Sarah S Kalia

List of Publications by Year in descending order

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1162889 1474057 3,886 10 8 9 citations h-index g-index papers 10 10 10 7540 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Focused revision: An addendum to a National Society of Genetic Counselors (NSGC) practice resource. Journal of Genetic Counseling, 2020, 29, 135-135.	0.9	3
2	Secondary findings: How did we get here, and where are we going?. Journal of Genetic Counseling, 2019, 28, 326-333.	0.9	20
3	Effect of communicating personalized rheumatoid arthritis risk on concern for developing RA: A randomized controlled trial. Patient Education and Counseling, 2019, 102, 976-983.	1.0	20
4	Personal Genomic Testing for Cancer Risk: Results From the Impact of Personal Genomics Study. Journal of Clinical Oncology, 2017, 35, 636-644.	0.8	34
5	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2016 update (ACMG SF v2.0): a policy statement of the American College of Medical Genetics and Genomics. Genetics in Medicine, 2017, 19, 249-255.	1.1	1,398
6	Parents are interested in newborn genomic testing during the early postpartum period. Genetics in Medicine, 2015, 17, 501-504.	1.1	38
7	Design, methods, and participant characteristics of the Impact of Personal Genomics (PGen) Study, a prospective cohort study of direct-to-consumer personal genomic testing customers. Genome Medicine, 2014, 6, 96.	3.6	37
8	O3-02-01: DIRECT-TO-CONSUMER GENETIC TESTING FOR RISK OF ALZHEIMER'S DISEASE (AD): THE PSYCHOLOGICAL AND BEHAVIORAL IMPACT OF APOE GENOTYPE DISCLOSURE. , 2014, 10, P209-P209.		1
9	ACMG recommendations for reporting of incidental findings in clinical exome and genome sequencing. Genetics in Medicine, 2013, 15, 565-574.	1.1	2,186
10	Exploring concordance and discordance for return of incidental findings from clinical sequencing. Genetics in Medicine, 2012, 14, 405-410.	1.1	149