

# Jan A Lubiński

## List of Publications by Year in descending order

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Version: 2024-02-01

535  
papers

42,316  
citations

2423

97  
h-index

3714

179  
g-index

556  
all docs

556  
docs citations

556  
times ranked

39874  
citing authors

#	ARTICLE	IF	CITATIONS
1	Polygenic risk modeling for prediction of epithelial ovarian cancer risk. <i>European Journal of Human Genetics</i> , 2022, 30, 349-362.	1.4	23
2	Contraceptive use and the risk of ovarian cancer among women with a BRCA1 or BRCA2 mutation. <i>Gynecologic Oncology</i> , 2022, 164, 514-521.	0.6	8
3	An Assessment of GPX1 (rs1050450), DIO2 (rs225014) and SEPP1 (rs7579) Gene Polymorphisms in Women with Endometrial Cancer. <i>Genes</i> , 2022, 13, 188.	1.0	5
4	Whole-Exome Sequencing Identifies a Novel Germline Variant in PTK7 Gene in Familial Colorectal Cancer. <i>International Journal of Molecular Sciences</i> , 2022, 23, 1295.	1.8	2
5	Germline Variants of CYBA and TRPM4 Predispose to Familial Colorectal Cancer. <i>Cancers</i> , 2022, 14, 670.	1.7	11
6	Germline HOXB13 mutation p.G84E do not confer an increased bladder or kidney cancer risk in polish population. <i>Hereditary Cancer in Clinical Practice</i> , 2022, 20, 1.	0.6	0
7	Pathology of Tumors Associated With Pathogenic Germline Variants in 9 Breast Cancer Susceptibility Genes. <i>JAMA Oncology</i> , 2022, 8, e216744.	3.4	51
8	Validated biomarker assays confirm that <i>ARID1A</i> loss is confounded with <i>MMR</i> deficiency, <i>CD8</i> <sup>+</sup> <i>TIL</i> infiltration, and provides no independent prognostic value in endometriosis-associated ovarian carcinomas. <i>Journal of Pathology</i> , 2022, 256, 388-401.	2.1	15
9	Risk of Second Primary Thyroid Cancer in Women with Breast Cancer. <i>Cancers</i> , 2022, 14, 957.	1.7	5
10	An Assessment of Serum Selenium Concentration in Women with Endometrial Cancer. <i>Nutrients</i> , 2022, 14, 958.	1.7	2
11	Spectrum and frequency of CHEK2 variants in breast cancer affected and general population in the Baltic states region, initial results and literature review. <i>European Journal of Medical Genetics</i> , 2022, 65, 104477.	0.7	5
12	Association of recurrent mutations in BRCA1, BRCA2, RAD51C, PALB2, and CHEK2 with the risk of borderline ovarian tumor. <i>Hereditary Cancer in Clinical Practice</i> , 2022, 20, 11.	0.6	4
13	The impact of oophorectomy on survival from breast cancer in patients with CHEK2 mutations. <i>British Journal of Cancer</i> , 2022, 127, 84-91.	2.9	4
14	Frequency of BRCA1 and BRCA2 mutations in ovarian cancer patients in South-East Poland. <i>Hereditary Cancer in Clinical Practice</i> , 2022, 20, 12.	0.6	0
15	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. <i>Nature</i> , 2022, 604, 502-508.	13.7	929
16	Germline BRCA1 and BRCA2 mutations and the risk of bladder or kidney cancer in Poland. <i>Hereditary Cancer in Clinical Practice</i> , 2022, 20, 13.	0.6	2
17	Common Variant in ALDH2 Modifies the Risk of Breast Cancer Among Carriers of the p.K3326* Variant in BRCA2. <i>JCO Precision Oncology</i> , 2022, 6, e2100450.	1.5	1
18	Germline MBD4 deficiency causes a multi-tumor predisposition syndrome. <i>American Journal of Human Genetics</i> , 2022, 109, 953-960.	2.6	23

