Janusz Limon

List of Publications by Year
in descending order

101 papers	3,586 citations	126907 33 h-index	144013 57 g-index
104 all docs	104 docs citations	104 times ranked	4089 citing authors

#	Article	IF	Citations
1	Comparative Genomic Hybridization to Microarrays in Fetuses with High-Risk Prenatal Indications: Polish Experience with 7400 Pregnancies. Genes, 2022, 13, 690.	2.4	3
2	Albert de la Chapelleâ€"pro memoriam. Journal of Applied Genetics, 2021, 62, 455-458.	1.9	0
3	Monitoring the Effects of Hypolipidemic Treatment in Children with Familial Hypercholesterolemia in Poland. Life, 2020, 10, 270.	2.4	1
4	Association of Genes Related to Oxidative Stress with the Extent of Coronary Atherosclerosis. Life, 2020, 10, 210.	2.4	3
5	NADPH Oxidase Gene Polymorphism is Associated with Mortality and Cardiovascular Events in 7-Year Follow-Up. Journal of Clinical Medicine, 2020, 9, 1475.	2.4	7
6	Cell-free DNA BRAF V600E measurements during BRAF inhibitor therapy of metastatic melanoma: long-term analysis. Tumori, 2020, 106, 241-248.	1.1	13
7	Bayesian multilevel model of micro RNA levels in ovarian-cancer and healthy subjects. PLoS ONE, 2019, 14, e0221764.	2.5	7
8	Folate/homocysteine metabolism and lung cancer risk among smokers. PLoS ONE, 2019, 14, e0214462.	2.5	18
9	Spectrum and Prevalence of Pathogenic Variants in Ovarian Cancer Susceptibility Genes in a Group of 333 Patients. Cancers, 2018, 10, 442.	3.7	30
10	Mitochondrial DNA levels in Huntington disease leukocytes and dermal fibroblasts. Metabolic Brain Disease, 2017, 32, 1237-1247.	2.9	19
11	Application of high-resolution genomic profiling in the differential diagnosis of liposarcoma. Molecular Cytogenetics, 2017, 10, 7.	0.9	6
12	Coincidence of PTPN22 c.1858CC and FCRL3 -169CC genotypes as a biomarker of preserved residual Î²-cell function in children with type 1 diabetes. Pediatric Diabetes, 2017, 18, 696-705.	2.9	7
13	Genomic findings in patients with clinical suspicion of 22q11.2 deletion syndrome. Journal of Applied Genetics, 2017, 58, 93-98.	1.9	21
14	Detection of BRCA1/2 mutations in circulating tumor DNA from patients with ovarian cancer. Oncotarget, 2017, 8, 101325-101332.	1.8	32
15	An open label phase II study evaluating first-line EGFR tyrosine kinase inhibitor erlotinib in non-small cell lung cancer patients with tumors showing high EGFR gene copy number. Oncotarget, 2017, 8, 17270-17278.	1.8	3
16	Antagonizing functions of BARD1 and its alternatively spliced variant BARD1Î´ in telomere stability. Oncotarget, 2017, 8, 9339-9353.	1.8	9
17	When do paediatric patients with familial hypercholesterolemia need statin therapy?. Medycyna Wieku Rozwojowego, 2017, 21, 43-50.	0.2	0
18	Chromosome 18q deletion syndrome with autoimmune diabetes mellitus: putative genomic loci for autoimmunity and immunodeficiency. Pediatric Diabetes, 2016, 17, 153-159.	2.9	7

