

Grzegorz Kurzawski

List of Publications by Year in descending order

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87
papers

2,710
citations

185998

28
h-index

189595

50
g-index

88
all docs

88
docs citations

88
times ranked

3818
citing authors

#	ARTICLE	IF	CITATIONS
1	A genetic variant in telomerase reverse transcriptase (TERT) modifies cancer risk in Lynch syndrome patients harbouring pathogenic MSH2 variants. <i>Scientific Reports</i> , 2021, 11, 11401.	1.6	6
2	CD36 gene polymorphism and plasma sCD36 as the risk factor in higher cholesterolemia. <i>Archives De Pediatrie</i> , 2018, 25, 177-181.	0.4	10
3	New <i>EPCAM</i> founder deletion in Polish population. <i>Clinical Genetics</i> , 2017, 92, 649-653.	1.0	6
4	Cumulative Small Effect Genetic Markers and the Risk of Colorectal Cancer in Poland, Estonia, Lithuania, and Latvia. <i>Gastroenterology Research and Practice</i> , 2015, 2015, 1-10.	0.7	5
5	Management of ovarian and endometrial cancers in women belonging to HNPCC carrier families: review of the literature and results of cancer risk assessment in Polish HNPCC families. <i>Hereditary Cancer in Clinical Practice</i> , 2015, 13, 3.	0.6	11
6	Is plasma soluble CD36 associated with cardiovascular risk factors in early onset coronary artery disease patients?. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2015, 75, 398-406.	0.6	18
7	Polymorphisms in nucleotide excision repair genes and susceptibility to colorectal cancer in the Polish population. <i>Molecular Biology Reports</i> , 2015, 42, 755-764.	1.0	32
8	Cumulative effects of genetic markers and the detection of advanced colorectal neoplasias by population screening. <i>Clinical Genetics</i> , 2015, 88, 234-240.	1.0	5
9	Lynch syndrome mutations shared by the Baltic States and Poland. <i>Clinical Genetics</i> , 2014, 86, 190-193.	1.0	5
10	Germline deletions in the EPCAM gene as a cause of Lynch syndrome – literature review. <i>Hereditary Cancer in Clinical Practice</i> , 2013, 11, 9.	0.6	104
11	Combined analysis of three lynch syndrome cohorts confirms the modifying effects of 8q23.3 and 11q23.1 in MLH1 mutation carriers. <i>International Journal of Cancer</i> , 2013, 132, 1556-1564.	2.3	33
12	Is CD36 gene polymorphism in region encoding lipid-binding domain associated with early onset CAD?. <i>Gene</i> , 2013, 530, 134-137.	1.0	6
13	Association of CD36 gene polymorphisms with echo- and electrocardiographic parameters in patients with early onset coronary artery disease. <i>Archives of Medical Science</i> , 2013, 4, 640-650.	0.4	7
14	DNA repair gene polymorphisms and risk of early onset colorectal cancer in Lynch syndrome. <i>Cancer Epidemiology</i> , 2012, 36, 183-189.	0.8	25
15	DNA and RNA analyses in detection of genetic predisposition to cancer. <i>Hereditary Cancer in Clinical Practice</i> , 2012, 10, 17.	0.6	6
16	Fast diagnostic test for the identification of an increased genetic predisposition to colon cancer (exemplified on a DNA test for recurrent mutations of the gene MMR). <i>Hereditary Cancer in Clinical Practice</i> , 2012, 10, A13.	0.6	0
17	Polymorphism of the CD36 Gene and Cardiovascular Risk Factors in Patients with Coronary Artery Disease Manifested at a Young Age. <i>Biochemical Genetics</i> , 2012, 50, 103-111.	0.8	22
18	Polymorphism of CD36 gene, carbohydrate metabolism and plasma CD36 concentration in obese children. A preliminary study. <i>Postepy Higieny I Medycyny Doswiadczalnej</i> , 2012, 66, 954-958.	0.1	5

