

Rajiv Kumar

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.

238
papers

13,271
citations

59
h-index

108
g-index

251
ext. papers

14,993
ext. citations

6.8
avg, IF

5.78
L-index

#	Paper	IF	Citations
238	Mutational Characterization of Cutaneous Melanoma Supports Divergent Pathways Model for Melanoma Development. <i>Cancers</i> , 2021 , 13,	6.6	1
237	Longitudinal study of prognostic factors for localized cutaneous melanoma in patients who have been disease-free for five years. <i>European Journal of Dermatology</i> , 2021 , 31, 192-198	0.8	1
236	Clinical, environmental and histological distribution of BRAF, NRAS and TERT promoter mutations among patients with cutaneous melanoma: a retrospective study of 563 patients. <i>British Journal of Dermatology</i> , 2021 , 184, 504-513	4	9
235	Polymorphisms in CXCR3 ligands predict early CXCL9 recovery and severe chronic GVHD. <i>Blood Cancer Journal</i> , 2021 , 11, 42	7	1
234	KIT mutational status does not constitute an independent prognostic marker in cutaneous melanoma. A study on 688 Spanish patients. <i>Melanoma Research</i> , 2021 , 31, 101-103	3.3	
233	Occurrence, functionality and abundance of the TERT promoter mutations. <i>International Journal of Cancer</i> , 2021 , 149, 1852-1862	7.5	2
232	Association of HERV-K and LINE-1 hypomethylation with reduced disease-free survival in melanoma patients. <i>Epigenomics</i> , 2020 , 12, 1689-1706	4.4	4
231	Telomerase Reverse Transcriptase Promoter Mutations Identify a Genomically Defined and Highly Aggressive Human Pleural Mesothelioma Subgroup. <i>Clinical Cancer Research</i> , 2020 , 26, 3819-3830	12.9	8
230	Mutation Signatures in Melanocytic Nevi Reveal Characteristics of Defective DNA Repair. <i>Journal of Investigative Dermatology</i> , 2020 , 140, 2093-2096.e2	4.3	2
229	Risk factors for the development of a second melanoma in patients with cutaneous melanoma. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2020 , 34, 2295-2302	4.6	7
228	Telomeres and Telomere Length: A General Overview. <i>Cancers</i> , 2020 , 12,	6.6	56
227	Assessment of polygenic architecture and risk prediction based on common variants across fourteen cancers. <i>Nature Communications</i> , 2020 , 11, 3353	17.4	32
226	TERT promoter mutations in actinic keratosis before and after treatment. <i>International Journal of Cancer</i> , 2020 , 146, 2932-2934	7.5	1
225	Genome-wide characterization of 5-hydroxymethylcytosine in melanoma reveals major differences with nevus. <i>Genes Chromosomes and Cancer</i> , 2020 , 59, 366-374	5	3
224	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	36.3	39
223	Chromosomal damage and telomere length in peripheral blood lymphocytes of cancer patients. <i>Oncology Reports</i> , 2020 , 44, 2219-2230	3.5	0
222	Coding and noncoding somatic mutations in candidate genes in basal cell carcinoma. <i>Scientific Reports</i> , 2020 , 10, 8005	4.9	13

