

Janusz Limon

List of Publications by Year in descending order

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Version: 2024-02-01

101
papers

3,586
citations

136885

32
h-index

149623

56
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. <i>Genes</i> , 2022, 13, 690. | 1.0 | 3 |
| 2 | Albert de la Chapelle – pro memoriam. <i>Journal of Applied Genetics</i> , 2021, 62, 455-458. | 1.0 | 0 |
| 3 | Monitoring the Effects of Hypolipidemic Treatment in Children with Familial Hypercholesterolemia in Poland. <i>Life</i> , 2020, 10, 270. | 1.1 | 1 |
| 4 | Association of Genes Related to Oxidative Stress with the Extent of Coronary Atherosclerosis. <i>Life</i> , 2020, 10, 210. | 1.1 | 3 |
| 5 | NADPH Oxidase Gene Polymorphism is Associated with Mortality and Cardiovascular Events in 7-Year Follow-Up. <i>Journal of Clinical Medicine</i> , 2020, 9, 1475. | 1.0 | 7 |
| 6 | Cell-free DNA BRAF V600E measurements during BRAF inhibitor therapy of metastatic melanoma: long-term analysis. <i>Tumori</i> , 2020, 106, 241-248. | 0.6 | 13 |
| 7 | Bayesian multilevel model of micro RNA levels in ovarian-cancer and healthy subjects. <i>PLoS ONE</i> , 2019, 14, e0221764. | 1.1 | 7 |
| 8 | Folate/homocysteine metabolism and lung cancer risk among smokers. <i>PLoS ONE</i> , 2019, 14, e0214462. | 1.1 | 18 |
| 9 | Spectrum and Prevalence of Pathogenic Variants in Ovarian Cancer Susceptibility Genes in a Group of 333 Patients. <i>Cancers</i> , 2018, 10, 442. | 1.7 | 30 |
| 10 | Mitochondrial DNA levels in Huntington disease leukocytes and dermal fibroblasts. <i>Metabolic Brain Disease</i> , 2017, 32, 1237-1247. | 1.4 | 19 |
| 11 | Application of high-resolution genomic profiling in the differential diagnosis of liposarcoma. <i>Molecular Cytogenetics</i> , 2017, 10, 7. | 0.4 | 6 |
| 12 | Coincidence of <i>PTPN22</i> c.1858CC and <i>FCRL3</i> -169CC genotypes as a biomarker of preserved residual β 2-cell function in children with type 1 diabetes. <i>Pediatric Diabetes</i> , 2017, 18, 696-705. | 1.2 | 7 |
| 13 | Genomic findings in patients with clinical suspicion of 22q11.2 deletion syndrome. <i>Journal of Applied Genetics</i> , 2017, 58, 93-98. | 1.0 | 21 |
| 14 | Detection of <i>BRCA1/2</i> mutations in circulating tumor DNA from patients with ovarian cancer. <i>Oncotarget</i> , 2017, 8, 101325-101332. | 0.8 | 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. <i>Oncotarget</i> , 2017, 8, 17270-17278. | 0.8 | 3 |
| 16 | Antagonizing functions of BARD1 and its alternatively spliced variant BARD1 Δ in telomere stability. <i>Oncotarget</i> , 2017, 8, 9339-9353. | 0.8 | 9 |
| 17 | When do paediatric patients with familial hypercholesterolemia need statin therapy?. <i>Medycyna Wieku Rozwojowego</i> , 2017, 21, 43-50. | 0.2 | 0 |
| 18 | Chromosome 18q deletion syndrome with autoimmune diabetes mellitus: putative genomic loci for autoimmunity and immunodeficiency. <i>Pediatric Diabetes</i> , 2016, 17, 153-159. | 1.2 | 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. | 0.9 | 0 |
| 20 | Efficacy of clinical diagnostic criteria for familial hypercholesterolemia genetic testing in Poland. Atherosclerosis, 2016, 249, 52-58. | 0.4 | 19 |
| 21 | The algorithm for Alzheimer risk assessment based on APOE promoter polymorphisms. Alzheimer's Research and Therapy, 2016, 8, 19. | 3.0 | 17 |
| 22 | Detection of somatic BRCA 1/2 mutations in ovarian cancer â€œ nextâ€œgeneration sequencing analysis of 100 cases. Cancer Medicine, 2016, 5, 1640-1646. | 1.3 | 36 |
| 23 | A simple modification to improve the accuracy of methylation-sensitive restriction enzyme quantitative polymerase chain reaction. Analytical Biochemistry, 2016, 500, 88-90. | 1.1 | 5 |
| 24 | Co-occurrence 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. | 0.8 | 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. | 1.1 | 11 |
| 27 | Cancer predisposing BARD1 mutations affect exon skipping and are associated with overexpression of specific BARD1 isoforms. Oncology Reports, 2015, 34, 2609-2617. | 1.2 | 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. | 1.0 | 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. | 1.6 | 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. | 0.9 | 12 |
| 31 | The Outcome of Targeted Therapy in Advanced Gastrointestinal Stromal Tumors (Gist) with Non-Exon 11 Kit Mutations. Polski Przegląd Chirurgiczny, 2014, 86, 325-32. | 0.2 | 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. | 3.2 | 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. | 0.6 | 30 |
| 34 | Rare cancers. International Journal of Biochemistry and Cell Biology, 2014, 53, 461. | 1.2 | 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. | 1.0 | 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. | 1.2 | 20 |

