Terri McVeigh

List of Publications by Year
in descending order

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	Clinical Presentation of a Complex Neurodevelopmental Disorder Caused by Mutations in ADNP. Biological Psychiatry, 2019, 85, 287-297.	0.7	108
2	The impact of Oncotype DX testing on breast cancer management and chemotherapy prescribing patterns in a tertiary referral centre. European Journal of Cancer, 2014, 50, 2763-2770.	1.3	71
3	Clinical use of the Oncotype DX genomic test to guide treatment decisions for patients with invasive breast cancer. Breast Cancer: Targets and Therapy, 2017, Volume 9, 393-400.	1.0	64
4	Classical galactosaemia: novel insights in IgG N-glycosylation and N-glycan biosynthesis. European Journal of Human Genetics, 2016, 24, 976-984.	1.4	60
5	Embryonal Rhabdomyosarcoma of the Ovary and Fallopian Tube. American Journal of Surgical Pathology, 2020, 44, 738-747.	2.1	42
6	â€œExcuse Me:â€•Teaching Interns to Speak Up. Joint Commission Journal on Quality and Patient Safety, 2013, 39, 426-431.	0.4	35
7	Increasing Reporting of Adverse Events to Improve the Educational Value of the Morbidity and Mortality Conference. Journal of the American College of Surgeons, 2013, 216, 50-56.	0.2	34
8	Assessing the impact of an ageing population on complication rates and in-patient length of stay. International Journal of Surgery, 2013, 11, 872-875.	1.1	32
9	The KRAS-Variant Is Associated with Risk of Developing Double Primary Breast and Ovarian Cancer. PLoS ONE, 2012, 7, e37891.	1.1	30
10	Screening for mismatch repair deficiency in colorectal cancer: data from three academic medical centers. Cancer Medicine, 2017, 6, 1465-1472.	1.3	30
11	Targeted resequencing of the microRNAome and 3â€²UTRome reveals functional germline DNA variants with altered prevalence in epithelial ovarian cancer. Oncogene, 2015, 34, 2125-2137.	2.6	24
12	Kabuki syndrome. Clinical Dysmorphology, 2015, 24, 135-139.	0.1	21
13	Effectiveness of contrast-associated acute kidney injury prevention methods; a systematic review and network meta-analysis. BMC Nephrology, 2018, 19, 323.	0.8	21
14	A pilot of Blood-First diagnostic cell free DNA (cfDNA) next generation sequencing (NGS) in patients with suspected advanced lung cancer. Lung Cancer, 2022, 165, 34-42.	0.9	20
15	Assessing awareness of colorectal cancer symptoms and screening in a peripheral colorectal surgical unit: a survey based study. BMC Surgery, 2013, 13, 20.	0.6	18
16	Assessing the Impact of Neoadjuvant Chemotherapy on the Management of the Breast and Axilla in Breast Cancer. Clinical Breast Cancer, 2014, 14, 20-25.	1.1	17
17	The 8q24 rs6983267G variant is associated with increased thyroid cancer risk. Endocrine-Related Cancer, 2015, 22, 841-849.	1.6	16
18	A germline mutation in the BRCA13â€™UTR predicts Stage IV breast cancer. BMC Cancer, 2014, 14, 421.	1.1	14