221	Telomere length in peripheral blood lymphocytes related to genetic variation in telomerase, prognosis and clinicopathological features in breast cancer patients. <i>Mutagenesis</i> , 2020 , 35, 491-497	2.8	2
220	Expression quantitative trait loci in ABC transporters are associated with survival in 5-FU treated colorectal cancer patients. <i>Mutagenesis</i> , 2020 , 35, 273-281	2.8	1
219	Informing patients about their mutation tests: CDKN2A c.256G>A in melanoma as an example. <i>Hereditary Cancer in Clinical Practice</i> , 2020 , 18, 15	2.3	0
218	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	3.3	2
217	Lymphatic and Hematogenous Dissemination in Patients With Primary Cutaneous Melanoma-Reply. <i>JAMA Dermatology</i> , 2019 , 155, 1323	5.1	
216	Risk Factors for Lymphatic and Hematogenous Dissemination in Patients With Stages I to II Cutaneous Melanoma. <i>JAMA Dermatology</i> , 2019 , 155, 679-687	5.1	13
215	Telomere length, arsenic exposure and risk of basal cell carcinoma of skin. <i>Carcinogenesis</i> , 2019 , 40, 715-723	4.3	10
214	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	14.5	8
213	Promoter Mutations are Associated with Visceral Spreading in Melanoma of the Trunk. <i>Cancers</i> , 2019 , 11,	6.6	11
212	TERT promoter mutation subtypes and survival in stage I and II melanoma patients. <i>International Journal of Cancer</i> , 2019 , 144, 1027-1036	7.5	27
211	TERT expression is susceptible to BRAF and ETS-factor inhibition in BRAF/TERT promoter double-mutated glioma. <i>Acta Neuropathologica Communications</i> , 2019 , 7, 128	7.3	18
210	Relationship of telomere length in colorectal cancer patients with cancer phenotype and patient prognosis. <i>British Journal of Cancer</i> , 2019 , 121, 344-350	8.7	14
209	The CXCL9 Polymorphism rs884304 Associates with Early CXCL9 Reconstitution and with Severe Chronic Graft-Versus-Host Disease (cGVHD) in Human Allograft Recipients. <i>Blood</i> , 2019 , 134, 873-873	2.2	
208	TERT promoter mutation subtypes in 20 in-situ melanomas. <i>Melanoma Research</i> , 2019 , 29, 347-348	3.3	
207	Asymmetric dimethylarginine serum levels are associated with early mortality after allogeneic stem cell transplantation. <i>Haematologica</i> , 2019 , 104, 827-834	6.6	3
206	Genome-wide association study identifies susceptibility loci for B-cell childhood acute lymphoblastic leukemia. <i>Nature Communications</i> , 2018 , 9, 1340	17.4	39
205	Bleomycin-induced chromosomal damage and shortening of telomeres in peripheral blood lymphocytes of incident cancer patients. <i>Genes Chromosomes and Cancer</i> , 2018 , 57, 61-69	5	10
204	Characteristics of Familial Melanoma in Valencia, Spain, Based on the Presence of CDKN2A Mutations and MC1R Variants. <i>Acta Dermato-Venereologica</i> , 2018 , 98, 512-516	2.2	

203	TERT promoter mutations are associated with poor prognosis and cell immortalization in meningioma. <i>Neuro-Oncology</i> , 2018 , 20, 1584-1593	1	58
202	Telomere length and survival in primary cutaneous melanoma patients. <i>Scientific Reports</i> , 2018 , 8, 109474-9	4.9	11
201	Leukocyte telomere length throughout the continuum of colorectal carcinogenesis. <i>Oncotarget</i> , 2018 , 9, 13582-13592	3.3	4
200	Genetic Risk of Severe Chronic Graft-Versus-Host Disease Defined By Host-Derived CXCR3 Ligands. <i>Blood</i> , 2018 , 132, 357-357	2.2	
199	Single Nucleotide Polymorphisms in CD40L Predict Endothelial Complications and Mortality After Allogeneic Stem-Cell Transplantation. <i>Journal of Clinical Oncology</i> , 2018 , 36, 789-800	2.2	14
198	Defective DNA damage repair leads to frequent catastrophic genomic events in murine and human tumors. <i>Nature Communications</i> , 2018 , 9, 4760	17.4	37
197	P02.10 Aggressiveness of meningiomas is predicted by cell immortalization in vitro and dependent on TERT promoter mutations. <i>Neuro-Oncology</i> , 2018 , 20, iii274-iii274	1	78
196	Telomere length, telomerase reverse transcriptase promoter mutations, and melanoma risk. <i>Genes Chromosomes and Cancer</i> , 2018 , 57, 564-572	5	24
195	Altered TERT promoter and other genomic regulatory elements: occurrence and impact. <i>International Journal of Cancer</i> , 2017 , 141, 867-876	7.5	14
194	TERT promoter mutations in telomere biology. <i>Mutation Research - Reviews in Mutation Research</i> , 2017 , 771, 15-31	7	93
193	TERT promoter mutations associate with MC1R variants in melanoma patients. <i>Pigment Cell and Melanoma Research</i> , 2017 , 30, 273-275	4.5	7
192	Distribution of TERT promoter mutations in primary and metastatic melanomas in Austrian patients. <i>Journal of Cancer Research and Clinical Oncology</i> , 2017 , 143, 613-617	4.9	12
191	TERT promoter mutations are not always associated with poor prognosis in atypical spitzoid tumors. <i>Pigment Cell and Melanoma Research</i> , 2017 , 30, 265-268	4.5	9
190	Genetic alterations in seborrheic keratoses. <i>Oncotarget</i> , 2017 , 8, 36639-36649	3.3	28
189	New prognostic factor telomerase reverse transcriptase promoter mutation presents without MR imaging biomarkers in primary glioblastoma. <i>Neuroradiology</i> , 2017 , 59, 1223-1231	3.2	9
188	A genome-wide association study identifies risk loci for childhood acute lymphoblastic leukemia at 10q26.13 and 12q23.1. <i>Leukemia</i> , 2017 , 31, 573-579	10.7	52
187	Cutaneous melanoma primary site is linked to nevus density. <i>Oncotarget</i> , 2017 , 8, 98876-98886	3.3	5
186	ETS1, nucleolar and non-nucleolar TERT expression in nevus to melanoma progression. <i>Oncotarget</i> , 2017 , 8, 104408-104417	3.3	6