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19	Bilateral Oophorectomy and the Risk of Breast Cancer in <i>BRCA1</i> Mutation Carriers: A Reappraisal. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2022, 31, 1351-1358.	1.1	3
20	Bladder cancer survival in patients with <i>NOD2</i> or <i>CDKN2A</i> variants. <i>Oncotarget</i> , 2022, 13, 628-640.	0.8	0
21	Whole exome sequencing identifies novel germline variants of <i>SLC15A4</i> gene as potentially cancer predisposing in familial colorectal cancer. <i>Molecular Genetics and Genomics</i> , 2022, , 1.	1.0	1
22	Breast cancer risks associated with missense variants in breast cancer susceptibility genes. <i>Genome Medicine</i> , 2022, 14, 51.	3.6	19
23	Population-based targeted sequencing of 54 candidate genes identifies <i>PALB2</i> as a susceptibility gene for high-grade serous ovarian cancer. <i>Journal of Medical Genetics</i> , 2021, 58, 305-313.	1.5	26
24	Lung Cancer Occurrence Correlation with Serum Chromium Levels and Genotypes. <i>Biological Trace Element Research</i> , 2021, 199, 1228-1236.	1.9	13
25	Blood cadmium levels as a marker for early lung cancer detection. <i>Journal of Trace Elements in Medicine and Biology</i> , 2021, 64, 126682.	1.5	28
26	Prevalence of germline <i>TP53</i> variants among early-onset breast cancer patients from Polish population. <i>Breast Cancer</i> , 2021, 28, 226-235.	1.3	10
27	<i>CYP3A7*1C</i> allele: linking premenopausal oestrone and progesterone levels with risk of hormone receptor-positive breast cancers. <i>British Journal of Cancer</i> , 2021, 124, 842-854.	2.9	5
28	Trans-ancestry genome-wide association meta-analysis of prostate cancer identifies new susceptibility loci and informs genetic risk prediction. <i>Nature Genetics</i> , 2021, 53, 65-75.	9.4	264
29	Recurrent <i>PALB2</i> mutations and the risk of cancers of bladder or kidney in Polish population. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 6.	0.6	1
30	Breast cancer risk after age 60 among <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2021, 187, 515-523.	1.1	5
31	Whole Exome Sequencing Identifies <i>APCDD1</i> and <i>HDAC5</i> Genes as Potentially Cancer Predisposing in Familial Colorectal Cancer. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1837.	1.8	6
32	Breast Cancer Risk Genes Association Analysis in More than 113,000 Women. <i>New England Journal of Medicine</i> , 2021, 384, 428-439.	13.9	532
33	Recurrent Mutations in <i>BRCA1</i> , <i>BRCA2</i> , <i>RAD51C</i> , <i>PALB2</i> and <i>CHEK2</i> in Polish Patients with Ovarian Cancer. <i>Cancers</i> , 2021, 13, 849.	1.7	13
34	Survival from breast cancer in women with a <i>BRCA2</i> mutation by treatment. <i>British Journal of Cancer</i> , 2021, 124, 1524-1532.	2.9	12
35	Proteogenomic insights into the biology and treatment of HPV-negative head and neck squamous cell carcinoma. <i>Cancer Cell</i> , 2021, 39, 361-379.e16.	7.7	189
36	Serum Selenium Level Predicts 10-Year Survival after Breast Cancer. <i>Nutrients</i> , 2021, 13, 953.	1.7	14

