Caroline Benjamin

List of Publications by Year in Descending Order

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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.

23 451 12 21 g-index

23 554 4 3.06 ext. papers ext. citations avg, IF L-index

#	Paper	IF	Citations
23	The genetic counseling profession in Austria: StakeholdersTperspectives. <i>Journal of Genetic Counseling</i> , 2021 , 30, 861-871	2.5	2
22	A Maturity Matrix for Nurse Leaders to Facilitate and Benchmark Progress in Genomic Healthcare Policy, Infrastructure, Education, and Delivery. <i>Journal of Nursing Scholarship</i> , 2020 , 52, 583-592	3.6	5
21	A Roadmap for Global Acceleration of Genomics Integration Across Nursing. <i>Journal of Nursing Scholarship</i> , 2020 , 52, 329-338	3.6	12
20	Exploring professional issues: the psychosocial component of genetic counseling in genomic healthcare. <i>Personalized Medicine</i> , 2020 , 17, 55-65	2.2	1
19	Recontacting patients in clinical genetics services: recommendations of the European Society of Human Genetics. <i>European Journal of Human Genetics</i> , 2019 , 27, 169-182	5.3	34
18	Recontacting or not recontacting? A survey of current practices in clinical genetics centres in Europe. <i>European Journal of Human Genetics</i> , 2018 , 26, 946-954	5.3	19
17	The Global Landscape of Nursing and Genomics. <i>Journal of Nursing Scholarship</i> , 2018 , 50, 249-256	3.6	31
16	Increasing nursing capacity in genomics: Overview of existing global genomics resources. <i>Nurse Education Today</i> , 2018 , 69, 53-59	3.7	12
15	A prospective cohort study assessing clinical referral management & workforce allocation within a UK regional medical genetics service. <i>European Journal of Human Genetics</i> , 2015 , 23, 996-1003	5.3	6
14	The role of the genetic counsellor: a systematic review of research evidence. <i>European Journal of Human Genetics</i> , 2015 , 23, 452-8	5.3	37
13	Position statement on opportunistic genomic screening from the Association of Genetic Nurses and Counsellors (UK and Ireland). <i>European Journal of Human Genetics</i> , 2014 , 22, 955-6	5.3	21
12	Confidence of primary care physicians in their ability to carry out basic medical genetic tasks-a European survey in five countries-Part 1. <i>Journal of Community Genetics</i> , 2011 , 2, 1-11	2.5	66
11	General practitioner management of genetic aspects of a cardiac disease: a scenario-based study to anticipate providersTpractices. <i>Journal of Community Genetics</i> , 2010 , 1, 83-90	2.5	6
10	Testing the children: do non-genetic health-care providers differ in their decision to advise genetic presymptomatic testing on minors? A cross-sectional study in five countries in the European Union. <i>Genetic Testing and Molecular Biomarkers</i> , 2009 , 13, 367-76	1.6	8
9	Educational priorities and current involvement in genetic practice: a survey of midwives in the Netherlands, UK and Sweden. <i>Midwifery</i> , 2009 , 25, 483-99	2.8	12
8	The use of the life course paradigm and life course charts to explore referral for family history of breast cancer. <i>International Journal of Nursing Studies</i> , 2008 , 45, 95-109	5.8	7
7	Genetics in clinical practice: general practitionersTeducational priorities in European countries. Genetics in Medicine, 2008, 10, 107-13	8.1	28

LIST OF PUBLICATIONS

6	Assessing educational priorities in genetics for general practitioners and specialists in five countries: factor structure of the Genetic-Educational Priorities (Gen-EP) scale. <i>Genetics in Medicine</i> , 2008 , 10, 99-106	8.1	26
5	Genetic education for non-geneticist health professionals. <i>Public Health Genomics</i> , 2006 , 9, 224-6	1.9	11
4	Genetic education and nongenetic health professionals: educational providers and curricula in Europe. <i>Genetics in Medicine</i> , 2005 , 7, 302-10	8.1	37
3	A Prospective Comparison Study of Different Methods of Gathering Self-Reported Family History Information for Breast Cancer Risk Assessment. <i>Journal of Genetic Counseling</i> , 2003 , 12, 151-70	2.5	11
2	United Kingdom experience with presymptomatic testing of individuals at 25% risk for Huntington's disease. <i>Clinical Genetics</i> , 2000 , 58, 41-9	4	21
1	Mapping of von Hippel-Lindau disease to chromosome 3p confirmed by genetic linkage analysis. Journal of the Neurological Sciences, 1990 , 100, 27-30	3.2	38