185	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	4.3	12
184	Identification of a novel susceptibility locus at 13q34 and refinement of the 20p12.2 region as a multi-signal locus associated with bladder cancer risk in individuals of European ancestry. <i>Human Molecular Genetics</i> , 2016 , 25, 1203-14	5.6	20
183	PARKIN Inactivation Links Parkinson's Disease to Melanoma. <i>Journal of the National Cancer Institute</i> , 2016 , 108,	9.7	46
182	Characterization of individuals at high risk of developing melanoma in Latin America: bases for genetic counseling in melanoma. <i>Genetics in Medicine</i> , 2016 , 18, 727-36	8.1	18
181	Risk factors for keratinocyte skin cancer in patients diagnosed with melanoma, a large retrospective study. <i>European Journal of Cancer</i> , 2016 , 53, 115-24	7.5	10
180	Mapping of deletion breakpoints at the CDKN2A locus in melanoma: detection of MTAP-ANRIL fusion transcripts. <i>Oncotarget</i> , 2016 , 7, 16490-504	3.3	18
179	Thrombosis in a population of phase I trial patients.. <i>Journal of Clinical Oncology</i> , 2016 , 34, e14054-e14054	5.4	4
178	MELADIAG: A panel of DNA biomarkers dedicated to melanoma diagnosis.. <i>Journal of Clinical Oncology</i> , 2016 , 34, e23258-e23258	2.2	2
177	TERT promoter mutations in melanoma render TERT expression dependent on MAPK pathway activation. <i>Oncotarget</i> , 2016 , 7, 53127-53136	3.3	42
176	SERPINB1 expression is predictive for sensitivity and outcome of cisplatin-based chemotherapy in melanoma. <i>Oncotarget</i> , 2016 , 7, 10117-32	3.3	12
175	TERT promoter mutations associate with fast-growing melanoma. <i>Pigment Cell and Melanoma Research</i> , 2016 , 29, 236-8	4.5	32
174	TERT promoter mutations in melanoma survival. <i>International Journal of Cancer</i> , 2016 , 139, 75-84	7.5	79
173	Genome-wide meta-analysis identifies five new susceptibility loci for cutaneous malignant melanoma. <i>Nature Genetics</i> , 2015 , 47, 987-995	36.3	162
172	A novel c. 204 Ile68Met germline variant in exon 2 of the mutL homolog 1 gene in a colorectal cancer patient. <i>Oncology Letters</i> , 2015 , 9, 183-186	2.6	2
171	MC1R gene variants and non-melanoma skin cancer: a pooled-analysis from the M-SKIP project. <i>British Journal of Cancer</i> , 2015 , 113, 354-63	8.7	26
170	New basal cell carcinoma susceptibility loci. <i>Nature Communications</i> , 2015 , 6, 6825	17.4	49
169	Genes involved in the WNT and vesicular trafficking pathways are associated with melanoma predisposition. <i>International Journal of Cancer</i> , 2015 , 136, 2109-19	7.5	21
168	Telomere length in circulating lymphocytes: Association with chromosomal aberrations. <i>Genes Chromosomes and Cancer</i> , 2015 , 54, 194-6	5	11

