

Saskia C Sanderson

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

1,924
citations

394421

19
h-index

377865

34
g-index

35
all docs

35
docs citations

35
times ranked

2996
citing authors

#	ARTICLE	IF	CITATIONS
1	Return of Genomic Results to Research Participants: The Floor, the Ceiling, and the Choices In Between. <i>American Journal of Human Genetics</i> , 2014, 94, 818-826.	6.2	342
2	A systematic literature review of individuals'™ perspectives on broad consent and data sharing in the United States. <i>Genetics in Medicine</i> , 2016, 18, 663-671.	2.4	181
3	Public Attitudes toward Consent and Data Sharing in Biobank Research: A Large Multi-site Experimental Survey in the US. <i>American Journal of Human Genetics</i> , 2017, 100, 414-427.	6.2	172
4	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.	6.3	146
5	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.	6.3	135
6	Awareness of lifestyle risk factors for cancer and heart disease among adults in the UK. <i>Patient Education and Counseling</i> , 2009, 74, 221-227.	2.2	106
7	Motivations, concerns and preferences of personal genome sequencing research participants: Baseline findings from the HealthSeq project. <i>European Journal of Human Genetics</i> , 2016, 24, 14-20.	2.8	94
8	Willingness to participate in genomics research and desire for personal results among underrepresented minority patients: a structured interview study. <i>Journal of Community Genetics</i> , 2013, 4, 469-482.	1.2	82
9	Psychological and Behavioural Impact of Genetic Testing Smokers for Lung Cancer Risk. <i>Journal of Health Psychology</i> , 2008, 13, 481-494.	2.3	69
10	Stakeholder engagement: a key component of integrating genomic information into electronic health records. <i>Genetics in Medicine</i> , 2013, 15, 792-801.	2.4	64
11	Experiences and concerns of patients with recurrent attacks of acute hepatic porphyria: A qualitative study. <i>Molecular Genetics and Metabolism</i> , 2016, 119, 278-283.	1.1	60
12	Psychological and behavioural impact of returning personal results from whole-genome sequencing: the HealthSeq project. <i>European Journal of Human Genetics</i> , 2017, 25, 280-292.	2.8	54
13	Development and preliminary evaluation of an online educational video about whole-genome sequencing for research participants, patients, and the general public. <i>Genetics in Medicine</i> , 2016, 18, 501-512.	2.4	51
14	Responses to Online <i>GSTM1</i> Genetic Test Results among Smokers Related to Patients with Lung Cancer: A Pilot Study. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2009, 18, 1953-1961.	2.5	46
15	Predispositional genome sequencing in healthy adults: design, participant characteristics, and early outcomes of the PeopleSeq Consortium. <i>Genome Medicine</i> , 2019, 11, 10.	8.2	41
16	Informed decision-making among students analyzing their personal genomes on a whole genome sequencing course: a longitudinal cohort study. <i>Genome Medicine</i> , 2013, 5, 113.	8.2	29
17	How do students react to analyzing their own genomes in a whole-genome sequencing course?: outcomes of a longitudinal cohort study. <i>Genetics in Medicine</i> , 2015, 17, 866-874.	2.4	29
18	Practical Guidance on Informed Consent for Pediatric Participants in a Biorepository. <i>Mayo Clinic Proceedings</i> , 2014, 89, 1471-1480.	3.0	27

#	ARTICLE	IF	CITATIONS
19	Public Awareness of Genetic Influence on Chronic Disease Risk: Are Genetic and Lifestyle Causal Beliefs Compatible?. <i>Public Health Genomics</i> , 2011, 14, 290-297.	1.0	25
20	Parents' attitudes toward consent and data sharing in biobanks: A multisite experimental survey. <i>AJOB Empirical Bioethics</i> , 2018, 9, 128-142.	1.6	25
21	Impacts of incorporating personal genome sequencing into graduate genomics education: a longitudinal study over three course years. <i>BMC Medical Genomics</i> , 2018, 11, 5.	1.5	17
22	Preparing the next generation of genomicists: a laboratory-style course in medical genomics. <i>BMC Medical Genomics</i> , 2015, 8, 47.	1.5	16
23	Impact of Genomic Counseling on Informed Decision-Making among ostensibly Healthy Individuals Seeking Personal Genome Sequencing: the HealthSeq Project. <i>Journal of Genetic Counseling</i> , 2016, 25, 1044-1053.	1.6	15
24	Development of the Knowledge of Genome Sequencing (KOGS) questionnaire. <i>Patient Education and Counseling</i> , 2018, 101, 1966-1972.	2.2	15
25	Increasing genomic literacy among adolescents. <i>Genetics in Medicine</i> , 2019, 21, 994-1000.	2.4	14
26	Young people's understanding, attitudes and involvement in decision-making about genome sequencing for rare diseases: A qualitative study with participants in the UK 100,000 Genomes Project. <i>European Journal of Medical Genetics</i> , 2020, 63, 104043.	1.3	13
27	Factors Affecting Recall of Different Types of Personal Genetic Information about Alzheimer's Disease Risk: The REVEAL Study. <i>Public Health Genomics</i> , 2015, 18, 78-86.	1.0	10
28	Development and mixed-methods evaluation of an online animation for young people about genome sequencing. <i>European Journal of Human Genetics</i> , 2020, 28, 896-906.	2.8	10
29	Genome sequencing for healthy individuals. <i>Trends in Genetics</i> , 2013, 29, 556-558.	6.7	9
30	Conducting a large, multi-site survey about patients' views on broad consent: challenges and solutions. <i>BMC Medical Research Methodology</i> , 2016, 16, 162.	3.1	9
31	Effects of Testing and Disclosing Ancestry-Specific Genetic Risk for Kidney Failure on Patients and Health Care Professionals. <i>JAMA Network Open</i> , 2022, 5, e221048.	5.9	9
32	Development of a measure of genome sequencing knowledge for young people: The kids-KOGS. <i>Clinical Genetics</i> , 2019, 96, 411-417.	2.0	4
33	Article does not prove that smoking cessation has an "effect" on mental health. <i>BMJ, The</i> , 2014, 348, g2018-g2018.	6.0	3
34	Animation or leaflet: Does it make a difference when educating young people about genome sequencing?. <i>Patient Education and Counseling</i> , 2021, 104, 2522-2530.	2.2	2
35	Decision-Making in the Age of Whole Genome Sequencing. , 2016, , 357-373.		0