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19	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
20	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
21	The acquisition and retention of urinary catheterisation skills using surgical simulator devices: teaching method or student traits. BMC Medical Education, 2014, 14, 264.	1.0	13
22	Teaching Radiology to Medical Studentsâ€"There Is a Need for Change to Better Prepare Students for Clinical Practice. Academic Radiology, 2017, 24, 506-513.	1.3	13
23	A systematic review and metaâ€analysis of telephone vs inâ€person genetic counseling in BRCA1/BRCA2 genetic testing. Journal of Genetic Counseling, 2021, 30, 563-573.	0.9	12
24	Lobular Breast Cancer in a CDH1 Splice Site Mutation Carrier: Case Report and Review of the Literature. Clinical Breast Cancer, 2014, 14, e47-e51.	1.1	11
25	Estrogen withdrawal, increased breast cancer risk and the KRAS-variant. Cell Cycle, 2015, 14, 2091-2099.	1.3	11
26	Fumarate Metabolic Signature for the Detection of Reed Syndrome in Humans. Clinical Cancer Research, 2020, 26, 391-396.	3.2	11
27	Impact of receptor phenotype on nodal burden in patients with breast cancer who have undergone neoadjuvant chemotherapy. BJS Open, 2017, 1, 39-45.	0.7	8
28	A qualitative analysis of the attitudes of Irish patients towards participation in genetic-based research. Irish Journal of Medical Science, 2016, 185, 825-831.	0.8	7
29	The impact of the Biomolecular Era on breast cancer surgery. Journal of the Royal College of Surgeons of Edinburgh, 2017, 15, 169-181.	0.8	7
30	Management strategies for the colonoscopic surveillance of people with Lynch syndrome during the COVID-19 pandemic. Gut, 2021, 70, 624-626.	6.1	7
31	A Review of Breast Cancer Risk Factors in Adolescents and Young Adults. Cancers, 2021, 13, 5552.	1.7	7
32	Differentiated Thyroid Cancer: How Do Current Practice Guidelines Affect Management?. European Thyroid Journal, 2018, 7, 319-326.	1.2	6
33	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
34	Managing uncertainty in inherited cardiac pathologiesâ€"an international multidisciplinary survey. European Journal of Human Genetics, 2019, 27, 1178-1185.	1.4	6
35	Towards establishing consistency in triage in a tertiary specialty. European Journal of Human Genetics, 2019, 27, 547-555.	1.4	6
36	Should All Individuals Be Screened for Genetic Predisposition to Cancer?. Genetical Research, 2021, 2021, 1-6.	0.3	6

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37	KIT-Associated Familial GIST Syndrome: Response to Tyrosine Kinase Inhibitors and Implications for Risk Management. Oncologist, 2022, 27, 615-620.	1.9	6
38	Recurrent large genomic rearrangements in BRCA1 and BRCA2 in an Irish case series. Cancer Genetics, 2017, 214-215, 1-8.	0.2	5
39	Investigating the association of rs2910164 with cancer predisposition in an Irish cohort. Endocrine Connections, 2017, 6, 614-624.	0.8	5
40	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 MAPT gene. Neurobiology of Aging, 2021, 106, 343.e1-343.e8.	1.5	5
41	Quantifying evidence toward pathogenicity for rare phenotypes: The case of succinate dehydrogenase genes, SDHB and SDHD. Genetics in Medicine, 2022, 24, 41-50.	1.1	5
42	Changing practices in the surgical management of hyperparathyroidism â€“ A 10-year review. Journal of the Royal College of Surgeons of Edinburgh, 2012, 10, 314-320.	0.8	4
43	Endovascular Aneurysm Repair for Multiple Aneurysms as a Sequel of Hypereosinophilic Syndrome. Vascular and Endovascular Surgery, 2014, 48, 277-280.	0.3	4
44	The benefits of a Neurogenetics clinic in an adult Academic Teaching Hospital. Irish Journal of Medical Science, 2018, 187, 1073-1076.	0.8	4
45	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
46	UK recommendations forSDHAgermline genetic testing and surveillance in clinical practice. Journal of Medical Genetics, 2023, 60, 107-111.	1.5	4
47	Familial breast cancer genetic testing in the West of Ireland. Irish Journal of Medical Science, 2014, 183, 199-206.	0.8	3
48	Sleep disordered breathing in children with Down syndrome in the Republic of Ireland. American Journal of Medical Genetics, Part A, 2020, 182, 2847-2856.	0.7	3
49	Non-invasive Technology Advances in Cancerâ€”A Review of the Advances in the Liquid Biopsy for Endometrial and Ovarian Cancers. Frontiers in Digital Health, 2020, 2, 573010.	1.5	3
50	Genetic Testing for Cancer Predisposition Syndromes in Adolescents and Young Adults (AYAs). Current Genetic Medicine Reports, 2020, 8, 61-71.	1.9	3
51	Approach to screening for Familial Adenomatous Polyposis (FAP) in a cohort of 226 patients with Desmoid-type Fibromatosis (DF): experience of a specialist center in the UK. Familial Cancer, 2022, 21, 69-74.	0.9	3
52	Gender differences in undergraduate medicine in Galway: a tale of two curricula. Irish Journal of Medical Science, 2014, 183, 103-110.	0.8	2
53	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
54	Postgraduate training in Cancer Geneticsâ€”a cross-specialty survey exploring experience of clinicians in Ireland. Irish Journal of Medical Science, 2021, , 1.	0.8	2