167	TERT promoter mutations in clear cell renal cell carcinoma. <i>International Journal of Cancer</i> , 2015 , 136, 2448-52	7.5	36
166	TERT promoter mutations: a novel independent prognostic factor in primary glioblastomas. <i>Neuro-Oncology</i> , 2015 , 17, 45-52	1	123
165	MC1R 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, 618-31	7.5	67
164	Genetic variation in arsenic (+3 oxidation state) methyltransferase (AS3MT), arsenic metabolism and risk of basal cell carcinoma in a European population. <i>Environmental and Molecular Mutagenesis</i> , 2015 , 56, 60-9	3.2	40
163	The 9p21.3 risk of childhood acute lymphoblastic leukaemia is explained by a rare high-impact variant in CDKN2A. <i>Scientific Reports</i> , 2015 , 5, 15065	4.9	17
162	Age-related characteristics of cutaneous melanoma in a Spanish Mediterranean population. <i>International Journal of Dermatology</i> , 2015 , 54, 778-84	1.7	11
161	Mutations in TERT promoter and FGFR3 and telomere length in bladder cancer. <i>International Journal of Cancer</i> , 2015 , 137, 1621-9	7.5	65
160	RICTOR involvement in the PI3K/AKT pathway regulation in melanocytes and melanoma. <i>Oncotarget</i> , 2015 , 6, 28120-31	3.3	24
159	BRAF, NRAS and MC1R status in a prospective series of primary cutaneous melanoma. <i>British Journal of Dermatology</i> , 2015 , 172, 1128-31	4	14
158	TERT promoter mutations and telomere length in adult malignant gliomas and recurrences. <i>Oncotarget</i> , 2015 , 6, 10617-33	3.3	60
157	Frequent DPH3 promoter mutations in skin cancers. <i>Oncotarget</i> , 2015 , 6, 35922-30	3.3	42
156	A retrospective comparative exploratory study on two methylenetetrahydrofolate reductase (MTHFR) polymorphisms in esophagogastric cancer: the A1298C MTHFR polymorphism is an independent prognostic factor only in neoadjuvantly treated gastric cancer patients. <i>BMC Cancer</i> , 2014 , 14, 58	4.8	15
155	Telomerase reverse transcriptase promoter mutations in primary cutaneous melanoma. <i>Nature Communications</i> , 2014 , 5, 3401	17.4	132
154	Genome-wide association study yields variants at 20p12.2 that associate with urinary bladder cancer. <i>Human Molecular Genetics</i> , 2014 , 23, 5545-57	5.6	29
153	Clinical Characteristics of Patients With Cutaneous Melanoma According to Variants in the Melanocortin 1 Receptor Gene. <i>Actas Dermo-sifiliograficas</i> , 2014 , 105, 159-171	0.5	
152	Inherited variability in a master regulator polymorphism (rs4846126) associates with survival in 5-FU treated colorectal cancer patients. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2014 , 766-767, 7-13	3.3	2
151	Germline sequence variants in TGM3 and RGS22 confer risk of basal cell carcinoma. <i>Human Molecular Genetics</i> , 2014 , 23, 3045-53	5.6	39
150	Polymorphisms in DNA repair genes XRCC1 and XRCC3, occupational exposure to arsenic and sunlight, and the risk of non-melanoma skin cancer in a European case-control study. <i>Environmental Research</i> , 2014 , 134, 382-9	7.9	6

