

# Terri McVeigh

## List of Publications by Year in descending order

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Version: 2024-02-01

74  
papers

898  
citations

567144

15  
h-index

526166

27  
g-index

77  
all docs

77  
docs citations

77  
times ranked

1961  
citing authors

#	ARTICLE	IF	CITATIONS
1	UK recommendations for <i>SDHA</i> germline genetic testing and surveillance in clinical practice. <i>Journal of Medical Genetics</i> , 2023, 60, 107-111.	1.5	4
2	Approach to screening for Familial Adenomatous Polyposis (FAP) in a cohort of 226 patients with Desmond-type Fibromatosis (DF): experience of a specialist center in the UK. <i>Familial Cancer</i> , 2022, 21, 69-74.	0.9	3
3	Uterine leiomyomatosis in adolescents and young adults (AYAs) may represent a narrow phenotypic variant of FH tumour predisposition syndrome. <i>Familial Cancer</i> , 2022, 21, 357-362.	0.9	1
4	Quantifying evidence toward pathogenicity for rare phenotypes: The case of succinate dehydrogenase genes, <i>SDHB</i> and <i>SDHD</i> . <i>Genetics in Medicine</i> , 2022, 24, 41-50.	1.1	5
5	A pilot of Blood-First diagnostic cell free DNA (cfDNA) next generation sequencing (NGS) in patients with suspected advanced lung cancer. <i>Lung Cancer</i> , 2022, 165, 34-42.	0.9	20
6	KIT-Associated Familial GIST Syndrome: Response to Tyrosine Kinase Inhibitors and Implications for Risk Management. <i>Oncologist</i> , 2022, 27, 615-620.	1.9	6
7	Management strategies for the colonoscopic surveillance of people with Lynch syndrome during the COVID-19 pandemic. <i>Gut</i> , 2021, 70, 624-626.	6.1	7
8	A systematic review and meta-analysis of telephone vs in-person genetic counseling in <i>BRCA1</i> / <i>BRCA2</i> genetic testing. <i>Journal of Genetic Counseling</i> , 2021, 30, 563-573.	0.9	12
9	Should All Individuals Be Screened for Genetic Predisposition to Cancer?. <i>Genetical Research</i> , 2021, 2021, 1-6.	0.3	6
10	FOXE1 polymorphism rs965513 predisposes to thyroid cancer in a European cohort. <i>Endocrine Oncology</i> , 2021, 1, 1-8.	0.1	0
11	Postgraduate training in Cancer Genetics—a cross-specialty survey exploring experience of clinicians in Ireland. <i>Irish Journal of Medical Science</i> , 2021, , 1.	0.8	2
12	Fibroadenoma in vulval ectopic breast tissue in a patient with PTEN Hamartoma Tumour Syndrome. <i>Familial Cancer</i> , 2021, , 1.	0.9	0
13	A clinical, molecular genetics and pathological study of a FTDP-17 family with a heterozygous splicing variant c.823-10G>T at the intron 9/exon 10 of the <i>MAPT</i> gene. <i>Neurobiology of Aging</i> , 2021, 106, 343.e1-343.e8.	1.5	5
14	A Review of Breast Cancer Risk Factors in Adolescents and Young Adults. <i>Cancers</i> , 2021, 13, 5552.	1.7	7
15	Fumarate Metabolic Signature for the Detection of Reed Syndrome in Humans. <i>Clinical Cancer Research</i> , 2020, 26, 391-396.	3.2	11
16	Sleep disordered breathing in children with Down syndrome in the Republic of Ireland. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 2847-2856.	0.7	3
17	Does Mode of Surgical Intervention Based on Oncotype DX Score Influence Disease Recurrence in Early Breast Cancer?. <i>The Surgery Journal</i> , 2020, 06, e135-e138.	0.3	0
18	Non-invasive Technology Advances in Cancer—A Review of the Advances in the Liquid Biopsy for Endometrial and Ovarian Cancers. <i>Frontiers in Digital Health</i> , 2020, 2, 573010.	1.5	3