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19	CD36 gene is associated with thickness of atheromatous plaque and ankle-brachial index in patients with early coronary artery disease. <i>Kardiologia Polska</i> , 2012, 70, 918-23.	0.3	12
20	Association of MMP8 gene variation with an increased risk of malignant melanoma. <i>Melanoma Research</i> , 2011, 21, 464-468.	0.6	19
21	AMPD1 gene mutations are associated with obesity and diabetes in Polish patients with cardiovascular diseases. <i>Journal of Applied Genetics</i> , 2011, 52, 67-76.	1.0	16
22	Colorectal cancer susceptibility loci on chromosome 8q23.3 and 11q23.1 as modifiers for disease expression in lynch syndrome. <i>Journal of Medical Genetics</i> , 2011, 48, 279-284.	1.5	44
23	BRCA1 mutations and colorectal cancer in Poland. <i>Familial Cancer</i> , 2010, 9, 541-544.	0.9	33
24	Variant alleles of the CYP1B1 gene are associated with colorectal cancer susceptibility. <i>BMC Cancer</i> , 2010, 10, 420.	1.1	28
25	CHEK2 mutations and HNPCC-related colorectal cancer. <i>International Journal of Cancer</i> , 2010, 126, 3005-3009.	2.3	28
26	Analysis of Human <i>CD36</i> Gene Sequence Alterations in the Oxidized Low-Density Lipoprotein-Binding Region Using Denaturing High-Performance Liquid Chromatography. <i>Genetic Testing and Molecular Biomarkers</i> , 2010, 14, 551-557.	0.3	6
27	Combined iPLEX and TaqMan Assays to Screen for 45 Common Mutations in Lynch Syndrome and FAP Patients. <i>Journal of Molecular Diagnostics</i> , 2010, 12, 82-90.	1.2	8
28	Nonalcoholic fatty liver disease and <i>HFE</i> gene mutations: A Polish study. <i>World Journal of Gastroenterology</i> , 2010, 16, 2531.	1.4	29
29	HFE gene mutations in patients with alcoholic liver disease. A prospective study from northwestern Poland. , 2010, 120, 127-31.		2
30	Haemochromatosis <i>HFE</i> gene polymorphisms as potential modifiers of hereditary nonpolyposis colorectal cancer risk and onset age. <i>International Journal of Cancer</i> , 2009, 125, 78-83.	2.3	39
31	MTHFR 677 C>T and 1298 A>C polymorphisms and the age of onset of colorectal cancer in hereditary nonpolyposis colorectal cancer. <i>European Journal of Human Genetics</i> , 2009, 17, 629-635.	1.4	17
32	Clinical characteristics of tumors derived from colorectal cancer patients who harbor the Tumor Necrosis Factor \pm -1031T/T and NOD2 3020insC polymorphism. <i>Cancer Epidemiology</i> , 2009, 33, 161-163.	0.8	2
33	Frequency of mutations related to hereditary haemochromatosis in northwestern Poland. <i>Journal of Applied Genetics</i> , 2008, 49, 105-107.	1.0	10
34	AuroraA and Cyclin D1 polymorphisms and the age of onset of colorectal cancer in hereditary nonpolyposis colorectal cancer. <i>International Journal of Cancer</i> , 2008, 122, 1273-1277.	2.3	28
35	IGF1 is a modifier of disease risk in hereditary nonpolyposis colorectal cancer. <i>International Journal of Cancer</i> , 2008, 123, 1339-1343.	2.3	25
36	Inflammatory response gene polymorphisms and their relationship with colorectal cancer risk. <i>BMC Cancer</i> , 2008, 8, 112.	1.1	41

