

Sandra Soo-Jin Lee

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

1,744
citations

331670

21
h-index

302126

39
g-index

64
all docs

64
docs citations

64
times ranked

1844
citing authors

#	ARTICLE	IF	CITATIONS
1	Health equality, race and pharmacogenomics. <i>British Journal of Clinical Pharmacology</i> , 2022, 88, 27-33.	2.4	24
2	Beyond inclusion: Enacting team equity in precision medicine research. <i>PLoS ONE</i> , 2022, 17, e0263750.	2.5	8
3	Laboratory-related outcomes from integrating an accessible delivery model for hereditary cancer risk assessment and genetic testing in populations with barriers to access. <i>Genetics in Medicine</i> , 2022, 24, 1196-1205.	2.4	6
4	Family secrets: Experiences and outcomes of participating in direct-to-consumer genetic relative-finder services. <i>American Journal of Human Genetics</i> , 2022, 109, 486-497.	6.2	12
5	Strategies of inclusion: The tradeoffs of pursuing "baked in" diversity through place-based recruitment. <i>Social Science and Medicine</i> , 2022, 306, 115132.	3.8	6
6	Challenges and potential solutions to health disparities in genomic medicine. <i>Cell</i> , 2022, 185, 2007-2010.	28.9	3
7	Participant Reactions to a Literacy-Focused, Web-Based Informed Consent Approach for a Genomic Implementation Study. <i>AJOB Empirical Bioethics</i> , 2021, 12, 1-11.	1.6	11
8	Is precision medicine relevant in the age of COVID-19?. <i>Genetics in Medicine</i> , 2021, 23, 999-1000.	2.4	17
9	Parental Attitudes Toward Clinical Genomic Sequencing in Children With Critical Cardiac Disease. <i>Pediatric Critical Care Medicine</i> , 2021, 22, e419-e426.	0.5	4
10	Response to Open Peer Commentaries: Distinguishing the "Gift" from "Donation" as a Path toward Reciprocity and Relational Ethics. <i>American Journal of Bioethics</i> , 2021, 21, W1-W3.	0.9	2
11	The Ethics of Consent in a Shifting Genomic Ecosystem. <i>Annual Review of Biomedical Data Science</i> , 2021, 4, 145-164.	6.5	6
12	Cancer Health Assessments Reaching Many (CHARM): A clinical trial assessing a multimodal cancer genetics services delivery program and its impact on diverse populations. <i>Contemporary Clinical Trials</i> , 2021, 106, 106432.	1.8	19
13	Obligations of the "Gift": Reciprocity and Responsibility in Precision Medicine. <i>American Journal of Bioethics</i> , 2021, 21, 57-66.	0.9	34
14	Feasibility of a Traceback Approach for Using Pathology Specimens to Facilitate Genetic Testing in the Genetic Risk Analysis in Ovarian Cancer (GRACE) Study Protocol. <i>Journal of Personalized Medicine</i> , 2021, 11, 1194.	2.5	1
15	Translating genomic testing results for pediatric critical care: Opportunities for genetic counselors. <i>Journal of Genetic Counseling</i> , 2020, 29, 78-87.	1.6	13
16	Excavating the Personal Genome: The Good Biocitizen in the Age of Precision Health. <i>Hastings Center Report</i> , 2020, 50, S54-S61.	1.0	4
17	Lessons learned about harmonizing survey measures for the CSER consortium. <i>Journal of Clinical and Translational Science</i> , 2020, 4, 537-546.	0.6	16
18	Integrating stakeholder feedback in translational genomics research: an ethnographic analysis of a study protocol's evolution. <i>Genetics in Medicine</i> , 2020, 22, 1094-1101.	2.4	11

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19	Ethics of inclusion: Cultivate trust in precision medicine. <i>Science</i> , 2019, 364, 941-942.	12.6	27
20	Anticipating uncertainty and irrevocable decisions: provider perspectives on implementing whole-genome sequencing in critically ill children with heart disease. <i>Genetics in Medicine</i> , 2018, 20, 1455-1461.	2.4	26
21	Beyond Consent: Building Trusting Relationships With Diverse Populations in Precision Medicine Research. <i>American Journal of Bioethics</i> , 2018, 18, 3-20.	0.9	152
22	Trustworthiness in Untrustworthy Times: Response to Open Peer Commentaries on <i>Beyond Consent</i>. <i>American Journal of Bioethics</i> , 2018, 18, W6-W8.	0.9	6
23	Studying "Friends": The Ethics of Using Social Media as Research Platforms. <i>American Journal of Bioethics</i> , 2017, 17, 1-2.	0.9	8
24	A randomized study of multimedia informational aids for research on medical practices: Implications for informed consent. <i>Clinical Trials</i> , 2017, 14, 94-102.	1.6	48
25	Consuming DNA: The Good Citizen in the Age of Precision Medicine. <i>Annual Review of Anthropology</i> , 2017, 46, 33-48.	1.5	9
26	Institutional culture is the key to team science. <i>Nature Biotechnology</i> , 2017, 35, 1212-1214.	17.5	9
27	Strategies for recruiting representative samples of Asian Americans for a population-based case-control study. <i>Journal of Epidemiology and Community Health</i> , 2016, 70, 974-982.	3.7	10
28	Introduction to the article collection "Translation in healthcare: ethical, legal, and social implications". <i>BMC Medical Ethics</i> , 2016, 17, 74.	2.4	14
29	Adrift in the gray zone: IRB perspectives on research in the learning health system. <i>AJOB Empirical Bioethics</i> , 2016, 7, 125-134.	1.6	29
30	A comparison of institutional review board professionals' and patients' views on consent for research on medical practices. <i>Clinical Trials</i> , 2016, 13, 555-565.	1.6	36
31	Enacting the molecular imperative: How gene-environment interaction research links bodies and environments in the post-genomic age. <i>Social Science and Medicine</i> , 2016, 155, 51-60.	3.8	52
32	The Role of Patient Perspectives in Clinical Research Ethics and Policy: Response to Open Peer Commentaries on "Patient Perspectives on the Learning Health System". <i>American Journal of Bioethics</i> , 2016, 16, W7-W9.	0.9	1
33	Accounting for Complexity. <i>Science Technology and Human Values</i> , 2016, 41, 194-218.	3.1	21
34	Direct-to-Consumer Personal Genetic Testing. , 2015, , 440-445.		0
35	Attitudes Toward Risk and Informed Consent for Research on Medical Practices. <i>Annals of Internal Medicine</i> , 2015, 162, 690-696.	3.9	87
36	Patient Perspectives on the Learning Health System: The Importance of Trust and Shared Decision Making. <i>American Journal of Bioethics</i> , 2015, 15, 4-17.	0.9	94