149	Single-Nucleotide Polymorphisms Within the Thrombomodulin Gene (THBD) Predict Mortality in Patients With Graft-Versus-Host Disease. <i>Journal of Clinical Oncology</i> , 2014 , 32, 3421-7	2.2	36
148	Inherited variation in the PARP1 gene and survival from melanoma. <i>International Journal of Cancer</i> , 2014 , 135, 1625-33	7.5	22
147	An inherited variant in the gene coding for vitamin D-binding protein and survival from cutaneous melanoma: a BioGenoMEL study. <i>Pigment Cell and Melanoma Research</i> , 2014 , 27, 234-43	4.5	21
146	Early epigenetic downregulation of WNK2 kinase during pancreatic ductal adenocarcinoma development. <i>Oncogene</i> , 2014 , 33, 3401-10	9.2	24
145	TERT promoter mutations in cancer development. <i>Current Opinion in Genetics and Development</i> , 2014 , 24, 30-7	4.9	167
144	Clinical characteristics of patients with cutaneous melanoma according to variants in the melanocortin 1 receptor gene. <i>Actas Dermo-sifiliograficas</i> , 2014 , 105, 159-71	0.5	3
143	Variants at chromosome 20 (ASIP locus) and melanoma risk. <i>International Journal of Cancer</i> , 2013 , 132, 42-54	7.5	19
142	Distribution of MC1R variants among melanoma subtypes: p.R163Q is associated with lentigo maligna melanoma in a Mediterranean population. <i>British Journal of Dermatology</i> , 2013 , 169, 804-11	4	20
141	Variants at the 9p21 locus and melanoma risk. <i>BMC Cancer</i> , 2013 , 13, 325	4.8	23
140	Occupational exposure to arsenic and risk of nonmelanoma skin cancer in a multinational European study. <i>International Journal of Cancer</i> , 2013 , 133, 2182-91	7.5	35
139	TERT promoter mutations in familial and sporadic melanoma. <i>Science</i> , 2013 , 339, 959-61	33.3	1261
138	TERT promoter mutations in bladder cancer affect patient survival and disease recurrence through modification by a common polymorphism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 17426-31	11.5	236
137	Variation at 10p12.2 and 10p14 influences risk of childhood B-cell acute lymphoblastic leukemia and phenotype. <i>Blood</i> , 2013 , 122, 3298-307	2.2	119
136	Occupational exposure to ultraviolet radiation and risk of non-melanoma skin cancer in a multinational European study. <i>PLoS ONE</i> , 2013 , 8, e62359	3.7	40
135	Gene expression variations: potentialities of master regulator polymorphisms in colorectal cancer risk. <i>Mutagenesis</i> , 2012 , 27, 161-7	2.8	12
134	Functional, genetic, and epigenetic aspects of base and nucleotide excision repair in colorectal carcinomas. <i>Clinical Cancer Research</i> , 2012 , 18, 5878-87	12.9	59
133	Inherited variants in the MC1R gene and survival from cutaneous melanoma: a BioGenoMEL study. <i>Pigment Cell and Melanoma Research</i> , 2012 , 25, 384-94	4.5	50
132	Melanocortin-1 receptor, skin cancer and phenotypic characteristics (M-SKIP) project: study design and methods for pooling results of genetic epidemiological studies. <i>BMC Medical Research Methodology</i> , 2012 , 12, 116	4.7	10