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37	The $\hat{A}^{\sim}149C>T$ SNP within the $\hat{I}^{\sim}DNMT3B$ gene, is not associated with early disease onset in hereditary non-polyposis colorectal cancer. <i>Cancer Letters</i> , 2008, 265, 39-44.	3.2	10
38	DNA and RNA analyses in detection of genetic predisposition to cancer. <i>Hereditary Cancer in Clinical Practice</i> , 2008, 6, 73.	0.6	0
39	MSH2 and MLH1 testing. <i>Hereditary Cancer in Clinical Practice</i> , 2008, 6, 83.	0.6	0
40	DNA testing for variants conferring low or moderate increase in the risk of cancer. <i>Hereditary Cancer in Clinical Practice</i> , 2008, 6, 84.	0.6	3
41	CARD15 variants in patients with sporadic Parkinson's disease. <i>Neuroscience Research</i> , 2007, 57, 473-476.	1.0	67
42	Germline CHEK2 mutations and colorectal cancer risk: different effects of a missense and truncating mutations?. <i>European Journal of Human Genetics</i> , 2007, 15, 237-241.	1.4	61
43	Nationwide study of clinical and molecular features of hereditary non-polyposis colorectal cancer (HNPCC) in Latvia. <i>Anticancer Research</i> , 2007, 27, 653-8.	0.5	4
44	Low-risk Genes and Multi-organ Cancer Risk in the Polish Population. <i>Hereditary Cancer in Clinical Practice</i> , 2006, 4, 52.	0.6	1
45	Some aspects of molecular diagnostics in Lynch syndrome. <i>Hereditary Cancer in Clinical Practice</i> , 2006, 4, 197.	0.6	1
46	Frequency and nature of hMSH6 germline mutations in Polish patients with colorectal, endometrial and ovarian cancers. <i>Clinical Genetics</i> , 2006, 70, 68-70.	1.0	6
47	Prevalence of the NOD2 3020insC mutation in aggregations of breast and lung cancer. <i>Breast Cancer Research and Treatment</i> , 2006, 95, 141-145.	1.1	31
48	CDKN2A common variant and multi-organ cancer risk – a population-based study. <i>International Journal of Cancer</i> , 2006, 118, 3180-3182.	2.3	26
49	MC1R common variants, CDKN2A and their association with melanoma and breast cancer risk. <i>International Journal of Cancer</i> , 2006, 119, 2597-2602.	2.3	38
50	Genetic Polymorphisms in Xenobiotic Clearance Genes and Their Influence on Disease Expression in Hereditary Nonpolyposis Colorectal Cancer Patients. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2006, 15, 2307-2310.	1.1	23
51	The 3020insC Allele of NOD2 Predisposes to Cancers of Multiple Organs. <i>Hereditary Cancer in Clinical Practice</i> , 2005, 3, 59.	0.6	26
52	NOD2 variants and the risk of malignant melanoma. <i>European Journal of Cancer Prevention</i> , 2005, 14, 143-146.	0.6	16
53	Germline MSH2 and MLH1 mutational spectrum including large rearrangements in HNPCC families from Poland (update study). <i>Clinical Genetics</i> , 2005, 69, 40-47.	1.0	34
54	Polymorphism in the P-glycoprotein drug transporter MDR1 gene in colon cancer patients. <i>European Journal of Clinical Pharmacology</i> , 2005, 61, 389-394.	0.8	79

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55	The 3020insC allele of NOD2 predisposes to early-onset breast cancer. <i>Breast Cancer Research and Treatment</i> , 2005, 89, 91-93.	1.1	47
56	CDKN2A common variants and their association with melanoma risk: a population-based study. <i>Cancer Research</i> , 2005, 65, 835-9.	0.4	43
57	Importance of microsatellite instability (MSI) in colorectal cancer: MSI as a diagnostic tool. <i>Annals of Oncology</i> , 2004, 15, iv283-iv284.	0.6	28
58	A Novel Founder CHEK2 Mutation is Associated with Increased Prostate Cancer Risk: Table 1. <i>Cancer Research</i> , 2004, 64, 2677-2679.	0.4	137
59	The NOD2 3020insC Mutation and the Risk of Colorectal Cancer: Table 1. <i>Cancer Research</i> , 2004, 64, 1604-1606.	0.4	105
60	Germline mutation and large deletion analysis of the CDKN2A and ARF genes in families with multiple melanoma or an aggregation of malignant melanoma and breast cancer. <i>International Journal of Cancer</i> , 2004, 110, 558-562.	2.3	24
61	CHEK2 Is a Multiorgan Cancer Susceptibility Gene. <i>American Journal of Human Genetics</i> , 2004, 75, 1131-1135.	2.6	426
62	Rarity of germline 1100delC mutation in CHK2 in patients with malignant melanoma of the skin. <i>Melanoma Research</i> , 2004, 14, 121-124.	0.6	8
63	Germline 657del5 mutation in the NBS1 gene in breast cancer patients. <i>International Journal of Cancer</i> , 2003, 106, 379-381.	2.3	80
64	Increased risk of breast cancer in relatives of malignant melanoma patients from families with strong cancer familial aggregation. <i>European Journal of Cancer Prevention</i> , 2003, 12, 241-245.	0.6	14
65	Germline 657del5 mutation in the NBS1 gene in patients with malignant melanoma of the skin. <i>Melanoma Research</i> , 2003, 13, 365-370.	0.6	30
66	Nuclear Pedigree Criteria of Suspected HNPCC. <i>Hereditary Cancer in Clinical Practice</i> , 2003, 1, 1.	0.6	8
67	Electro-Oculographic and Electroretinographic Studies in HNPCC Gene Mutation Carriers. <i>Ophthalmic Research</i> , 2003, 35, 281-294.	1.0	2
68	Germline MSH2 and MLH1 mutational spectrum in HNPCC families from Poland and the Baltic States. <i>Journal of Medical Genetics</i> , 2002, 39, 65e-65.	1.5	26
69	Germline mutations in the von Hippel-Lindau (VHL) gene in patients from Poland: disease presentation in patients with deletions of the entire VHL gene. <i>Journal of Medical Genetics</i> , 2002, 39, 38e-38.	1.5	47
70	Mutation analysis of MLH1 and MSH2 genes performed by denaturing high-performance liquid chromatography. <i>Journal of Proteomics</i> , 2002, 51, 89-100.	2.4	44
71	Ovarian cancer of endometrioid type as part of the MSH6 gene mutation phenotype. <i>Journal of Human Genetics</i> , 2002, 47, 0529-0531.	1.1	12
72	Molecular basis of inherited predispositions for tumors.. <i>Acta Biochimica Polonica</i> , 2002, 49, 571-581.	0.3	3