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37	The Biobank as Political Artifact. <i>Annals of the American Academy of Political and Social Science</i> , 2015, 661, 143-159.	1.6	7
38	Protecting Posted Genes: Social Networking and the Limits of GINA. <i>American Journal of Bioethics</i> , 2014, 14, 32-44.	0.9	17
39	Homogeneity and heterogeneity as situational properties: Producing "and moving beyond?" race in post-genomic science. <i>Social Studies of Science</i> , 2014, 44, 579-599.	2.5	25
40	Race and Ancestry in the Age of Inclusion: Technique and Meaning in Post-Genomic Science. <i>Journal of Health and Social Behavior</i> , 2014, 55, 504-518.	4.8	18
41	Views of Genetics Health Professionals on the Return of Genomic Results. <i>Journal of Genetic Counseling</i> , 2014, 23, 531-538.	1.6	43
42	The time is ripe for an ethics of entrepreneurship. <i>Nature Biotechnology</i> , 2014, 32, 316-318.	17.5	5
43	Personal genome testing in medical education: student experiences with genotyping in the classroom. <i>Genome Medicine</i> , 2013, 5, 24.	8.2	53
44	American DNA. <i>Current Anthropology</i> , 2013, 54, S77-S86.	1.6	17
45	Reflections on the Cost of "Low-Cost" Whole Genome Sequencing: Framing the Health Policy Debate. <i>PLoS Biology</i> , 2013, 11, e1001699.	5.6	67
46	Race, Risk, and Recreation in Personal Genomics: The Limits of Play. <i>Medical Anthropology Quarterly</i> , 2013, 27, 550-569.	1.4	25
47	Attitudes towards Social Networking and Sharing Behaviors among Consumers of Direct-to-Consumer Personal Genomics. <i>Journal of Personalized Medicine</i> , 2013, 3, 275-287.	2.5	7
48	Lessons Learned From the U.S. Public Health Service Syphilis Study at Tuskegee: Incorporating a Discourse on Relationships Into the Ethics of Research Participation Among Asian Americans. <i>Ethics and Behavior</i> , 2012, 22, 489-492.	1.8	4
49	Assessing the Pedagogical Goals of Self-Testing in Evaluating the Consultation Needs of Different Student Populations. <i>American Journal of Bioethics</i> , 2012, 12, 41-43.	0.9	4
50	Research 2.0: Social Networking and Direct-To-Consumer (DTC) Genomics. <i>American Journal of Bioethics</i> , 2009, 9, 35-44.	0.9	62
51	The Illusive Gold Standard in Genetic Ancestry Testing. <i>Science</i> , 2009, 325, 38-39.	12.6	62
52	Response to Open Peer Commentaries on "Research 2.0: Social Networking and Direct-to-Consumer Personal Genomics" <i>American Journal of Bioethics</i> , 2009, 9, W1-W3.	0.9	1
53	Racing Forward: The Genomics and Personalized Medicine Act. <i>Science</i> , 2009, 323, 342-342.	12.6	27
54	The ethics of characterizing difference: guiding principles on using racial categories in human genetics. <i>Genome Biology</i> , 2008, 9, 404.	9.6	75

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55	Biobanks of a "racial kind": mining for difference in the new genetics. <i>Patterns of Prejudice</i> , 2006, 40, 443-460.	0.6	20
56	Racializing Drug Design: Implications of Pharmacogenomics for Health Disparities. <i>American Journal of Public Health</i> , 2005, 95, 2133-2138.	2.7	82
57	personalized medicine and pharmacogenomics: ethical and social challenges. <i>Personalized Medicine</i> , 2005, 2, 29-35.	1.5	6
58	Genetic Research and Health Disparities. <i>JAMA - Journal of the American Medical Association</i> , 2004, 291, 2985.	7.4	227
59	Race, Distributive Justice and the Promise of Pharmacogenomics. <i>Molecular Diagnosis and Therapy</i> , 2003, 3, 385-392.	3.3	37
60	Dys-appearing Tongues and Bodily Memories: The Aging of First-Generation Resident Koreans in Japan. <i>Ethos</i> , 2000, 28, 198-223.	0.2	20
61	Ethical Considerations for Pharmacogenomics: Privacy and Confidentiality. , 0, , 66-73.		1
62	Race and the science of difference in the age of genomics. , 0, , 26-39.		6