

Caroline Benjamin

List of Publications by Year in descending order

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Version: 2024-02-01

23
papers

633
citations

623734

14
h-index

642732

23
g-index

23
all docs

23
docs citations

23
times ranked

758
citing authors

#	ARTICLE	IF	CITATIONS
1	The genetic counseling profession in Austria: Stakeholders'™ perspectives. <i>Journal of Genetic Counseling</i> , 2021, 30, 861-871.	1.6	5
2	Exploring professional issues: the psychosocial component of genetic counseling in genomic healthcare. <i>Personalized Medicine</i> , 2020, 17, 55-65.	1.5	1
3	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.	2.4	14
4	A Roadmap for Global Acceleration of Genomics Integration Across Nursing. <i>Journal of Nursing Scholarship</i> , 2020, 52, 329-338.	2.4	24
5	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.	2.8	65
6	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.	2.8	33
7	The Global Landscape of Nursing and Genomics. <i>Journal of Nursing Scholarship</i> , 2018, 50, 249-256.	2.4	59
8	Increasing nursing capacity in genomics: Overview of existing global genomics resources. <i>Nurse Education Today</i> , 2018, 69, 53-59.	3.3	32
9	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.	2.8	8
10	The role of the genetic counsellor: a systematic review of research evidence. <i>European Journal of Human Genetics</i> , 2015, 23, 452-458.	2.8	57
11	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, 955-956.	2.8	25
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.	1.2	77
13	General practitioner management of genetic aspects of a cardiac disease: a scenario-based study to anticipate providers™ practices. <i>Journal of Community Genetics</i> , 2010, 1, 83-90.	1.2	9
14	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, 367-376.	0.7	8
15	Educational priorities and current involvement in genetic practice: a survey of midwives in the Netherlands, UK and Sweden. <i>Midwifery</i> , 2009, 25, 483-499.	2.3	16
16	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.	5.6	7
17	Genetics in clinical practice: general practitioners' educational priorities in European countries. <i>Genetics in Medicine</i> , 2008, 10, 107-113.	2.4	35
18	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.	2.4	30

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19	Genetic Education for Non-Geneticist Health Professionals. <i>Public Health Genomics</i> , 2006, 9, 224-226.	1.0	12
20	Genetic education and nongenetic health professionals: Educational providers and curricula in Europe. <i>Genetics in Medicine</i> , 2005, 7, 302-310.	2.4	39
21	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, 151-170.	1.6	11
22	United Kingdom experience with presymptomatic testing of individuals at 25% risk for Huntington's disease. <i>Clinical Genetics</i> , 2001, 58, 41-49.	2.0	22
23	Mapping of Von Hippel-Lindau disease to chromosome 3p confirmed by genetic linkage analysis. <i>Journal of the Neurological Sciences</i> , 1990, 100, 27-30.	0.6	44