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37	A Novel Low-Risk Germline Variant in the SH2 Domain of the SRC Gene Affects Multiple Pathways in Familial Colorectal Cancer. <i>Journal of Personalized Medicine</i> , 2021, 11, 262.	1.1	0
38	PALB2 mutations and prostate cancer risk and survival. <i>British Journal of Cancer</i> , 2021, 125, 569-575.	2.9	18
39	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
40	A mosaic PIK3CA variant in a young adult with diffuse gastric cancer: case report. <i>European Journal of Human Genetics</i> , 2021, 29, 1354-1358.	1.4	9
41	The predictive ability of the 313 variant-based polygenic risk score for contralateral breast cancer risk prediction in women of European ancestry with a heterozygous BRCA1 or BRCA2 pathogenic variant. <i>Genetics in Medicine</i> , 2021, 23, 1726-1737.	1.1	16
42	Survival of Laryngeal Cancer Patients Depending on Zinc Serum Level and Oxidative Stress Genotypes. <i>Biomolecules</i> , 2021, 11, 865.	1.8	13
43	Abstract 878: Contraceptive use and ovarian cancer risk in BRCA1 and BRCA2 mutation carriers: A prospective cohort study. <i>Cancer Research</i> , 2021, 81, 878-878.	0.4	1
44	Functional annotation of the 2q35 breast cancer risk locus implicates a structural variant in influencing activity of a long-range enhancer element. <i>American Journal of Human Genetics</i> , 2021, 108, 1190-1203.	2.6	6
45	Blood Arsenic Levels as a Marker of Breast Cancer Risk among BRCA1 Carriers. <i>Cancers</i> , 2021, 13, 3345.	1.7	6
46	Abstract 857: Evaluating the relationship between arsenic exposure and cancer risk in Canada. , 2021, , .		0
47	Serum Selenium Level and 10-Year Survival after Melanoma. <i>Biomedicines</i> , 2021, 9, 991.	1.4	8
48	Low Blood-As Levels and Selected Genotypes Appears to Be Promising Biomarkers for Occurrence of Colorectal Cancer in Women. <i>Biomedicines</i> , 2021, 9, 1105.	1.4	0
49	Association of germline genetic variants with breast cancer-specific survival in patient subgroups defined by clinic-pathological variables related to tumor biology and type of systemic treatment. <i>Breast Cancer Research</i> , 2021, 23, 86.	2.2	7
50	Mendelian randomisation study of smoking exposure in relation to breast cancer risk. <i>British Journal of Cancer</i> , 2021, 125, 1135-1145.	2.9	9
51	Genetic predisposition to male breast cancer in Poland. <i>BMC Cancer</i> , 2021, 21, 975.	1.1	7
52	Weight Gain and the Risk of Ovarian Cancer in BRCA1 and BRCA2 Mutation Carriers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021, 30, 2038-2043.	1.1	6
53	Influence of the Levels of Arsenic, Cadmium, Mercury and Lead on Overall Survival in Lung Cancer. <i>Biomolecules</i> , 2021, 11, 1160.	1.8	23
54	Survival of bladder or renal cancer in patients with CHEK2 mutations. <i>PLoS ONE</i> , 2021, 16, e0257132.	1.1	1

