

John J Mulvihill

List of Publications by Year in descending order

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Version: 2024-02-01

87
papers

11,222
citations

81743

39
h-index

71532

76
g-index

91
all docs

91
docs citations

91
times ranked

9412
citing authors

#	ARTICLE	IF	CITATIONS
1	Projecting Individualized Probabilities of Developing Breast Cancer for White Females Who Are Being Examined Annually. <i>Journal of the National Cancer Institute</i> , 1989, 81, 1879-1886.	3.0	2,934
2	Study design and cohort characteristics of the childhood cancer survivor study: A multi-institutional collaborative project. <i>Medical and Pediatric Oncology</i> , 2002, 38, 229-239.	1.0	632
3	Long-Term Follow-up of von Recklinghausen Neurofibromatosis. <i>New England Journal of Medicine</i> , 1986, 314, 1010-1015.	13.9	609
4	The Childhood Cancer Survivor Study: A National Cancer Instituteâ€œSupported Resource for Outcome and Intervention Research. <i>Journal of Clinical Oncology</i> , 2009, 27, 2308-2318.	0.8	551
5	Ovarian Failure and Reproductive Outcomes After Childhood Cancer Treatment: Results From the Childhood Cancer Survivor Study. <i>Journal of Clinical Oncology</i> , 2009, 27, 2374-2381.	0.8	444
6	Effects of Treatment on Fertility in Long-Term Survivors of Childhood or Adolescent Cancer. <i>New England Journal of Medicine</i> , 1987, 317, 1315-1321.	13.9	437
7	Germline mutations in the Von Hippel-Lindau disease (VHL) gene in families from North America, Europe, and Japan. <i>Human Mutation</i> , 1996, 8, 348-357.	1.1	436
8	Developmental defects in gorlin syndrome related to a putative tumor suppressor gene on chromosome 9. <i>Cell</i> , 1992, 69, 111-117.	13.5	396
9	Early menopause in long-term survivors of cancer during adolescence. <i>American Journal of Obstetrics and Gynecology</i> , 1992, 166, 788-793.	0.7	395
10	Neurofibromatosis 1 (Recklinghausen Disease) and Neurofibromatosis 2 (Bilateral Acoustic) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 382 T	2.0	366
11	Impact of nine common type 2 diabetes risk polymorphisms in Asian Indian Sikhs: PPARG2 (Pro12Ala), IGF2BP2, TCF7L2 and FTO variants confer a significant risk. <i>BMC Medical Genetics</i> , 2008, 9, 59.	2.1	235
12	Patterns of inheritance in hypertrophic cardiomyopathy: Assessment by m-mode and two-dimensional echocardiography. <i>American Journal of Cardiology</i> , 1984, 53, 1087-1094.	0.7	233
13	Genetic Disease in Offspring of Long-Term Survivors of Childhood and Adolescent Cancer. <i>American Journal of Human Genetics</i> , 1998, 62, 45-52.	2.6	229
14	Pregnancy outcome following cancer chemotherapy. <i>American Journal of Medicine</i> , 1980, 69, 828-832.	0.6	193
15	Genome-Wide Association Study Identifies a Novel Locus Contributing to Type 2 Diabetes Susceptibility in Sikhs of Punjabi Origin From India. <i>Diabetes</i> , 2013, 62, 1746-1755.	0.3	167
16	Stillbirth and neonatal death in relation to radiation exposure before conception: a retrospective cohort study. <i>Lancet, The</i> , 2010, 376, 624-630.	6.3	144
17	Pregnancy outcome in cancer patients. Experience in a large cooperative group. <i>Cancer</i> , 1987, 60, 1143-1150.	2.0	141
18	The NIH Undiagnosed Diseases Program and Network: Applications to modern medicine. <i>Molecular Genetics and Metabolism</i> , 2016, 117, 393-400.	0.5	140