131	Inorganic arsenic and basal cell carcinoma in areas of Hungary, Romania, and Slovakia: a case-control study. <i>Environmental Health Perspectives</i> , 2012 , 120, 721-6	8.4	77
130	Mutations and polymorphisms in TP53 gene--an overview on the role in colorectal cancer. <i>Mutagenesis</i> , 2012 , 27, 211-8	2.8	80
129	POMC and TP53 genetic variability and risk of basal cell carcinoma of skin: Interaction between host and genetic factors. <i>Journal of Dermatological Science</i> , 2011 , 63, 47-54	4.3	13
128	DNA repair gene and MTHFR gene polymorphisms as prognostic markers in locally advanced adenocarcinoma of the esophagus or stomach treated with cisplatin and 5-fluorouracil-based neoadjuvant chemotherapy. <i>Annals of Surgical Oncology</i> , 2011 , 18, 2688-98	3.1	46
127	A germline variant in the TP53 polyadenylation signal confers cancer susceptibility. <i>Nature Genetics</i> , 2011 , 43, 1098-103	36.3	203
126	Exposure to low environmental levels of benzene: evaluation of micronucleus frequencies and S-phenylmercapturic acid excretion in relation to polymorphisms in genes encoding metabolic enzymes. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , 2011 , 719, 7-13	3	38
125	MTHFR and MTRR genotype and haplotype analysis and colorectal cancer susceptibility in a case-control study from the Czech Republic. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , 2011 , 721, 74-80	3	40
124	Pancreatic cancer susceptibility loci and their role in survival. <i>PLoS ONE</i> , 2011 , 6, e27921	3.7	41
123	Familial bladder cancer and the related genes. <i>Current Opinion in Urology</i> , 2011 , 21, 386-92	2.8	14
122	Defining fast-growing melanomas: reappraisal of epidemiological, clinical, and histological features. <i>Melanoma Research</i> , 2011 , 21, 131-8	3.3	35
121	Rationale for an international consortium to study inherited genetic susceptibility to childhood acute lymphoblastic leukemia. <i>Haematologica</i> , 2011 , 96, 1049-54	6.6	32
120	5-Fluorouracil-based chemotherapy for colorectal cancer and MTHFR/MTRR genotypes. <i>British Journal of Clinical Pharmacology</i> , 2011 , 72, 162-3	3.8	52
119	European genome-wide association study identifies SLC14A1 as a new urinary bladder cancer susceptibility gene. <i>Human Molecular Genetics</i> , 2011 , 20, 4268-81	5.6	105
118	DNA damage, DNA repair rates and mRNA expression levels of cell cycle genes (TP53, p21(CDKN1A), BCL2 and BAX) with respect to occupational exposure to styrene. <i>Carcinogenesis</i> , 2011 , 32, 74-9	4.6	5
117	Interaction between functional polymorphic variants in cytokine genes, established risk factors and susceptibility to basal cell carcinoma of skin. <i>Carcinogenesis</i> , 2011 , 32, 1849-54	4.6	16
116	Epigenetic deregulation of TCF21 inhibits metastasis suppressor KISS1 in metastatic melanoma. <i>Carcinogenesis</i> , 2011 , 32, 1467-73	4.6	54
115	ERCC5 p.Asp1104His and ERCC2 p.Lys751Gln polymorphisms are independent prognostic factors for the clinical course of melanoma. <i>Journal of Investigative Dermatology</i> , 2011 , 131, 1280-90	4.3	23
114	A sequence variant at 4p16.3 confers susceptibility to urinary bladder cancer. <i>Nature Genetics</i> , 2010 , 42, 415-9	36.3	138

113	Variation in CDKN2A at 9p21.3 influences childhood acute lymphoblastic leukemia risk. <i>Nature Genetics</i> , 2010 , 42, 492-4	36.3	214
112	A multi-stage genome-wide association study of bladder cancer identifies multiple susceptibility loci. <i>Nature Genetics</i> , 2010 , 42, 978-84	36.3	408
111	Association between the germline MC1R variants and somatic BRAF/NRAS mutations in melanoma tumors. <i>Journal of Investigative Dermatology</i> , 2010 , 130, 2844-8	4.3	25
110	Single-nucleotide polymorphisms in DNA-repair genes and cutaneous melanoma. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , 2010 , 702, 8-16	3	26
109	Verification of the susceptibility loci on 7p12.2, 10q21.2, and 14q11.2 in precursor B-cell acute lymphoblastic leukemia of childhood. <i>Blood</i> , 2010 , 115, 1765-7	2.2	107
108	Lifetime exposure to arsenic in residential drinking water in Central Europe. <i>International Archives of Occupational and Environmental Health</i> , 2010 , 83, 471-81	3.2	23
107	Genotypes, haplotypes and diplotypes of three XPC polymorphisms in urinary-bladder cancer patients. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2010 , 694, 39-44	3.3	16
106	Genetics of pigmentation in skin cancer--a review. <i>Mutation Research - Reviews in Mutation Research</i> , 2010 , 705, 141-153	7	121
105	Modulation of DNA repair capacity and mRNA expression levels of XRCC1, hOGG1 and XPC genes in styrene-exposed workers. <i>Toxicology and Applied Pharmacology</i> , 2010 , 248, 194-200	4.6	22
104	Somatic alterations in the melanoma genome: a high-resolution array-based comparative genomic hybridization study. <i>Genes Chromosomes and Cancer</i> , 2010 , 49, 733-45	5	79
103	XPC 939 and XRCC3 Polymorphismen sind Prognosefaktoren bei lokal fortgeschrittenen Adenokarzinomen des oberen Gastrointestinaltraktes nach neoadjuvanter 5-FU und Cisplatin basierter Chemotherapie. <i>Langenbecks Archiv Für Chirurgie Supplement</i> , 2009 , 99-101		
102	Polymorphisms in DNA repair genes, smoking, and bladder cancer risk: findings from the international consortium of bladder cancer. <i>Cancer Research</i> , 2009 , 69, 6857-64	10.1	94
101	Single nucleotide polymorphisms in DNA repair genes XRCC1 and APEX1 in progression and survival of primary cutaneous melanoma patients. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2009 , 661, 78-84	3.3	21
100	Genotype and haplotype analysis of cell cycle genes in sporadic colorectal cancer in the Czech Republic. <i>Human Mutation</i> , 2009 , 30, 661-8	4.7	41
99	Melanocortin receptor 1 variants and melanoma risk: a study of 2 European populations. <i>International Journal of Cancer</i> , 2009 , 125, 1868-75	7.5	56
98	A case-control study of childhood acute lymphoblastic leukaemia and polymorphisms in the TGF-beta and receptor genes. <i>Pediatric Blood and Cancer</i> , 2009 , 52, 819-23	3	8
97	Sequence variants at the TERT-CLPTM1L locus associate with many cancer types. <i>Nature Genetics</i> , 2009 , 41, 221-7	36.3	509
96	Genome-wide association study identifies five susceptibility loci for glioma. <i>Nature Genetics</i> , 2009 , 41, 899-904	36.3	640