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19	B26â€...Differential mitochondrial DNA levels in HD patients depending on the cell type. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, A18.1-A18.	1.9	0
20	Efficacy of clinical diagnostic criteria for familial hypercholesterolemia genetic testing in Poland. Atherosclerosis, 2016, 249, 52-58.	0.8	19
21	The algorithm for Alzheimer risk assessment based on APOE promoter polymorphisms. Alzheimer's Research and Therapy, 2016, 8, 19.	6.2	17
22	Detection of somatic BRCA 1/2 mutations in ovarian cancer â€“ nextâ€generation sequencing analysis of 100 cases. Cancer Medicine, 2016, 5, 1640-1646.	2.8	36
23	A simple modification to improve the accuracy of methylation-sensitive restriction enzyme quantitative polymerase chain reaction. Analytical Biochemistry, 2016, 500, 88-90.	2.4	5
24	Co-incidence of Turner syndrome and Duchenne muscular dystrophy - an important problem for the clinician. Medycyna Wieku Rozwojowego, 2016, 20, 273-278.	0.2	2
25	A novel splicing mutation in the SLC9A3R1 gene in tumors from ovarian cancer patients. Oncology Letters, 2015, 10, 3722-3726.	1.8	7
26	Concurrent DNA Copy-Number Alterations and Mutations in Genes Related to Maintenance of Genome Stability in Uninvolved Mammary Glandular Tissue from Breast Cancer Patients. Human Mutation, 2015, 36, 1088-1099.	2.5	11
27	Cancer predisposing BARD1 mutations affect exon skipping and are associated with overexpression of specific BARD1 isoforms. Oncology Reports, 2015, 34, 2609-2617.	2.6	18
28	Mutational analysis of BRCA1/2 in a group of 134 consecutive ovarian cancer patients. Novel and recurrent BRCA1/2 alterations detected by next generation sequencing. Journal of Applied Genetics, 2015, 56, 193-198.	1.9	19
29	Analysis of large mutations in BARD1 in patients with breast and/or ovarian cancer: the Polish population as an example. Scientific Reports, 2015, 5, 10424.	3.3	28
30	Prevalence of the BLM nonsense mutation, p.Q548X, in ovarian cancer patients from Central and Eastern Europe. Familial Cancer, 2015, 14, 145-149.	1.9	12
31	The Outcome of Targeted Therapy in Advanced Gastrointestinal Stromal Tumors (Gist) with Non-Exon 11 Kit Mutations. Polski Przeglad Chirurgiczny, 2014, 86, 325-32.	0.4	2
32	Tumor Genotype Is an Independent Prognostic Factor in Primary Gastrointestinal Stromal Tumors of Gastric Origin: A European Multicenter Analysis Based on ConticaGIST. Clinical Cancer Research, 2014, 20, 6105-6116.	7.0	129
33	Management of familial hypercholesterolemia in children and adolescents. Position paper of the Polish Lipid Expert Forum. Journal of Clinical Lipidology, 2014, 8, 173-180.	1.5	30
34	Rare cancers. International Journal of Biochemistry and Cell Biology, 2014, 53, 461.	2.8	0
35	Mutational analysis in podocin-associated hereditary nephrotic syndrome in Polish patients: founder effect in the Kashubian population. Journal of Applied Genetics, 2013, 54, 327-333.	1.9	17
36	What are the current outcomes of advanced gastrointestinal stromal tumors: who are the long-term survivors treated initially with imatinib?. Medical Oncology, 2013, 30, 765.	2.5	20

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37	Spectrum of NIPBL gene mutations in Polish patients with Cornelia de Lange syndrome. Journal of Applied Genetics, 2013, 54, 27-33.	1.9	11
38	Stanowisko dotyczÄ…ce postÄ™powania w rodzinnej hipercholesterolemii u dzieci i mÅ‚odzieÅ¼y. Stanowisko Forum EkspertÃ³w Lipidowych. Pediatria Polska, 2013, 88, 567-574.	0.2	1
39	Management of familial heterozygous hypercholesterolemia: Position Paper of the Polish Lipid Expert Forum. Journal of Clinical Lipidology, 2013, 7, 217-221.	1.5	28
40	Isolated NIBPL missense mutations that cause Cornelia de Lange syndrome alter MAU2 interaction. European Journal of Human Genetics, 2012, 20, 271-276.	2.8	24
41	Limited significance of family history for presence of BRCA1 gene mutation in Polish breast and ovarian cancer cases. Familial Cancer, 2012, 11, 351-354.	1.9	20
42	Cornelia de Lange syndrome with NIPBL mutation and mosaic Turner syndrome in the same individual. BMC Medical Genetics, 2012, 13, 43.	2.1	12
43	The outcome and predictive factors of sunitinib therapy in advanced gastrointestinal stromal tumors (GIST) after imatinib failure - one institution study. BMC Cancer, 2012, 12, 107.	2.6	62
44	Angiotensin converting enzyme gene polymorphism is associated with severity of coronary artery disease in men with high total cholesterol levels. Journal of Applied Genetics, 2012, 53, 175-182.	1.9	22
45	Cancer predisposing BARD1 mutations in breastâ€“ovarian cancer families. Breast Cancer Research and Treatment, 2012, 131, 89-97.	2.5	88
46	A case of mast cell leukaemia with exon 9 KIT mutation and good response to imatinib. European Journal of Haematology, 2011, 86, 531-535.	2.2	46
47	Novel, activating KIT-N822I mutation in familial cutaneous mastocytosis. Experimental Hematology, 2011, 39, 859-865.e2.	0.4	42
48	Prevalence of the most frequent BRCA1 mutations in Polish population. Journal of Applied Genetics, 2011, 52, 325-330.	1.9	38
49	Clinical utility of the new American Joint Committee on Cancer staging system for gastrointestinal stromal tumors. Cancer, 2011, 117, 4916-4924.	4.1	47
50	Cornelia de Lange syndrome associated with a de-novo novel NIPBL splice-site mutation and a coincidental inherited translocation t(3;5)(p13;q11). Clinical Dysmorphology, 2011, 20, 222-224.	0.3	1
51	Cornelia de Lange syndrome case due to genomic rearrangements including NIPBL. European Journal of Medical Genetics, 2010, 53, 378-382.	1.3	18
52	Activity of Dasatinib Against Novel KIT-N822I Mutation In Familial Cutaneous Mastocytosis. Blood, 2010, 116, 4088-4088.	1.4	0
53	c.1810C>T Polymorphism of NTRK1Gene is associated with reduced Survival in Neuroblastoma Patients. BMC Cancer, 2009, 9, 436.	2.6	7
54	High frequency of BRCA1/2 germline mutations in consecutive ovarian cancer patients in Poland. Gynecologic Oncology, 2008, 108, 433-437.	1.4	50

