Caroline Benjamin

List of Publications by Year in descending order

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623734 642732 23 633 14 23 citations g-index h-index papers 23 23 23 758 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Confidence of primary care physicians in their ability to carry out basic medical genetic tasks—a European survey in five countries—Part 1. Journal of Community Genetics, 2011, 2, 1-11.	1.2	77
2	Recontacting patients in clinical genetics services: recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2019, 27, 169-182.	2.8	65
3	The Global Landscape of Nursing and Genomics. Journal of Nursing Scholarship, 2018, 50, 249-256.	2.4	59
4	The role of the genetic counsellor: a systematic review of research evidence. European Journal of Human Genetics, 2015, 23, 452-458.	2.8	57
5	Mapping of Von Hippel-Lindau disease to chromosome 3p confirmed by genetic linkage analysis. Journal of the Neurological Sciences, 1990, 100, 27-30.	0.6	44
6	Genetic education and nongenetic health professionals: Educational providers and curricula in Europe. Genetics in Medicine, 2005, 7, 302-310.	2.4	39
7	Genetics in clinical practice: general practitioners' educational priorities in European countries. Genetics in Medicine, 2008, 10, 107-113.	2.4	35
8	Recontacting or not recontacting? A survey of current practices in clinical genetics centres in Europe. European Journal of Human Genetics, 2018, 26, 946-954.	2.8	33
9	Increasing nursing capacity in genomics: Overview of existing global genomics resources. Nurse Education Today, 2018, 69, 53-59.	3.3	32
10	Assessing educational priorities in genetics for general practitioners and specialists in five countries: factor structure of the Genetic-Educational Priorities (Gen-EP) scale. Genetics in Medicine, 2008, 10, 99-106.	2.4	30
11	Position statement on opportunistic genomic screening from the Association of Genetic Nurses and Counsellors (UK and Ireland). European Journal of Human Genetics, 2014, 22, 955-956.	2.8	25
12	A Roadmap for Global Acceleration of Genomics Integration Across Nursing. Journal of Nursing Scholarship, 2020, 52, 329-338.	2.4	24
13	United Kingdom experience with presymptomatic testing of individuals at 25% risk for Huntington's disease. Clinical Genetics, 2001, 58, 41-49.	2.0	22
14	Educational priorities and current involvement in genetic practice: a survey of midwives in the Netherlands, UK and Sweden. Midwifery, 2009, 25, 483-499.	2.3	16
15	A Maturity Matrix for Nurse Leaders to Facilitate and Benchmark Progress in Genomic Healthcare Policy, Infrastructure, Education, and Delivery. Journal of Nursing Scholarship, 2020, 52, 583-592.	2.4	14
16	Genetic Education for Non-Geneticist Health Professionals. Public Health Genomics, 2006, 9, 224-226.	1.0	12
17	A Prospective Comparison Study of Different Methods of Gathering Selfâ€Reported Family History Information for Breast Cancer Risk Assessment. Journal of Genetic Counseling, 2003, 12, 151-170.	1.6	11
18	General practitioner management of genetic aspects of a cardiac disease: a scenario-based study to anticipate providers' practices. Journal of Community Genetics, 2010, 1, 83-90.	1.2	9

#	Article	lF	CITATIONS
19	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. Genetic Testing and Molecular Biomarkers, 2009, 13, 367-376.	0.7	8
20	A prospective cohort study assessing clinical referral management & morkforce allocation within a UK regional medical genetics service. European Journal of Human Genetics, 2015, 23, 996-1003.	2.8	8
21	The use of the life course paradigm and life course charts to explore referral for family history of breast cancer. International Journal of Nursing Studies, 2008, 45, 95-109.	5.6	7
22	The genetic counseling profession in Austria: Stakeholders' perspectives. Journal of Genetic Counseling, 2021, 30, 861-871.	1.6	5
23	Exploring professional issues: the psychosocial component of genetic counseling in genomic healthcare. Personalized Medicine, 2020, 17, 55-65.	1.5	1