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19	Congenital Anomalies in the Children of Cancer Survivors: A Report From the Childhood Cancer Survivor Study. <i>Journal of Clinical Oncology</i> , 2012, 30, 239-245.	0.8	129
20	Recommendations for the integration of genomics into clinical practice. <i>Genetics in Medicine</i> , 2016, 18, 1075-1084.	1.1	125
21	Small head size after atomic irradiation. <i>Teratology</i> , 1976, 14, 355-357.	1.8	116
22	GENETIC EFFECTS OF RADIOTHERAPY FOR CHILDHOOD CANCER. <i>Health Physics</i> , 2003, 85, 65-80.	0.3	112
23	Reproductive problems and birth defects in survivors of Wilms' tumor and their relatives. <i>Medical and Pediatric Oncology</i> , 1988, 16, 233-240.	1.0	109
24	Chromosomal Abnormalities among Offspring of Childhood-Cancer Survivors in Denmark: A Population-Based Study. <i>American Journal of Human Genetics</i> , 2004, 74, 1282-1285.	2.6	107
25	Late effects of therapy in adult survivors of osteosarcoma and Ewing's sarcoma. <i>Medical and Pediatric Oncology</i> , 1992, 20, 6-12.	1.0	99
26	Genetic Disease in the Children of Danish Survivors of Childhood and Adolescent Cancer. <i>Journal of Clinical Oncology</i> , 2012, 30, 27-33.	0.8	99
27	Cholesterol and bile acid replacement therapy in children and adults with Smith-Lemli-Opitz (SLO/RSH) syndrome. , 1997, 68, 315-321.		97
28	Smoking habits in survivors of childhood and adolescent cancer. <i>Medical and Pediatric Oncology</i> , 1992, 20, 301-306.	1.0	89
29	A gene for Crouzon craniofacial dysostosis maps to the long arm of chromosome 10. <i>Nature Genetics</i> , 1994, 7, 149-153.	9.4	86
30	Craniofacial syndromes: no such thing as a single gene disease. <i>Nature Genetics</i> , 1995, 9, 101-103.	9.4	82
31	A dyadic approach to the delineation of diagnostic entities in clinical genomics. <i>American Journal of Human Genetics</i> , 2021, 108, 8-15.	2.6	71
32	Variants in KCNQ1 increase type II diabetes susceptibility in South Asians: A study of 3,310 subjects from India and the US. <i>BMC Medical Genetics</i> , 2011, 12, 18.	2.1	59
33	Reporting genomic secondary findings: ACMG members weigh in. <i>Genetics in Medicine</i> , 2015, 17, 27-35.	1.1	57
34	Ethical issues of CRISPR technology and gene editing through the lens of solidarity. <i>British Medical Bulletin</i> , 2017, 122, 17-29.	2.7	57
35	SC phocomelia syndrome, premature centromere separation, and congenital cranial nerve paralysis in two sisters, one with malignant melanoma. <i>American Journal of Medical Genetics Part A</i> , 1986, 24, 653-672.	2.4	51
36	Caffeine as teratogen and mutagen. <i>Teratology</i> , 1973, 8, 69-72.	1.8	44

