Network of Investigator Networks in Human Genome Epidemiology. <i>American Journal of Epidemiology</i> , 2005, 162, 302-304.	1.6	104
165	No Evidence for BRAF as a Melanoma/Nevus Susceptibility Gene. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2005, 14, 913-918.	1.1	24
166	Malignant Melanoma in Pregnancy. <i>Obstetrics and Gynecology Clinics of North America</i> , 2005, 32, 559-568.	0.7	20
167	The genetics of susceptibility to cutaneous melanoma. <i>Drugs of Today</i> , 2005, 41, 193.	2.4	33
168	Effect of Pregnancy on Survival in Women With Cutaneous Malignant Melanoma. <i>Journal of Clinical Oncology</i> , 2004, 22, 4369-4375.	0.8	164
169	The Relationship Between the Epidermal Growth Factor (EGF) 5'UTR Variant A61G and Melanoma/Nevus Susceptibility. <i>Journal of Investigative Dermatology</i> , 2004, 123, 755-759.	0.3	39
170	An Assessment of a Variant of the DNA Repair Gene XRCC3 as a Possible Nevus or Melanoma Susceptibility Genotype. <i>Journal of Investigative Dermatology</i> , 2004, 122, 429-432.	0.3	25
171	A Quality-of-Life Study in High-Risk (Thickness ≥ 2 mm) Cutaneous Melanoma Patients in a Randomized Trial of 1-cm versus 3-cm Surgical Excision Margins. <i>Journal of Investigative Dermatology Symposium Proceedings</i> , 2004, 9, 152-159.	0.8	45
172	Surgical margins in cutaneous melanoma (2 cm versus 5 cm for lesions measuring less than 2.1-mm) T_j ETQq0 0 0 rBT /Overlock 10 Tf	2.0	4
173	Excision Margins in High-Risk Malignant Melanoma. <i>New England Journal of Medicine</i> , 2004, 350, 757-766.	13.9	424
174	Biallelic Mutations in p16 INK4a Confer Resistance to Ras- and Ets-Induced Senescence in Human Diploid Fibroblasts. <i>Molecular and Cellular Biology</i> , 2002, 22, 8135-8143.	1.1	112
175	Excision Margins in the Treatment of Primary Cutaneous Melanoma. <i>Archives of Surgery</i> , 2002, 137, 1101-5.	2.3	73
176	UK guidelines for the management of cutaneous melanoma. <i>Journal of Plastic, Reconstructive and Aesthetic Surgery</i> , 2002, 55, 46-54.	1.1	80
177	Management of regional lymph nodes in patients with malignant melanoma: questionnaire survey of UK current practice. <i>Journal of Plastic, Reconstructive and Aesthetic Surgery</i> , 2002, 55, 372-375.	1.1	13
178	An Assessment of the CDKN2A Variant Ala148Thr as a Nevus/Melanoma Susceptibility Allele. <i>Journal of Investigative Dermatology</i> , 2002, 119, 961-965.	0.3	36
179	Heritability and Gene-Environment Interactions for Melanocytic Nevus Density Examined in a U.K. Adolescent Twin Study. <i>Journal of Investigative Dermatology</i> , 2001, 117, 348-352.	0.3	112
180	Melanoma. <i>British Journal of Hospital Medicine</i> , 2000, 61, 103-107.	0.3	1

#	ARTICLE	IF	CITATIONS
181	Genotype/Phenotype and Penetrance Studies in Melanoma Families with Germline CDKN2A Mutations. Journal of Investigative Dermatology, 2000, 114, 28-33.	0.3	102
182	Mutation testing in melanoma families: INK4A, CDK4 and INK4D. British Journal of Cancer, 1999, 80, 295-300.	2.9	57
183	Counseling and DNA Testing for Individuals Perceived to Be Genetically Predisposed to Melanoma: A Consensus Statement of the Melanoma Genetics Consortium. Journal of Clinical Oncology, 1999, 17, 3245-3251.	0.8	209
184	Haplotype analysis of two recurrent CDKN2A mutations in 10 melanoma families: Evidence for common founders and independent mutations. , 1998, 11, 424-431.		61
185	The Atypical-Mole Syndrome and Predisposition to Melanoma. New England Journal of Medicine, 1998, 339, 348-349.	13.9	39
186	Molecular pathology of melanoma. , 1997, 16, 141-154.		25
187	Naevi and pigmentary characteristics as risk factors for melanoma in a high-risk population: A case-control study in new South Wales, Australia. , 1996, 67, 485-491.		114
188	Risk of ocular melanoma in relation to cutaneous and IRIS naevi. International Journal of Cancer, 1995, 60, 622-626.	2.3	58
189	Genetic heterogeneity in familial malignant melanoma. Human Molecular Genetics, 1994, 3, 2195-2200.	1.4	66
190	Melanoma and reproductive health. , 0, , 257-264.		0
191	Surgical excision methods for skin cancer involving the nail unit. The Cochrane Library, 0, , .	1.5	1