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|----|---|-----|-----------|
| 37 | Spectrum of NIPBL gene mutations in Polish patients with Cornelia de Lange syndrome. <i>Journal of Applied Genetics</i> , 2013, 54, 27-33. | 1.0 | 11 |
| 38 | Stanowisko dotyczące postępowania w rodzinnej hipercholesterolemii u dzieci i młodzieży. Stanowisko Forum Ekspertów w Lipidowych. <i>Pediatrics Polska</i> , 2013, 88, 567-574. | 0.1 | 1 |
| 39 | Management of familial heterozygous hypercholesterolemia: Position Paper of the Polish Lipid Expert Forum. <i>Journal of Clinical Lipidology</i> , 2013, 7, 217-221. | 0.6 | 28 |
| 40 | Isolated NIPBL missense mutations that cause Cornelia de Lange syndrome alter MAU2 interaction. <i>European Journal of Human Genetics</i> , 2012, 20, 271-276. | 1.4 | 24 |
| 41 | Limited significance of family history for presence of BRCA1 gene mutation in Polish breast and ovarian cancer cases. <i>Familial Cancer</i> , 2012, 11, 351-354. | 0.9 | 20 |
| 42 | Cornelia de Lange syndrome with NIPBL mutation and mosaic Turner syndrome in the same individual. <i>BMC Medical Genetics</i> , 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. <i>BMC Cancer</i> , 2012, 12, 107. | 1.1 | 62 |
| 44 | Angiotensin converting enzyme gene polymorphism is associated with severity of coronary artery disease in men with high total cholesterol levels. <i>Journal of Applied Genetics</i> , 2012, 53, 175-182. | 1.0 | 22 |
| 45 | Cancer predisposing BARD1 mutations in breast-ovarian cancer families. <i>Breast Cancer Research and Treatment</i> , 2012, 131, 89-97. | 1.1 | 88 |
| 46 | A case of mast cell leukaemia with exon 9 KIT mutation and good response to imatinib. <i>European Journal of Haematology</i> , 2011, 86, 531-535. | 1.1 | 46 |
| 47 | Novel, activating KIT-N822I mutation in familial cutaneous mastocytosis. <i>Experimental Hematology</i> , 2011, 39, 859-865.e2. | 0.2 | 42 |
| 48 | Prevalence of the most frequent BRCA1 mutations in Polish population. <i>Journal of Applied Genetics</i> , 2011, 52, 325-330. | 1.0 | 38 |
| 49 | Clinical utility of the new American Joint Committee on Cancer staging system for gastrointestinal stromal tumors. <i>Cancer</i> , 2011, 117, 4916-4924. | 2.0 | 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). <i>Clinical Dysmorphology</i> , 2011, 20, 222-224. | 0.1 | 1 |
| 51 | Cornelia de Lange syndrome case due to genomic rearrangements including NIPBL. <i>European Journal of Medical Genetics</i> , 2010, 53, 378-382. | 0.7 | 18 |
| 52 | Activity of Dasatinib Against Novel KIT-N822I Mutation In Familial Cutaneous Mastocytosis. <i>Blood</i> , 2010, 116, 4088-4088. | 0.6 | 0 |
| 53 | c.1810C>T Polymorphism of NTRK1 Gene is associated with reduced Survival in Neuroblastoma Patients. <i>BMC Cancer</i> , 2009, 9, 436. | 1.1 | 7 |
| 54 | High frequency of BRCA1/2 germline mutations in consecutive ovarian cancer patients in Poland. <i>Gynecologic Oncology</i> , 2008, 108, 433-437. | 0.6 | 50 |