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37	Familial Sotos syndrome (cerebral gigantism): Craniofacial and psychological characteristics. American Journal of Medical Genetics Part A, 1985, 20, 613-624.	2.4	44
38	Sister chromatid exchanges and chromosomes in chronic myelogenous leukemia and cancer families. International Journal of Cancer, 1979, 23, 8-13.	2.3	43
39	Risk of cancer among children of cancer patients—a nationwide study in Finland. International Journal of Cancer, 2010, 126, 1196-1205.	2.3	41
40	Prevention in familial breast cancer: counseling and prophylactic mastectomy. Preventive Medicine, 1982, 11, 500-511.	1.6	39
41	Discussion: Genetics of multiple primary tumors. A clinical etiologic approach illustrated by three patients. Cancer, 1977, 40, 1867-1871.	2.0	38
42	Multiple childhood osteosarcomas in an American Indian family with erythroid macrocytosis and skeletal anomalies. Cancer, 1977, 40, 3115-3122.	2.0	37
43	Congenital heart disease in dogs: Epidemiologic similarities to man. Teratology, 1973, 7, 73-77.	1.8	36
44	Testing the association of novel meta-analysis-derived diabetes risk genes with type II diabetes and related metabolic traits in Asian Indian Sikhs. Journal of Human Genetics, 2009, 54, 162-168.	1.1	36
45	Replication of Association Between a Common Variant Near <i>Melanocortin-4 Receptor</i> Gene and Obesity-related Traits in Asian Sikhs. Obesity, 2010, 18, 425-429.	1.5	36
46	Familial eosinophilia: Clinical and laboratory results on a U.S. Kindred. American Journal of Medical Genetics Part A, 1998, 76, 229-237.	2.4	35
47	Germline mutations in the Von Hippel-Lindau disease (VHL) gene in families from North America, Europe, and Japan. Human Mutation, 1996, 8, 348-357.	1.1	33
48	Cleft palate in domestic animals: Epidemiologic features. Teratology, 1980, 21, 109-112.	1.8	31
49	Hospitalizations among children of survivors of childhood and adolescent cancer: A population-based cohort study. International Journal of Cancer, 2010, 127, 2879-2887.	2.3	31
50	Cancer chemotherapeutic agents as human teratogens. Birth Defects Research Part A: Clinical and Molecular Teratology, 2012, 94, 626-650.	1.6	31
51	Lymphangiosarcoma in late-onset hereditary lymphedema: Case report and nosological implications. American Journal of Medical Genetics Part A, 1995, 56, 72-75.	2.4	29
52	Cancer Risk Assessment and Genetic Counseling in an Academic Medical Center: Consultants' Satisfaction, Knowledge, and Behavior in the First Year. Journal of Genetic Counseling, 1998, 7, 279-297.	0.9	29
53	Surgical Management of Spinal Cord Compression from Plexiform Neurofibromas in Patients with Neurofibromatosis 1. Neurosurgery, 1998, 43, 248-255.	0.6	29
54	Harnessing genomics to identify environmental determinants of heritable disease. Mutation Research - Reviews in Mutation Research, 2013, 752, 6-9.	2.4	25