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55	Breast Cancer Risk Factors and Survival by Tumor Subtype: Pooled Analyses from the Breast Cancer Association Consortium. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021, 30, 623-642.	1.1	19
56	A rare large duplication of MLH1 identified in Lynch syndrome. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 10.	0.6	2
57	BRCA1 and BRCA2 mutations in ovarian cancer patients from Belarus: update. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 13.	0.6	8
58	Germline variants and breast cancer survival in patients with distant metastases at primary breast cancer diagnosis. <i>Scientific Reports</i> , 2021, 11, 19787.	1.6	2
59	A prospective prostate cancer screening programme for men with pathogenic variants in mismatch repair genes (IMPACT): initial results from an international prospective study. <i>Lancet Oncology</i> , The, 2021, 22, 1618-1631.	5.1	48
60	Do BARD1 Mutations Confer an Elevated Risk of Prostate Cancer?. <i>Cancers</i> , 2021, 13, 5464.	1.7	1
61	Association of ABCA4 Gene Polymorphisms with Cleft Lip with or without Cleft Palate in the Polish Population. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 11483.	1.2	4
62	Blood Copper Levels and the Occurrence of Colorectal Cancer in Poland. <i>Biomedicines</i> , 2021, 9, 1628.	1.4	19
63	Blood arsenic levels and the risk of familial breast cancer in Poland. <i>International Journal of Cancer</i> , 2020, 146, 2721-2727.	2.3	18
64	BRCA1 promoter methylation in peripheral blood is associated with the risk of triple-negative breast cancer. <i>International Journal of Cancer</i> , 2020, 146, 1293-1298.	2.3	33
65	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. <i>Nature Genetics</i> , 2020, 52, 56-73.	9.4	120
66	Does preventive oophorectomy increase the risk of depression in BRCA mutation carriers?. <i>Menopause</i> , 2020, 27, 156-161.	0.8	5
67	Constitutional variants in POT1, TERF2IP, and ACD genes in patients with melanoma in the Polish population. <i>European Journal of Cancer Prevention</i> , 2020, 29, 511-519.	0.6	3
68	Cancer Predisposition Genes in Cancer-Free Families. <i>Cancers</i> , 2020, 12, 2770.	1.7	2
69	Breastfeeding and the risk of epithelial ovarian cancer among women with a BRCA1 or BRCA2 mutation. <i>Gynecologic Oncology</i> , 2020, 159, 820-826.	0.6	10
70	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. <i>Genetics in Medicine</i> , 2020, 22, 1653-1666.	1.1	82
71	Polymorphisms in MMP-1, MMP-2, MMP-7, MMP-13 and MT2A do not contribute to breast, lung and colon cancer risk in polish population. <i>Hereditary Cancer in Clinical Practice</i> , 2020, 18, 16.	0.6	14
72	Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. <i>American Journal of Human Genetics</i> , 2020, 107, 837-848.	2.6	39

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73	The Influence of Maternal BMI on Adverse Pregnancy Outcomes in Older Women. <i>Nutrients</i> , 2020, 12, 2838.	1.7	16
74	Prevalence of Recurrent Mutations Predisposing to Breast Cancer in Early-Onset Breast Cancer Patients from Poland. <i>Cancers</i> , 2020, 12, 2321.	1.7	11
75	Monoallelic NTHL1 Loss-of-Function Variants and Risk of Polyposis and Colorectal Cancer. <i>Gastroenterology</i> , 2020, 159, 2241-2243.e6.	0.6	20
76	Pre-Pregnancy Obesity vs. Other Risk Factors in Probability Models of Preeclampsia and Gestational Hypertension. <i>Nutrients</i> , 2020, 12, 2681.	1.7	21
77	Mutations in ATM , NBN and BRCA2 predispose to aggressive prostate cancer in Poland. <i>International Journal of Cancer</i> , 2020, 147, 2793-2800.	2.3	27
78	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. <i>Nature Genetics</i> , 2020, 52, 572-581.	9.4	265
79	Germline HOXB13 mutations p.G84E and p.R217C do not confer an increased breast cancer risk. <i>Scientific Reports</i> , 2020, 10, 9688.	1.6	2
80	Development and Validation of the Gene Expression Predictor of High-grade Serous Ovarian Carcinoma Molecular SubTYPE (PrOTYPE). <i>Clinical Cancer Research</i> , 2020, 26, 5411-5423.	3.2	43
81	Cancer prevention with aspirin in hereditary colorectal cancer (Lynch syndrome), 10-year follow-up and registry-based 20-year data in the CAPP2 study: a double-blind, randomised, placebo-controlled trial. <i>Lancet, The</i> , 2020, 395, 1855-1863.	6.3	220
82	Integrated proteogenomic deep sequencing and analytics accurately identify non-canonical peptides in tumor immunopeptidomes. <i>Nature Communications</i> , 2020, 11, 1293.	5.8	196
83	Identification of Familial Hodgkin Lymphoma Predisposing Genes Using Whole Genome Sequencing. <i>Frontiers in Bioengineering and Biotechnology</i> , 2020, 8, 179.	2.0	12
84	Transcriptome-wide association study of breast cancer risk by estrogen receptor status. <i>Genetic Epidemiology</i> , 2020, 44, 442-468.	0.6	32
85	A network analysis to identify mediators of germline-driven differences in breast cancer prognosis. <i>Nature Communications</i> , 2020, 11, 312.	5.8	30
86	Prediction of contralateral breast cancer: external validation of risk calculators in 20 international cohorts. <i>Breast Cancer Research and Treatment</i> , 2020, 181, 423-434.	1.1	14
87	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
88	First Trimester Microelements and Their Relationships with Pregnancy Outcomes and Complications. <i>Nutrients</i> , 2020, 12, 1108.	1.7	33
89	Serum Microelements in Early Pregnancy and their Risk of Large-for-Gestational Age Birth Weight. <i>Nutrients</i> , 2020, 12, 866.	1.7	7
90	Influence of the selenium level on overall survival in lung cancer. <i>Journal of Trace Elements in Medicine and Biology</i> , 2019, 56, 46-51.	1.5	25