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19	Diagnostic yield of a custom-designed multi-gene cancer panel in Irish patients with breast cancer. Irish Journal of Medical Science, 2020, 189, 849-864.	0.8	1
20	Genetic Testing for Cancer Predisposition Syndromes in Adolescents and Young Adults (AYAs). Current Genetic Medicine Reports, 2020, 8, 61-71.	1.9	3
21	Embryonal Rhabdomyosarcoma of the Ovary and Fallopian Tube. American Journal of Surgical Pathology, 2020, 44, 738-747.	2.1	42
22	Neoadjuvant chemoradiation and breast reconstruction: the potential for improved outcomes in the treatment of breast cancer. Irish Journal of Medical Science, 2019, 188, 75-83.	0.8	14
23	Multidisciplinary interventions in a specialist Drug Development Unit to improve family history documentation and onward referral of patients with advanced cancer to cancer genetics services. European Journal of Cancer, 2019, 114, 97-106.	1.3	4
24	Managing uncertainty in inherited cardiac pathologies—an international multidisciplinary survey. European Journal of Human Genetics, 2019, 27, 1178-1185.	1.4	6
25	P489—Cancer predisposition screening in children with cancer in Ireland. , 2019, , .		0
26	Towards establishing consistency in triage in a tertiary specialty. European Journal of Human Genetics, 2019, 27, 547-555.	1.4	6
27	Clinical Presentation of a Complex Neurodevelopmental Disorder Caused by Mutations in ADNP. Biological Psychiatry, 2019, 85, 287-297.	0.7	108
28	The role of genomic profiling in adolescents and young adults (AYAs) with advanced cancer participating in phase I clinical trials. European Journal of Cancer, 2018, 95, 20-29.	1.3	14
29	Non-syndromic bilateral ulnar aplasia with humero-radial synostosis and oligo-dactyly. American Journal of Medical Genetics, Part A, 2018, 176, 1180-1183.	0.7	1
30	The benefits of a Neurogenetics clinic in an adult Academic Teaching Hospital. Irish Journal of Medical Science, 2018, 187, 1073-1076.	0.8	4
31	Successful Repatriation of Breast Cancer Surveillance for High-Risk Women to the UK National Health Service Breast Screening Programme. Clinical Breast Cancer, 2018, 18, 282-288.	1.1	2
32	Effectiveness of contrast-associated acute kidney injury prevention methods; a systematic review and network meta-analysis. BMC Nephrology, 2018, 19, 323.	0.8	21
33	Differentiated Thyroid Cancer: How Do Current Practice Guidelines Affect Management?. European Thyroid Journal, 2018, 7, 319-326.	1.2	6
34	Clinical outcomes of adolescents and young adults with advanced solid tumours participating in phase I trials. European Journal of Cancer, 2018, 101, 55-61.	1.3	6
35	Teaching Radiology to Medical Students—There Is a Need for Change to Better Prepare Students for Clinical Practice. Academic Radiology, 2017, 24, 506-513.	1.3	13
36	Recurrent large genomic rearrangements in BRCA1 and BRCA2 in an Irish case series. Cancer Genetics, 2017, 214-215, 1-8.	0.2	5

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37	Screening for mismatch repair deficiency in colorectal cancer: data from three academic medical centers. <i>Cancer Medicine</i> , 2017, 6, 1465-1472.	1.3	30
38	Investigating the association of rs2910164 with cancer predisposition in an Irish cohort. <i>Endocrine Connections</i> , 2017, 6, 614-624.	0.8	5
39	The impact of the Biomolecular Era on breast cancer surgery. <i>Journal of the Royal College of Surgeons of Edinburgh</i> , 2017, 15, 169-181.	0.8	7
40	Follicular variant papillary thyroid carcinoma in Western Europe: A distinct clinical entity. <i>European Journal of Surgical Oncology</i> , 2017, 43, 2399-2400.	0.5	0
41	Impact of receptor phenotype on nodal burden in patients with breast cancer who have undergone neoadjuvant chemotherapy. <i>BJS Open</i> , 2017, 1, 39-45.	0.7	8
42	Clinical use of the Oncotype DX genomic test to guide treatment decisions for patients with invasive breast cancer. <i>Breast Cancer: Targets and Therapy</i> , 2017, Volume 9, 393-400.	1.0	64
43	Clinical outcomes of adolescents and young adults (AYA) with advanced solid tumors participating in phase I trials.. <i>Journal of Clinical Oncology</i> , 2017, 35, 10536-10536.	0.8	1
44	Personalisation of Therapy – clinical impact and relevance of genetic mutations in tumours. <i>Cancer Research Frontiers</i> , 2017, 3, 29-50.	0.2	2
45	THE EXPERIENCE OF THE FIRST ADULT NEUROGENETIC CLINIC IN IRELAND. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, e1.22-e1.	0.9	0
46	A qualitative analysis of the attitudes of Irish patients towards participation in genetic-based research. <i>Irish Journal of Medical Science</i> , 2016, 185, 825-831.	0.8	7
47	Classical galactosaemia: novel insights in IgG N-glycosylation and N-glycan biosynthesis. <i>European Journal of Human Genetics</i> , 2016, 24, 976-984.	1.4	60
48	MUTYH-Associated Polyposis: The Irish Experience. <i>Irish Medical Journal</i> , 2016, 109, 485.	0.0	1
49	21. FOXE1 – A potential thyroid cancer predisposition gene: A case-control study. <i>European Journal of Surgical Oncology</i> , 2015, 41, S85.	0.5	0
50	22. Investigating the role of polymorphism rs2910164 in mir146a in cancer predisposition. <i>European Journal of Surgical Oncology</i> , 2015, 41, S85-S86.	0.5	0
51	Kabuki syndrome. <i>Clinical Dysmorphology</i> , 2015, 24, 135-139.	0.1	21
52	A genetic variant at 12p11 significantly modifies breast cancer risk in a genetically homogenous island population. <i>Breast Cancer Research and Treatment</i> , 2015, 149, 41-47.	1.1	1
53	The 8q24 rs6983267C variant is associated with increased thyroid cancer risk. <i>Endocrine-Related Cancer</i> , 2015, 22, 841-849.	1.6	16
54	Estrogen withdrawal, increased breast cancer risk and the KRAS-variant. <i>Cell Cycle</i> , 2015, 14, 2091-2099.	1.3	11