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55	Sister chromatid exchange and chromosome fragility in the nevoid basal cell carcinoma syndrome. <i>Cancer Genetics and Cytogenetics</i> , 1989, 42, 273-279.	1.0	23
56	Spectrum of malignancy and premalignancy in Carney syndrome. , 1997, 73, 369-377.		21
57	Further delineation of the Baller-Gerold syndrome. <i>American Journal of Medical Genetics Part A</i> , 1993, 45, 519-524.	2.4	20
58	Genetic counseling of the cancer survivor. <i>Seminars in Oncology Nursing</i> , 1989, 5, 29-35.	0.7	15
59	Smith-Lemli-Opitz syndrome: Biochemical before clinical diagnosis; early dietary management. <i>American Journal of Medical Genetics Part A</i> , 1994, 50, 375-376.	2.4	15
60	Multisystem burden of neurofibromatosis 1 in Denmark: registry- and population-based rates of hospitalizations over the life span. <i>Genetics in Medicine</i> , 2020, 22, 1069-1078.	1.1	15
61	The Floating Harbor syndrome with cardiac septal defect. , 1996, 66, 300-302.		13
62	Falling giants and the rise of gene editing: ethics, private interests and the public good. <i>Human Genomics</i> , 2017, 11, 20.	1.4	12
63	Preconception exposure to mutagens: medical and other exposures to radiation and chemicals. <i>Journal of Community Genetics</i> , 2012, 3, 205-211.	0.5	11
64	Late effects of therapy in survivors of childhood and adolescent osteosarcoma. <i>Cancer Treatment and Research</i> , 1993, 62, 45-48.	0.2	10
65	Neurofibromatosis.. <i>Annals of the New York Academy of Sciences</i> , 1986, 486, 38-44.	1.8	9
66	Chromosomal Abnormalities in Offspring of Young Cancer Survivors: A Population-Based Cohort Study in Denmark. <i>Journal of the National Cancer Institute</i> , 2018, 110, 534-538.	3.0	9
67	Psychiatric disorders in individuals with neurofibromatosis 1 in Denmark: A nationwide register-based cohort study. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 3706-3716.	0.7	8
68	The concurrence of saethre-chotzen syndrome and malignancy in a family with in vitro immune dysfunction. <i>Cancer</i> , 1984, 54, 2946-2951.	2.0	7
69	Statement on bioinformatics and capturing the benefits of genome sequencing for society. <i>Human Genomics</i> , 2019, 13, 24.	1.4	6
70	Forming and ending marital or cohabiting relationships in a Danish population-based cohort of individuals with neurofibromatosis 1. <i>European Journal of Human Genetics</i> , 2020, 28, 1028-1033.	1.4	5
71	Breast Cancer Risk Analysis and Counseling. <i>Clinical Obstetrics and Gynecology</i> , 1996, 39, 851-859.	0.6	5
72	AN EXAMPLE OF SAMPLE SIZE DETERMINATION FOR FAMILY STUDIES. <i>American Journal of Epidemiology</i> , 1981, 114, 299-303.	1.6	4

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73	A steroid metabolizing gene variant in a polyfactorial model improves risk prediction in a high incidence breast cancer population. <i>BBA Clinical</i> , 2014, 2, 94-102.	4.1	4
74	Pregnancy outcomes in women with neurofibromatosis 1: a Danish population-based cohort study. <i>Journal of Medical Genetics</i> , 2022, 59, 237-242.	1.5	3
75	Familial breast cancer in black Americans. <i>Cancer</i> , 1987, 60, 1657-1660.	2.0	2
76	Interstitial lung disease in an adult with Fanconi anemia: Clues to the pathogenesis. , 1997, 69, 315-319.		2
77	The View of CRISPR Patents Through the Lens of Solidarity and the Public Good. <i>American Journal of Bioethics</i> , 2018, 18, 54-56.	0.5	2
78	Reproductive Outcomes among Men Treated for Cancer. , 1994, , 197-203.		2
79	Encomium: Robert Warwick Miller: Mentor, synthesizer, and international interdisciplinary initiator. , 1998, 76, 1-8.		1
80	Familial eosinophilia: Clinical and laboratory results on a U.S. Kindred. , 1998, 76, 229.		1
81	Clinical Genetics of Human Cancer. , 1984, , 13-34.		1
82	Expanding metabolic screening of newborns: can the health care industry do better than public health?. <i>Journal - Oklahoma State Medical Association</i> , 2003, 96, 477-81.	0.4	1
83	Permanent Committee of the International Congresses of Human Genetics. , 1998, 79, 79-81.		0
84	The joy and duty of a marginal teratologist. <i>Birth Defects Research</i> , 2020, 112, 918-928.	0.8	0
85	Response to Hamosh etÅal.. <i>American Journal of Human Genetics</i> , 2021, 108, 1809-1810.	2.6	0
86	Cancer Genetics: <i>Inheritance of Susceptibility to Cancer in Man</i> . W. F. Bodmer, Ed. Oxford University Press, New York, 1983. vi, 192 pp., illus. \$26.95. Originally published in <i>Cancer Surveys</i> , vol. I, no. 1.. <i>Science</i> , 1984, 223, 162-162.	6.0	0
87	Celebrating the structure of DNA: 50 years and beyond. <i>Journal - Oklahoma State Medical Association</i> , 2003, 96, 184.	0.4	0