

Chris Jacobs

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

30
papers

663
citations

11
h-index

25
g-index

50
ext. papers

832
ext. citations

4.5
avg, IF

3.68
L-index

#	Paper	IF	Citations
30	The Angelina Jolie effect: how high celebrity profile can have a major impact on provision of cancer related services. <i>Breast Cancer Research</i> , 2014 , 16, 442	8.3	207
29	Population testing for cancer predisposing BRCA1/BRCA2 mutations in the Ashkenazi-Jewish community: a randomized controlled trial. <i>Journal of the National Cancer Institute</i> , 2015 , 107, 379	9.7	112
28	Cost-effectiveness of population screening for BRCA mutations in Ashkenazi jewish women compared with family history-based testing. <i>Journal of the National Cancer Institute</i> , 2015 , 107, 380	9.7	111
27	Comparing family members' motivations and attitudes towards genetic testing for hereditary breast and ovarian cancer: a qualitative analysis. <i>European Journal of Human Genetics</i> , 2010 , 18, 1289-95	5.3	34
26	Cluster-randomised non-inferiority trial comparing DVD-assisted and traditional genetic counselling in systematic population testing for BRCA1/2 mutations. <i>Journal of Medical Genetics</i> , 2016 , 53, 472-80	5.8	32
25	Current detection rates and time-to-detection of all identifiable carriers in the Greater London population. <i>Journal of Medical Genetics</i> , 2018 , 55, 538-545	5.8	30
24	Mainstreaming genetics and genomics: a systematic review of the barriers and facilitators for nurses and physicians in secondary and tertiary care. <i>Genetics in Medicine</i> , 2020 , 22, 1149-1155	8.1	23
23	Communication about genetic testing with breast and ovarian cancer patients: a scoping review. <i>European Journal of Human Genetics</i> , 2019 , 27, 511-524	5.3	22
22	Accuracy of recall of information about a cancer-predisposing BRCA1/2 gene mutation among patients and relatives. <i>European Journal of Human Genetics</i> , 2015 , 23, 147-51	5.3	17
21	Risk reducing salpingectomy and delayed oophorectomy in high risk women: views of cancer geneticists, genetic counsellors and gynaecological oncologists in the UK. <i>Familial Cancer</i> , 2015 , 14, 521-30	3.0	14
20	Patient perspectives on molecular tumor profiling: "Why wouldn't you?". <i>BMC Cancer</i> , 2019 , 19, 753	4.8	12
19	The PiGeOn project: protocol of a longitudinal study examining psychosocial and ethical issues and outcomes in germline genomic sequencing for cancer. <i>BMC Cancer</i> , 2018 , 18, 454	4.8	10
18	Cancer patients' views and understanding of genome sequencing: a qualitative study. <i>Journal of Medical Genetics</i> , 2020 , 57, 671-676	5.8	7
17	The PiGeOn project: protocol for a longitudinal study examining psychosocial, behavioural and ethical issues and outcomes in cancer tumour genomic profiling. <i>BMC Cancer</i> , 2018 , 18, 389	4.8	7
16	Key messages for communicating information about BRCA1 and BRCA2 to women with breast or ovarian cancer: Consensus across health professionals and service users. <i>Psycho-Oncology</i> , 2017 , 26, 1818-1824	3.9	6
15	Identification of germline missense mutations and rare allelic variants in the ATM gene in early-onset breast cancer 1999 , 26, 286		4
14	Opportunities for the Implementation of Immersive Virtual Reality in Rehabilitation 2020 ,		3

13	Patient and Relative Experiences and Decision-making About Genetic Testing and Counseling for Familial ALS and FTD: A Systematic Scoping Review. <i>Alzheimer Disease and Associated Disorders</i> , 2021 , 35, 374-385	2.5	2
12	Family communication about genomic sequencing: A qualitative study with cancer patients and relatives. <i>Patient Education and Counseling</i> , 2021 , 104, 944-952	3.1	2
11	Systematic review of outcomes in studies of reproductive genetic carrier screening: Towards development of a core outcome set.. <i>Genetics in Medicine</i> , 2021 ,	8.1	1
10	Who should access germline genome sequencing? A mixed methods study of patient views. <i>Clinical Genetics</i> , 2020 , 97, 329-337	4	1
9	Advanced cancer patient preferences for receiving molecular profiling results. <i>Psycho-Oncology</i> , 2020 , 29, 1533-1539	3.9	1
8	Genetic counseling and testing practices for late-onset neurodegenerative disease: a systematic review. <i>Journal of Neurology</i> , 2021 , 1	5.5	1
7	Adapting to the challenges of the global pandemic on genetic counselor education: Evaluating students' satisfaction with virtual clinical experiences. <i>Journal of Genetic Counseling</i> , 2021 , 30, 1074-1083	3.5	1
6	Preparing the genetic counseling workforce for the future in Australasia. <i>Journal of Genetic Counseling</i> , 2021 , 30, 55-60	2.5	1
5	The Core Outcome DEvelopment for Carrier Screening (CODECS) study: protocol for development of a core outcome set. <i>Trials</i> , 2021 , 22, 480	2.8	1
4	One size does not fit all: The case for targeted education in genetics and genomics for cancer nurses. <i>European Journal of Cancer Care</i> , 2021 , 30, e13480	2.4	0
3	Australian human research ethics committee members' confidence in reviewing genomic research applications. <i>European Journal of Human Genetics</i> , 2021 , 29, 1811-1818	5.3	0
2	Toward genetic counseling practice standards for diagnostic testing in amyotrophic lateral sclerosis and frontotemporal dementia.. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2022 , 1-13	3.6	
1	Collaborative co-design and evaluation of an immersive virtual reality application prototype for communication rehabilitation (DISCOVER prototype).. <i>Disability and Rehabilitation: Assistive Technology</i> , 2022 , 1-10	1.8	