Chris Jacobs

List of Publications by Year in Descending Order

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	Toward genetic counseling practice standards for diagnostic testing in amyotrophic lateral sclerosis and frontotemporal dementia.. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration*, **2022**, 1-13	3.6	
29	Collaborative co-design and evaluation of an immersive virtual reality application prototype for communication rehabilitation (DISCOVR prototype).. *Disability and Rehabilitation: Assistive Technology*, **2022**, 1-10	1.8	
28	Systematic review of outcomes in studies of reproductive genetic carrier screening: Towards development of a core outcome set.. *Genetics in Medicine*, **2021**,	8.1	1
27	Genetic counseling and testing practices for late-onset neurodegenerative disease: a systematic review. *Journal of Neurology*, **2021**, 1	5.5	1
26	Patient and Relative Experiences and Decision-making About Genetic Testing and Counseling for Familial ALS and FTD: A Systematic Scoping Review. *Alzheimer Disease and Associated Disorders*, **2021**, 35, 374-385	2.5	2
25	One size does not fit all: The case for targeted education in genetics and genomics for cancer nurses. *European Journal of Cancer Care*, **2021**, 30, e13480	2.4	0
24	Adapting to the challenges of the global pandemic on genetic counselor education: Evaluating students' satisfaction with virtual clinical experiences. *Journal of Genetic Counseling*, **2021**, 30, 1074-1083	2.5	1
23	Preparing the genetic counseling workforce for the future in Australasia. *Journal of Genetic Counseling*, **2021**, 30, 55-60	2.5	1
22	Family communication about genomic sequencing: A qualitative study with cancer patients and relatives. *Patient Education and Counseling*, **2021**, 104, 944-952	3.1	2
21	The Core Outcome DEvelopment for Carrier Screening (CODECS) study: protocol for development of a core outcome set. *Trials*, **2021**, 22, 480	2.8	1
20	Australian human research ethics committee members' confidence in reviewing genomic research applications. *European Journal of Human Genetics*, **2021**, 29, 1811-1818	5.3	0
19	Mainstreaming genetics and genomics: a systematic review of the barriers and facilitators for nurses and physicians in secondary and tertiary care. *Genetics in Medicine*, **2020**, 22, 1149-1155	8.1	23
18	Cancer patients' views and understanding of genome sequencing: a qualitative study. *Journal of Medical Genetics*, **2020**, 57, 671-676	5.8	7
17	Opportunities for the Implementation of Immersive Virtual Reality in Rehabilitation **2020**,		3
16	Who should access germline genome sequencing? A mixed methods study of patient views. *Clinical Genetics*, **2020**, 97, 329-337	4	1
15	Advanced cancer patient preferences for receiving molecular profiling results. *Psycho-Oncology*, **2020**, 29, 1533-1539	3.9	1
14	Patient perspectives on molecular tumor profiling: "Why wouldn't you?". *BMC Cancer*, **2019**, 19, 753	4.8	12

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13	Communication about genetic testing with breast and ovarian cancer patients: a scoping review. *European Journal of Human Genetics*, **2019**, 27, 511-524	5.3	22
12	Current detection rates and time-to-detection of all identifiable carriers in the Greater London population. *Journal of Medical Genetics*, **2018**, 55, 538-545	5.8	30
11	The PiGeOn project: protocol for a longitudinal study examining psychosocial, behavioural and ethical issues and outcomes in cancer tumour genomic profiling. *BMC Cancer*, **2018**, 18, 389	4.8	7
10	The PiGeOn project: protocol of a longitudinal study examining psychosocial and ethical issues and outcomes in germline genomic sequencing for cancer. *BMC Cancer*, **2018**, 18, 454	4.8	10
9	Key messages for communicating information about BRCA1 and BRCA2 to women with breast or ovarian cancer: Consensus across health professionals and service users. *Psycho-Oncology*, **2017**, 26, 1818-1824	3.9	6
8	Cluster-randomised non-inferiority trial comparing DVD-assisted and traditional genetic counselling in systematic population testing for BRCA1/2 mutations. *Journal of Medical Genetics*, **2016**, 53, 472-80	5.8	32
7	Population testing for cancer predisposing BRCA1/BRCA2 mutations in the Ashkenazi-Jewish community: a randomized controlled trial. *Journal of the National Cancer Institute*, **2015**, 107, 379	9.7	112
6	Cost-effectiveness of population screening for BRCA mutations in Ashkenazi jewish women compared with family history-based testing. *Journal of the National Cancer Institute*, **2015**, 107, 380	9.7	111
5	Risk reducing salpingectomy and delayed oophorectomy in high risk women: views of cancer geneticists, genetic counsellors and gynaecological oncologists in the UK. *Familial Cancer*, **2015**, 14, 521-30	3	14
4	Accuracy of recall of information about a cancer-predisposing BRCA1/2 gene mutation among patients and relatives. *European Journal of Human Genetics*, **2015**, 23, 147-51	5.3	17
3	The Angelina Jolie effect: how high celebrity profile can have a major impact on provision of cancer related services. *Breast Cancer Research*, **2014**, 16, 442	8.3	207
2	Comparing family members' motivations and attitudes towards genetic testing for hereditary breast and ovarian cancer: a qualitative analysis. *European Journal of Human Genetics*, **2010**, 18, 1289-95	5.3	34
1	Identification of germline missense mutations and rare allelic variants in the ATM gene in early-onset breast cancer **1999**, 26, 286		4