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|----|--|-----|-----------|
| 55 | BRCA1 and BRCA2 point mutations and large rearrangements in breast and ovarian cancer families in Northern Poland. <i>Oncology Reports</i> , 2008, , . | 1.2 | 21 |
| 56 | BRCA1 and BRCA2 point mutations and large rearrangements in breast and ovarian cancer families in Northern Poland. <i>Oncology Reports</i> , 2008, 19, 263-8. | 1.2 | 51 |
| 57 | Risk Criteria and Prognostic Factors for Predicting Recurrences After Resection of Primary Gastrointestinal Stromal Tumor. <i>Annals of Surgical Oncology</i> , 2007, 14, 2018-2027. | 0.7 | 227 |
| 58 | Predictive factors for long-term effects of imatinib therapy in patients with inoperable/metastatic CD117(+) gastrointestinal stromal tumors (GISTs). <i>Journal of Cancer Research and Clinical Oncology</i> , 2007, 133, 589-597. | 1.2 | 31 |
| 59 | Mutational and genotypeâ€“phenotype correlation analyses in 28 Polish patients with Cornelia de Lange syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 1531-1541. | 0.7 | 55 |
| 60 | C-344T polymorphism of the aldosterone synthase gene and blood pressure in the elderly: a population-based study. <i>Journal of Hypertension</i> , 2005, 23, 1991-1996. | 0.3 | 44 |
| 61 | Microdeletion and IGF2 loss of imprinting in a cascade causing Beckwith-Wiedemann syndrome with Wilms' tumor. <i>Nature Genetics</i> , 2005, 37, 785-786. | 9.4 | 40 |
| 62 | Cytogenetic and molecular findings in 75 clear cell renal cell carcinomas. <i>Oncology Reports</i> , 2005, 13, 949. | 1.2 | 9 |
| 63 | Microdeletion of target sites for insulator protein CTCF in a chromosome 11p15 imprinting center in Beckwith-Wiedemann syndrome and Wilms' tumor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 4085-4090. | 3.3 | 124 |
| 64 | Prevalence and clinical correlations of BRCA1/BRCA2 unclassified variant carriers among unselected primary ovarian cancer cases â€“ preliminary report. <i>European Journal of Cancer</i> , 2005, 41, 143-150. | 1.3 | 22 |
| 65 | The magic human 46 chromosomes were immortalised on a bronze plaque at Lund University in Sweden. <i>Journal of Applied Genetics</i> , 2004, 45, 1-2. | 1.0 | 16 |
| 66 | Title is missing!. <i>Journal of Neuro-Oncology</i> , 2003, 64, 284-284. | 1.4 | 0 |
| 67 | BRCA1 and BRCA2 mutation analysis in breast-ovarian cancer families from northeastern Poland. <i>Human Mutation</i> , 2003, 21, 553-554. | 1.1 | 39 |
| 68 | Association between the PIA platelet glycoprotein GPIIIa polymorphism and extent of coronary artery disease. <i>International Journal of Cardiology</i> , 2003, 88, 229-237. | 0.8 | 15 |
| 69 | Calretinin and Other Mesothelioma Markers in Synovial Sarcoma. <i>American Journal of Surgical Pathology</i> , 2001, 25, 610-617. | 2.1 | 147 |
| 70 | Spectral karyotyping reveals 17;22 fusions in a cytogenetically atypical dermatofibrosarcoma protuberans with a large marker chromosome as a sole abnormality. <i>Genes Chromosomes and Cancer</i> , 2001, 31, 182-186. | 1.5 | 10 |
| 71 | Clinical impact of molecular and cytogenetic findings in synovial sarcoma. <i>Genes Chromosomes and Cancer</i> , 2001, 31, 362-372. | 1.5 | 108 |
| 72 | Cytogenetics of hepatoblastoma: Further characterization of 1q rearrangements by fluorescence in situ hybridization: An international collaborative study. , 2000, 34, 165-170. | | 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. <i>Human Mutation</i> , 2000, 15, 480-481. | 1.1 | 33 |
| 74 | Patterns of keratin polypeptides in 110 biphasic, monophasic, and poorly differentiated synovial sarcomas. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2000, 437, 275-283. | 1.4 | 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 Product "Ectodysplasin-A. <i>Fetal and Pediatric Pathology</i> , 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 gene HMGIC 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. <i>Cancer Genetics and Cytogenetics</i> , 1998, 101, 159-161. | 1.0 | 21 |
| 80 | Cytogenetic Findings in an Embryonal Sarcoma of the Liver. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Human Genetics</i> , 1997, 101, 295-298. | 1.8 | 16 |
| 83 | Evidence of somatic mutations in osteoarthritis. <i>Human Genetics</i> , 1996, 98, 651-656. | 1.8 | 30 |
| 84 | Clear cell sarcoma of tendons and aponeuroses with t(12;22) (q13;q12) diagnosed initially as malignant melanoma. <i>Cancer Genetics and Cytogenetics</i> , 1996, 91, 37-39. | 1.0 | 18 |
| 85 | Cytogenetic and Immunohistochemical Profile of Myxoid Liposarcoma. <i>American Journal of Clinical Pathology</i> , 1995, 103, 20-26. | 0.4 | 53 |
| 86 | Chromosome abnormalities in bilateral breast carcinomas. Cytogenetic evaluation of the clonal origin of multiple primary tumors. <i>Cancer</i> , 1995, 76, 250-258. | 2.0 | 76 |
| 87 | Cytogenetic findings in malignant peripheral nerve sheath tumors. <i>International Journal of Cancer</i> , 1995, 61, 793-798. | 2.3 | 75 |
| 88 | Abnormal karyotypes in three carcinomas of the gallbladder. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Genes Chromosomes and Cancer</i> , 1993, 6, 151-155. | 1.5 | 62 |
| 90 | Trisomy 12 and 4 in a thecoma of the ovary. <i>Gynecologic Oncology</i> , 1992, 45, 66-68. | 0.6 | 16 |