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55	BRCA1 and BRCA2 point mutations and large rearrangements in breast and ovarian cancer families in Northern Poland. Oncology Reports, 2008, , .	2.6	21
56	BRCA1 and BRCA2 point mutations and large rearrangements in breast and ovarian cancer families in Northern Poland. Oncology Reports, 2008, 19, 263-8.	2.6	51
57	Risk Criteria and Prognostic Factors for Predicting Recurrences After Resection of Primary Gastrointestinal Stromal Tumor. Annals of Surgical Oncology, 2007, 14, 2018-2027.	1.5	227
58	Predictive factors for long-term effects of imatinib therapy in patients with inoperable/metastatic CD117(+) gastrointestinal stromal tumors (GISTs). Journal of Cancer Research and Clinical Oncology, 2007, 133, 589-597.	2.5	31
59	Mutational and genotypeâ€“phenotype correlation analyses in 28 Polish patients with Cornelia de Lange syndrome. American Journal of Medical Genetics, Part A, 2006, 140A, 1531-1541.	1.2	55
60	C-344T polymorphism of the aldosterone synthase gene and blood pressure in the elderly: a population-based study. Journal of Hypertension, 2005, 23, 1991-1996.	0.5	44
61	Microdeletion and IGF2 loss of imprinting in a cascade causing Beckwith-Wiedemann syndrome with Wilms' tumor. Nature Genetics, 2005, 37, 785-786.	21.4	40
62	Cytogenetic and molecular findings in 75 clear cell renal cell carcinomas. Oncology Reports, 2005, 13, 949.	2.6	9
63	Microdeletion of target sites for insulator protein CTCF in a chromosome 11p15 imprinting center in Beckwith-Wiedemann syndrome and Wilms' tumor. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 4085-4090.	7.1	124
64	Prevalence and clinical correlations of BRCA1/BRCA2 unclassified variant carriers among unselected primary ovarian cancer cases â€“ preliminary report. European Journal of Cancer, 2005, 41, 143-150.	2.8	22
65	The magic human 46 chromosomes were immortalised on a bronze plaque at Lund University in Sweden. Journal of Applied Genetics, 2004, 45, 1-2.	1.9	16
66	Title is missing!. Journal of Neuro-Oncology, 2003, 64, 284-284.	2.9	0
67	BRCA1 and BRCA2 mutation analysis in breast-ovarian cancer families from northeastern Poland. Human Mutation, 2003, 21, 553-554.	2.5	39
68	Association between the PlA platelet glycoprotein GPIIIa polymorphism and extent of coronary artery disease. International Journal of Cardiology, 2003, 88, 229-237.	1.7	15
69	Calretinin and Other Mesothelioma Markers in Synovial Sarcoma. American Journal of Surgical Pathology, 2001, 25, 610-617.	3.7	147
70	Spectral karyotyping reveals 17;22 fusions in a cytogenetically atypical dermatofibrosarcoma protuberans with a large marker chromosome as a sole abnormality. Genes Chromosomes and Cancer, 2001, 31, 182-186.	2.8	10
71	Clinical impact of molecular and cytogenetic findings in synovial sarcoma. Genes Chromosomes and Cancer, 2001, 31, 362-372.	2.8	108
72	Cytogenetics of hepatoblastoma: Further characterization of 1q rearrangements by fluorescence in situ hybridization: An international collaborative study. Medical and Pediatric Oncology, 2000, 34, 165-170.	1.0	41