95	New common variants affecting susceptibility to basal cell carcinoma. <i>Nature Genetics</i> , 2009 , 41, 909-14	36.3	275
94	Genetic variation in the prostate stem cell antigen gene PSCA confers susceptibility to urinary bladder cancer. <i>Nature Genetics</i> , 2009 , 41, 991-5	36.3	270
93	Malignant Melanoma— Genetic Overview. <i>Actas Dermo-sifiliograficas</i> , 2009 , 100, 38-51	0.5	12
92	Malignant melanoma—a genetic overview. <i>Actas Dermo-sifiliograficas</i> , 2009 , 100 Suppl 1, 38-51	0.5	5
91	ASIP and TYR pigmentation variants associate with cutaneous melanoma and basal cell carcinoma. <i>Nature Genetics</i> , 2008 , 40, 886-91	36.3	265
90	Sequence variant on 8q24 confers susceptibility to urinary bladder cancer. <i>Nature Genetics</i> , 2008 , 40, 1307-12	36.3	332
89	Common variants on 1p36 and 1q42 are associated with cutaneous basal cell carcinoma but not with melanoma or pigmentation traits. <i>Nature Genetics</i> , 2008 , 40, 1313-8	36.3	93
88	Inherited susceptibility to bleomycin-induced micronuclei: correlating polymorphisms in GSTT1, GSTM1 and DNA repair genes with mutagen sensitivity. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2008 , 638, 90-7	3.3	23
87	DNA repair genetic polymorphisms and risk of colorectal cancer in the Czech Republic. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2008 , 638, 146-53	3.3	95
86	Mutation analysis of the EGFR-NRAS-BRAF pathway in melanomas from black Africans and other subgroups of cutaneous melanoma. <i>Melanoma Research</i> , 2008 , 18, 29-35	3.3	66
85	MC1R variants associated susceptibility to basal cell carcinoma of skin: interaction with host factors and XRCC3 polymorphism. <i>International Journal of Cancer</i> , 2008 , 122, 1787-93	7.5	49
84	Three novel mutations of the EDA gene in Chinese patients with X-linked hypohidrotic ectodermal dysplasia. <i>British Journal of Dermatology</i> , 2008 , 158, 614-7	4	2
83	Influence of DNA repair gene polymorphisms on the initial repair of MMS-induced DNA damage in human lymphocytes as measured by the alkaline comet assay. <i>Environmental and Molecular Mutagenesis</i> , 2008 , 49, 669-75	3.2	7
82	B-RAF and N-RAS mutations are preserved during short time in vitro propagation and differentially impact prognosis. <i>PLoS ONE</i> , 2007 , 2, e236	3.7	107
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2	Telomere length, arsenic exposure and risk of basal cell carcinoma of skin		2
1	Assessment of Polygenic Architecture and Risk Prediction based on Common Variants Across Fourteen Cancers		1