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|-----|---|-----|-----------|
| 91 | Cytogenetic analysis of 57 primary prostatic adenocarcinomas. <i>Genes Chromosomes and Cancer</i> , 1992, 4, 16-24. | 1.5 | 155 |
| 92 | Improved technique for short-term culture and cytogenetic analysis of human breast cancer. <i>Genes Chromosomes and Cancer</i> , 1992, 5, 14-20. | 1.5 | 94 |
| 93 | AgNOR staining in benign hyperplasia and carcinoma of the prostate. <i>Prostate</i> , 1991, 18, 155-162. | 1.2 | 15 |
| 94 | Trisomy of chromosome 12 in a case of thecoma of the ovary. <i>Gynecologic Oncology</i> , 1990, 36, 413-416. | 0.6 | 36 |
| 95 | An improved technique for short-term culturing of human prostatic adenocarcinoma tissue for cytogenetic analysis. <i>Cancer Genetics and Cytogenetics</i> , 1990, 46, 191-199. | 1.0 | 26 |
| 96 | Double minutes in two primary adenocarcinomas of the prostate. <i>Cancer Genetics and Cytogenetics</i> , 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. <i>Cancer Genetics and Cytogenetics</i> , 1986, 23, 291-299. | 1.0 | 235 |
| 98 | Chromosome abnormalities in two benign adipose tumors. <i>Cancer Genetics and Cytogenetics</i> , 1986, 22, 55-61. | 1.0 | 66 |
| 99 | Recurrent chromosome translocations in liposarcoma. <i>Cancer Genetics and Cytogenetics</i> , 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). <i>Cancer Genetics and Cytogenetics</i> , 1986, 23, 87-91. | 1.0 | 98 |
| 101 | Translocation X;18 in synovial sarcoma. <i>Cancer Genetics and Cytogenetics</i> , 1986, 23, 93. | 1.0 | 143 |