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73	Founder BRCA1 mutations and two novel germline BRCA2 mutations in breast and/or ovarian cancer families from North-Eastern Poland. Human Mutation, 2000, 15, 480-481.	2.5	33
74	Patterns of keratin polypeptides in 110 biphasic, monophasic, and poorly differentiated synovial sarcomas. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2000, 437, 275-283.	2.8	74
75	Recurrent Deletion of the Region Encoding Two (Gly-X-Y) Repeats in Patients with Anhidrotic Ectodermal Dysplasia Indicates Important Role for Collagen-Like Domain of the EDA Gene Productâ€"Ectodysplasin-A. Fetal and Pediatric Pathology, 2000, 19, 425-432.	0.3	1
76	Founder BRCA1 mutations and two novel germline BRCA2 mutations in breast and/or ovarian cancer families from North-Eastern Poland. , 2000, 15, 480.		1
77	Rearrangement of the neoplasia-associated geneHMGIC in synovia from patients with osteoarthritis. , 1999, 24, 278-282.		12
78	Nonrandom chromosomal aberrations and cytogenetic heterogeneity in gallbladder carcinomas. , 1999, 26, 312-321.		17
79	Translocation (X;1)(p11.2;q21) in a Papillary Renal Cell Carcinoma in a 14-Year-Old Girl. Cancer Genetics and Cytogenetics, 1998, 101, 159-161.	1.0	21
80	Cytogenetic Findings in an Embryonal Sarcoma of the Liver. Cancer Genetics and Cytogenetics, 1998, 102, 142-144.	1.0	18
81	Recurrent chromosome changes in two adult fibrosarcomas. , 1998, 21, 119-123.		18
82	Clonal chromosome aberrations are present in vivo in synovia and osteophytes from patients with osteoarthritis. Human Genetics, 1997, 101, 295-298.	3.8	16
83	Evidence of somatic mutations in osteoarthritis. Human Genetics, 1996, 98, 651-656.	3.8	30
84	Clear cell sarcoma of tendons and aponeuroses with t(12;22) (q13;q12) diagnosed initially as malignant melanoma. Cancer Genetics and Cytogenetics, 1996, 91, 37-39.	1.0	18
85	Cytogenetic and Immunohistochemical Profile of Myxoid Liposarcoma. American Journal of Clinical Pathology, 1995, 103, 20-26.	0.7	53
86	Chromosome abnormalities in bilateral breast carcinomas. Cytogenetic evaluation of the clonal origin of multiple primary tumors. Cancer, 1995, 76, 250-258.	4.1	76
87	Cytogenetic findings in malignant peripheral nerve sheath tumors. International Journal of Cancer, 1995, 61, 793-798.	5.1	75
88	Abnormal karyotypes in three carcinomas of the gallbladder. Cancer Genetics and Cytogenetics, 1994, 76, 15-18.	1.0	9
89	Interstitial deletion of the short arm of chromosome 3 as a primary chromosome abnormality in carcinomas of the breast. Genes Chromosomes and Cancer, 1993, 6, 151-155.	2.8	62
90	Trisomy 12 and 4 in a thecoma of the ovary. Gynecologic Oncology, 1992, 45, 66-68.	1.4	16

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91	Cytogenetic analysis of 57 primary prostatic adenocarcinomas. Genes Chromosomes and Cancer, 1992, 4, 16-24.	2.8	155
92	Improved technique for short-term culture and cytogenetic analysis of human breast cancer. Genes Chromosomes and Cancer, 1992, 5, 14-20.	2.8	94
93	AgNOR staining in benign hyperplasia and carcinoma of the prostate. Prostate, 1991, 18, 155-162.	2.3	15
94	Trisomy of chromosome 12 in a case of thecoma of the ovary. Gynecologic Oncology, 1990, 36, 413-416.	1.4	36
95	An improved technique for short-term culturing of human prostatic adenocarcinoma tissue for cytogenetic analysis. Cancer Genetics and Cytogenetics, 1990, 46, 191-199.	1.0	26
96	Double minutes in two primary adenocarcinomas of the prostate. Cancer Genetics and Cytogenetics, 1989, 39, 191-194.	1.0	31
97	Cytogenetic studies of adipose tissue tumors. II. Recurrent reciprocal translocation t(12;16)(q13;p11) in myxoid liposarcomas. Cancer Genetics and Cytogenetics, 1986, 23, 291-299.	1.0	235
98	Chromosome abnormalities in two benign adipose tumors. Cancer Genetics and Cytogenetics, 1986, 22, 55-61.	1.0	66
99	Recurrent chromosome translocations in liposarcoma. Cancer Genetics and Cytogenetics, 1986, 22, 93-94.	1.0	57
100	Translocations involving the X chromosome in solid tumors: Presentation of two sarcomas with t(X;18)(q13;p11). Cancer Genetics and Cytogenetics, 1986, 23, 87-91.	1.0	98
101	Translocation X;18 in synovial sarcoma. Cancer Genetics and Cytogenetics, 1986, 23, 93.	1.0	143