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91	Allelic modification of breast cancer risk in women with an NBN mutation. <i>Breast Cancer Research and Treatment</i> , 2019, 178, 427-431.	1.1	6
92	Sex specific associations in genome wide association analysis of renal cell carcinoma. <i>European Journal of Human Genetics</i> , 2019, 27, 1589-1598.	1.4	27
93	First Trimester Serum Copper or Zinc Levels, and Risk of Pregnancy-Induced Hypertension. <i>Nutrients</i> , 2019, 11, 2479.	1.7	38
94	Inherited Variants in BLM and the Risk and Clinical Characteristics of Breast Cancer. <i>Cancers</i> , 2019, 11, 1548.	1.7	11
95	The FANCM:p.Arg658* truncating variant is associated with risk of triple-negative breast cancer. <i>Npj Breast Cancer</i> , 2019, 5, 38.	2.3	28
96	Integrated Proteogenomic Characterization of Clear Cell Renal Cell Carcinoma. <i>Cell</i> , 2019, 179, 964-983.e31.	13.5	430
97	Prevalence and spectrum of MLH1, MSH2, and MSH6 pathogenic germline variants in Pakistani colorectal cancer patients. <i>Hereditary Cancer in Clinical Practice</i> , 2019, 17, 29.	0.6	9
98	The Role of Early Pregnancy Maternal Selenium Levels on the Risk for Small-for-Gestational Age Newborns. <i>Nutrients</i> , 2019, 11, 2298.	1.7	24
99	Inherited variants in XRCC2 and the risk of breast cancer. <i>Breast Cancer Research and Treatment</i> , 2019, 178, 657-663.	1.1	13
100	Two truncating variants in FANCC and breast cancer risk. <i>Scientific Reports</i> , 2019, 9, 12524.	1.6	5
101	Interim Results from the IMPACT Study: Evidence for Prostate-specific Antigen Screening in BRCA2 Mutation Carriers. <i>European Urology</i> , 2019, 76, 831-842.	0.9	148
102	A comparison of ovarian cancer mortality in women with BRCA1 mutations undergoing annual ultrasound screening or preventive oophorectomy. <i>Gynecologic Oncology</i> , 2019, 155, 270-274.	0.6	15
103	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019, 10, 431.	5.8	88
104	Mendelian randomisation study of height and body mass index as modifiers of ovarian cancer risk in 22,588 BRCA1 and BRCA2 mutation carriers. <i>British Journal of Cancer</i> , 2019, 121, 180-192.	2.9	19
105	A combination of the immunohistochemical markers CK7 and SATB2 is highly sensitive and specific for distinguishing primary ovarian mucinous tumors from colorectal and appendiceal metastases. <i>Modern Pathology</i> , 2019, 32, 1834-1846.	2.9	54
106	The spectrum of mutations predisposing to familial breast cancer in Poland. <i>International Journal of Cancer</i> , 2019, 145, 3311-3320.	2.3	39
107	BARD1 is a Low/Moderate Breast Cancer Risk Gene: Evidence Based on an Association Study of the Central European p.Q564X Recurrent Mutation. <i>Cancers</i> , 2019, 11, 740.	1.7	25
108	Can Serum Iron Concentrations in Early Healthy Pregnancy Be Risk Marker of Pregnancy-Induced Hypertension?. <i>Nutrients</i> , 2019, 11, 1086.	1.7	22