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73	Comparison of Alu-PCR, microsatellite instability, and immunohistochemical analyses in finding features characteristic for hereditary nonpolyposis colorectal cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2001, 127, 565-569.	1.2	8
74	Fluorescence In Situ Detection of Human Cutaneous Melanoma: Study of Diagnostic Parameters of the Method. <i>Journal of Investigative Dermatology</i> , 2001, 117, 1449-1451.	0.3	26
75	Optimization of experimental conditions for RNA-based sequencing of MLH1 and MSH2 genes. <i>Human Mutation</i> , 2001, 17, 52-60.	1.1	19
76	Value of pedigree/clinical data, immunohistochemistry and microsatellite instability analyses in reducing the cost of determining hMLH1 and hMSH2 gene mutations in patients with colorectal cancer. <i>European Journal of Cancer</i> , 2000, 36, 49-54.	1.3	99
77	Molecular Analyses in Diagnosis of High Genetic Predispositions to Malignancies.. <i>Journal of Clinical Biochemistry and Nutrition</i> , 2000, 28, 159-165.	0.6	0
78	Relationship between acetylation polymorphism and risk of atopic diseases. <i>Clinical Pharmacology and Therapeutics</i> , 1999, 65, 562-569.	2.3	29
79	Polymorphism of GSTM1 gene in patients with colorectal cancer and colonic polyps. <i>Experimental and Toxicologic Pathology</i> , 1999, 51, 321-325.	2.1	22
80	Frequency and nature of germline Rb-1 gene mutations in a series of patients with sporadic unilateral retinoblastoma. <i>European Journal of Cancer</i> , 1999, 35, 1824-1827.	1.3	12
81	Long polymerase chain reaction in detection of germline deletions in the von Hippel-Lindau tumour suppressor gene. <i>Human Genetics</i> , 1999, 105, 333-336.	1.8	18
82	Age at diagnosis to discriminate those patients for whom constitutional DNA sequencing is appropriate in sporadic unilateral retinoblastoma. <i>European Journal of Cancer</i> , 1998, 34, 1919-1921.	1.3	17
83	Leptin receptor isoforms expressed in human adipose tissue. <i>Metabolism: Clinical and Experimental</i> , 1998, 47, 844-847.	1.5	110
84	Hereditary breast cancer. <i>Polish Journal of Pathology</i> , 1998, 49, 59-66.	0.1	2
85	Losses at 3p common deletion sites in subtypes of kidney tumours: histopathological correlations. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 1996, 429, 37-42.	1.4	24
86	Mechanism of the Inhibitory Effect of Curdlan Sulfate on HIV-1 Infection in Vitro. <i>Virology</i> , 1994, 202, 735-745.	1.1	101
87	Analysis of CD4+ T cells in peripheral blood of children with perinatal human immunodeficiency virus (HIV) infection. <i>Journal of Clinical Immunology</i> , 1993, 13, 193-203.	2.0	20