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55	Targeted resequencing of the microRNAome and 3'UTRome reveals functional germline DNA variants with altered prevalence in epithelial ovarian cancer. <i>Oncogene</i> , 2015, 34, 2125-2137.	2.6	24
56	Abstract P6-10-02: The KRAS-variant, multiple breast cancer risk, and estrogen withdrawal. , 2015, , .		0
57	Estrogen withdrawal, breast cell transformation, and multiple breast cancer risk in women with the KRAS-variant.. <i>Journal of Clinical Oncology</i> , 2015, 33, e12541-e12541.	0.8	0
58	A qualitative analysis of the attitudes of Irish patients towards participation in genetic-based research.. <i>Journal of Clinical Oncology</i> , 2015, 33, e20564-e20564.	0.8	1
59	Immunohistochemistry to initiate a complex screening cascade in the detection of Lynch syndrome.. <i>Journal of Clinical Oncology</i> , 2015, 33, 3571-3571.	0.8	0
60	The acquisition and retention of urinary catheterisation skills using surgical simulator devices: teaching method or student traits. <i>BMC Medical Education</i> , 2014, 14, 264.	1.0	13
61	The impact of Oncotype DX testing on breast cancer management and chemotherapy prescribing patterns in a tertiary referral centre. <i>European Journal of Cancer</i> , 2014, 50, 2763-2770.	1.3	71
62	Endovascular Aneurysm Repair for Multiple Aneurysms as a Sequel of Hypereosinophilic Syndrome. <i>Vascular and Endovascular Surgery</i> , 2014, 48, 277-280.	0.3	4
63	Lobular Breast Cancer in a CDH1 Splice Site Mutation Carrier: Case Report and Review of the Literature. <i>Clinical Breast Cancer</i> , 2014, 14, e47-e51.	1.1	11
64	Familial breast cancer genetic testing in the West of Ireland. <i>Irish Journal of Medical Science</i> , 2014, 183, 199-206.	0.8	3
65	Gender differences in undergraduate medicine in Galway: a tale of two curricula. <i>Irish Journal of Medical Science</i> , 2014, 183, 103-110.	0.8	2
66	Assessing the Impact of Neoadjuvant Chemotherapy on the Management of the Breast and Axilla in Breast Cancer. <i>Clinical Breast Cancer</i> , 2014, 14, 20-25.	1.1	17
67	A germline mutation in the BRCA13'UTR predicts Stage IV breast cancer. <i>BMC Cancer</i> , 2014, 14, 421.	1.1	14
68	Sequencing of Therapy in Breast Cancer. <i>Oncology &amp; Hematology Review</i> , 2014, 10, 33.	0.2	0
69	Assessing awareness of colorectal cancer symptoms and screening in a peripheral colorectal surgical unit: a survey based study. <i>BMC Surgery</i> , 2013, 13, 20.	0.6	18
70	Increasing Reporting of Adverse Events to Improve the Educational Value of the Morbidity and Mortality Conference. <i>Journal of the American College of Surgeons</i> , 2013, 216, 50-56.	0.2	34
71	Assessing the impact of an ageing population on complication rates and in-patient length of stay. <i>International Journal of Surgery</i> , 2013, 11, 872-875.	1.1	32
72	Excuse Me: Teaching Interns to Speak Up. <i>Joint Commission Journal on Quality and Patient Safety</i> , 2013, 39, 426-431.	0.4	35

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73	Changing practices in the surgical management of hyperparathyroidism – A 10-year review. Journal of the Royal College of Surgeons of Edinburgh, 2012, 10, 314-320.	0.8	4
74	The KRAS-Variant Is Associated with Risk of Developing Double Primary Breast and Ovarian Cancer. PLoS ONE, 2012, 7, e37891.	1.1	30