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109	Serum Selenium Level in Early Healthy Pregnancy as a Risk Marker of Pregnancy Induced Hypertension. <i>Nutrients</i> , 2019, 11, 1028.	1.7	33
110	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. <i>Nature Communications</i> , 2019, 10, 1741.	5.8	90
111	Selected features of breast and peritoneal cancers diagnosed in BRCA1 carriers after risk-reducing salpingo-oophorectomy. <i>Hereditary Cancer in Clinical Practice</i> , 2019, 17, 10.	0.6	1
112	International trends in the uptake of cancer risk reduction strategies in women with a BRCA1 or BRCA2 mutation. <i>British Journal of Cancer</i> , 2019, 121, 15-21.	2.9	101
113	Genome-wide association study of germline variants and breast cancer-specific mortality. <i>British Journal of Cancer</i> , 2019, 120, 647-657.	2.9	52
114	Population-based identity-by-descent mapping combined with exome sequencing to detect rare risk variants for schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2019, 180, 223-231.	1.1	2
115	Mutational Signature Analysis Reveals NTHL1 Deficiency to Cause a Multi-tumor Phenotype. <i>Cancer Cell</i> , 2019, 35, 256-266.e5.	7.7	123
116	Oophorectomy and risk of contralateral breast cancer among BRCA1 and BRCA2 mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2019, 175, 443-449.	1.1	12
117	Serum selenium level and cancer risk: a nested case-control study. <i>Hereditary Cancer in Clinical Practice</i> , 2019, 17, 33.	0.6	15
118	Prediction and clinical utility of a contralateral breast cancer risk model. <i>Breast Cancer Research</i> , 2019, 21, 144.	2.2	24
119	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. <i>American Journal of Human Genetics</i> , 2019, 104, 21-34.	2.6	711
120	The influence of obesity-related factors in the etiology of renal cell carcinoma—A mendelian randomization study. <i>PLoS Medicine</i> , 2019, 16, e1002724.	3.9	59
121	The intron 3 16Âbp duplication polymorphism of p53 (rs17878362) is not associated with increased risk of developing triple-negative breast cancer. <i>Breast Cancer Research and Treatment</i> , 2019, 173, 727-733.	1.1	5
122	Iron levels, genes involved in iron metabolism and antioxidative processes and lung cancer incidence. <i>PLoS ONE</i> , 2019, 14, e0208610.	1.1	41
123	Age-specific risks of incident, contralateral and ipsilateral breast cancer among 1776 Polish BRCA1 mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2019, 174, 769-774.	1.1	7
124	Height and Body Mass Index as Modifiers of Breast Cancer Risk in BRCA1 Mutation Carriers: A Mendelian Randomization Study. <i>Journal of the National Cancer Institute</i> , 2019, 111, 350-364.	3.0	30
125	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. <i>International Journal of Epidemiology</i> , 2019, 48, 795-806.	0.9	81
126	Founder Mutations for Early Onset Melanoma as Revealed by Whole Exome Sequencing Suggests That This is Not Associated with the Increasing Incidence of Melanoma in Poland. <i>Cancer Research and Treatment</i> , 2019, 51, 337-344.	1.3	8