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55	Personalisation of Therapy â€“ clinical impact and relevance of genetic mutations in tumours. Cancer Research Frontiers, 2017, 3, 29-50.	0.2	2
56	A genetic variant at 12p11 significantly modifies breast cancer risk in a genetically homogenous island population. Breast Cancer Research and Treatment, 2015, 149, 41-47.	1.1	1
57	Nonâ€syndromic bilateral ulnar aplasia with humeroâ€radial synostosis and oligoâ€ectroâ€dactyly. American Journal of Medical Genetics, Part A, 2018, 176, 1180-1183.	0.7	1
58	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
59	Uterine leiomyomatosis in adolescents and young adults (AYAs) may represent a narrow phenotypic variant of FH tumour predisposition syndrome. Familial Cancer, 2022, 21, 357-362.	0.9	1
60	Clinical outcomes of adolescents and young adults (AYA) with advanced solid tumors participating in phase I trials.. Journal of Clinical Oncology, 2017, 35, 10536-10536.	0.8	1
61	A qualitative analysis of the attitudes of Irish patients towards participation in genetic-based research.. Journal of Clinical Oncology, 2015, 33, e20564-e20564.	0.8	1
62	MUTYH-Associated Polyposis: The Irish Experience›. Irish Medical Journal, 2016, 109, 485.	0.0	1
63	21. FOXE1 â€“ A potential thyroid cancer predisposition gene: A case-control study. European Journal of Surgical Oncology, 2015, 41, S85.	0.5	0
64	22. Investigating the role of polymorphism rs2910164 in mir146a in cancer predisposition. European Journal of Surgical Oncology, 2015, 41, S85-S86.	0.5	0
65	THE EXPERIENCE OF THE FIRST ADULT NEUROGENETIC CLINIC IN IRELAND. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, e1.22-e1.	0.9	0
66	Follicular variant papillary thyroid carcinoma in Western Europe: A distinct clinical entity. European Journal of Surgical Oncology, 2017, 43, 2399-2400.	0.5	0
67	P489â€…Cancer predisposition screening in children with cancer in ireland. , 2019, , .		0
68	Does Mode of Surgical Intervention Based on Oncotype DX Score Influence Disease Recurrence in Early Breast Cancer?. The Surgery Journal, 2020, 06, e135-e138.	0.3	0
69	FOXE1 polymorphism rs965513 predisposes to thyroid cancer in a European cohort. Endocrine Oncology, 2021, 1, 1-8.	0.1	0
70	Fibroadenoma in vulval ectopic breast tissue in a patient with PTENÂHamartoma Tumour Syndrome. Familial Cancer, 2021, , 1.	0.9	0
71	Sequencing of Therapy in Breast Cancer. Oncology & Hematology Review, 2014, 10, 33.	0.2	0
72	Abstract P6-10-02: The KRAS-variant, multiple breast cancer risk, and estrogen withdrawal. , 2015, , .		0

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73	Estrogen withdrawal, breast cell transformation, and multiple breast cancer risk in women with the KRAS-variant.. Journal of Clinical Oncology, 2015, 33, e12541-e12541.	0.8	0
74	Immunohistochemistry to initiate a complex screening cascade in the detection of Lynch syndrome.. Journal of Clinical Oncology, 2015, 33, 3571-3571.	0.8	0