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127	Inherited NBN Mutations and Prostate Cancer Risk and Survival. <i>Cancer Research and Treatment</i> , 2019, 51, 1180-1187.	1.3	21
128	Abstract 4172: Identification of familial Hodgkin lymphoma predisposing genes by whole genome sequencing. , 2019, , .		0
129	The <i>BRCA2</i> c.68-7T>A variant is not pathogenic: A model for clinical calibration of spliceogenicity. <i>Human Mutation</i> , 2018, 39, 729-741.	1.1	19
130	MyD88 and TLR4 Expression in Epithelial Ovarian Cancer. <i>Mayo Clinic Proceedings</i> , 2018, 93, 307-320.	1.4	22
131	Hormone Replacement Therapy After Oophorectomy and Breast Cancer Risk Among <i>BRCA1</i> Mutation Carriers. <i>JAMA Oncology</i> , 2018, 4, 1059.	3.4	121
132	Adult height is associated with increased risk of ovarian cancer: a Mendelian randomisation study. <i>British Journal of Cancer</i> , 2018, 118, 1123-1129.	2.9	15
133	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> or <i>BRCA2</i> mutations. <i>Human Mutation</i> , 2018, 39, 593-620.	1.1	224
134	Physical activity during adolescence and young adulthood and the risk of breast cancer in <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2018, 169, 561-571.	1.1	25
135	Predictors of survival for breast cancer patients with a <i>BRCA1</i> mutation. <i>Breast Cancer Research and Treatment</i> , 2018, 168, 513-521.	1.1	20
136	The association between smoking and cancer incidence in <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>International Journal of Cancer</i> , 2018, 142, 2263-2272.	2.3	20
137	Role of germline aberrations affecting <i>CTNNA1</i> , <i>MAP3K6</i> and <i>MYD88</i> in gastric cancer susceptibility. <i>Journal of Medical Genetics</i> , 2018, 55, 669-674.	1.5	37
138	Prostate-specific antigen velocity in a prospective prostate cancer screening study of men with genetic predisposition. <i>British Journal of Cancer</i> , 2018, 118, 266-276.	2.9	12
139	Frequency of <i>BRCA1</i> and <i>BRCA2</i> causative founder variants in ovarian cancer patients in South-East Poland. <i>Hereditary Cancer in Clinical Practice</i> , 2018, 16, 6.	0.6	12
140	Assessment of moderate coffee consumption and risk of epithelial ovarian cancer: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2018, 47, 450-459.	0.9	15
141	Prospective evaluation of body size and breast cancer risk among <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>International Journal of Epidemiology</i> , 2018, 47, 987-997.	0.9	11
142	Serum selenium levels are associated with age-related cataract. <i>Annals of Agricultural and Environmental Medicine</i> , 2018, 25, 443-448.	0.5	12
143	<i>BRCA1/2</i> mutations are not a common cause of malignant melanoma in the Polish population. <i>PLoS ONE</i> , 2018, 13, e0204768.	1.1	6
144	Reply to "Mutations in <i>RECQL</i> are not associated with breast cancer risk in an Australian population"™. <i>Nature Genetics</i> , 2018, 50, 1348-1349.	9.4	2

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145	Genetic polymorphisms may influence the vertical growth rate of melanoma. <i>Journal of Cancer</i> , 2018, 9, 3078-3083.	1.2	2
146	Age-specific ovarian cancer risks among women with a BRCA1 or BRCA2 mutation. <i>Gynecologic Oncology</i> , 2018, 150, 85-91.	0.6	65
147	Age at first full-term birth and breast cancer risk in BRCA1 and BRCA2 mutation carriers. <i>Breast Cancer Research and Treatment</i> , 2018, 171, 421-426.	1.1	10
148	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. <i>American Journal of Human Genetics</i> , 2018, 102, 1185-1194.	2.6	119
149	Association of p16 expression with prognosis varies across ovarian carcinoma histotypes: an Ovarian Tumor Tissue Analysis consortium study. <i>Journal of Pathology: Clinical Research</i> , 2018, 4, 250-261.	1.3	70
150	Association of zinc level and polymorphism in MMP-7 gene with prostate cancer in Polish population. <i>PLoS ONE</i> , 2018, 13, e0201065.	1.1	30
151	Variants in genes encoding small GTPases and association with epithelial ovarian cancer susceptibility. <i>PLoS ONE</i> , 2018, 13, e0197561.	1.1	9
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