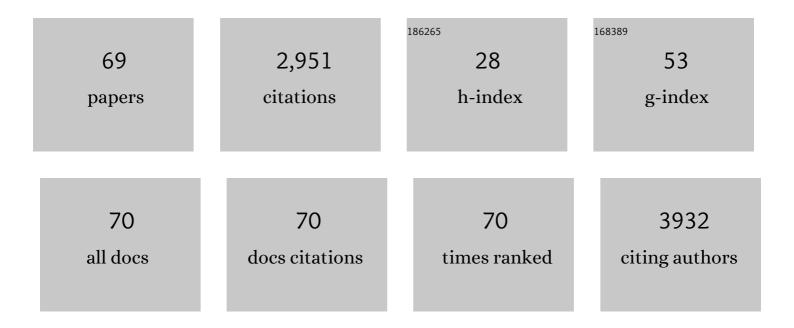
## Gail E Henderson

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4774306/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	International AIDS Society global scientific strategy: towards an HIV cure 2016. Nature Medicine, 2016, 22, 839-850.	30.7	395
2	Clinical Trials and Medical Care: Defining the Therapeutic Misconception. PLoS Medicine, 2007, 4, e324.	8.4	376
3	Recommendations for returning genomic incidental findings? We need to talk!. Genetics in Medicine, 2013, 15, 854-859.	2.4	272
4	HIV/AIDS Risk Among Brothel-Based Female Sex Workers in China: Assessing the Terms, Content, and Knowledge of Sex Work. Sexually Transmitted Diseases, 2004, 31, 695-700.	1.7	166
5	Characterizing biobank organizations in the U.S.: results from a national survey. Genome Medicine, 2013, 5, 3.	8.2	157
6	Clinical Sequencing Exploratory Research Consortium: Accelerating Evidence-Based Practice of Genomic Medicine. American Journal of Human Genetics, 2016, 98, 1051-1066.	6.2	137
7	Therapeutic misconception in early phase gene transfer trials. Social Science and Medicine, 2006, 62, 239-253.	3.8	113
8	Neglected ethical issues in biobank management: Results from a U.S. study. Life Sciences, Society and Policy, 2013, 9, 1.	3.2	77
9	Public Willingness to Participate in and Public Opinions About Genetic Variation Research: A Review of the Literature. American Journal of Public Health, 2006, 96, 1971-1978.	2.7	62
10	Is Informed Consent Broken?. American Journal of the Medical Sciences, 2011, 342, 267-272.	1.1	55
11	Framing expectations in early HIV cure research. Trends in Microbiology, 2014, 22, 547-549.	7.7	54
12	The promise and peril of genomic screening in the general population. Genetics in Medicine, 2016, 18, 593-599.	2.4	53
13	Ethics of treatment interruption trials in HIV cure research: addressing the conundrum of risk/benefit assessment. Journal of Medical Ethics, 2018, 44, medethics-2017-104433.	1.8	51
14	Great expectations: views of genetic research participants regarding current and future genetic studies. Genetics in Medicine, 2008, 10, 193-200.	2.4	49
15	Uncertain benefit: investigators' views and communications in early phase gene transfer trials. Molecular Therapy, 2004, 10, 225-231.	8.2	44
16	The Ethics of HIV "Cure―Research: What Can We Learn from Consent Forms?. AIDS Research and Human Retroviruses, 2015, 31, 56-63.	1.1	43
17	A comparison of respondent-driven and venue-based sampling of female sex workers in Liuzhou, China. Sexually Transmitted Infections, 2012, 88, i95-i101.	1.9	41
18	"Possibly positive or certainly uncertain?â€: participants' responses to uncertain diagnostic results from exome sequencing. Genetics in Medicine, 2018, 20, 313-319.	2.4	39

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19	Ethical considerations in global HIV phylogenetic research. Lancet HIV,the, 2018, 5, e656-e666.	4.7	39
20	Parents' perceptions of personal utility of exome sequencing results. Genetics in Medicine, 2020, 22, 752-757.	2.4	37
21	Indirect Benefits in HIV Cure Clinical Research: A Qualitative Analysis. AIDS Research and Human Retroviruses, 2019, 35, 100-107.	1.1	35
22	Stewardship Practices of U.S. Biobanks. Science Translational Medicine, 2013, 5, 215cm7.	12.4	34
23	What motivates participation in HIV cure trials? A call for real-time assessment to improve informed consent. Journal of Virus Eradication, 2015, 1, 51-53.	0.5	34
24	ls "incidental finding―the best term?: a study of patients' preferences. Genetics in Medicine, 2017, 19, 176-181.	2.4	34
25	Navigating the research–clinical interface in genomic medicine: analysis from the CSER Consortium. Genetics in Medicine, 2018, 20, 545-553.	2.4	34
26	Returning negative results to individuals in a genomic screening program: lessons learned. Genetics in Medicine, 2019, 21, 409-416.	2.4	33
27	Distributive justice, diversity, and inclusion in precision medicine: what will success look like?. Genetics in Medicine, 2017, 19, 157-159.	2.4	32
28	What motivates participation in HIV cure trials? A call for real-time assessment to improve informed consent. Journal of Virus Eradication, 2015, 1, 51-53.	0.5	31
29	Biobanks containing clinical specimens: Defining characteristics, policies, and practices. Clinical Biochemistry, 2014, 47, 245-251.	1.9	28
30	Social Contexts of Heterosexual Transmission of HIV/STI in Liuzhou City, China. AIDS and Behavior, 2014, 18, 111-117.	2.7	27
31	What Research Ethics Should Learn from Genomics and Society Research: Lessons from the ELSI Congress of 2011. Journal of Law, Medicine and Ethics, 2012, 40, 1008-1024.	0.9	24
32	Underutilization of specimens in biobanks: an ethical as well as a practical concern?. Genetics in Medicine, 2014, 16, 738-740.	2.4	23
33	Going off antiretroviral treatment in a closely monitored HIV "cure―trial: longitudinal assessments of acutely diagnosed trial participants and decliners. Journal of the International AIDS Society, 2019, 22, e25260.	3.0	23
34	Biobanks in the United States: How to Identify an Undefined and Rapidly Evolving Population. Biopreservation and Biobanking, 2012, 10, 511-517.	1.0	21
35	Asking the Right Questions: Views on Genetic Variation Research Among Black and White Research Participants. Journal of General Internal Medicine, 2009, 24, 299-304.	2.6	20
36	Genomic screening of the general adult population: key concepts for assessing net benefit with systematic evidence reviews. Genetics in Medicine, 2015, 17, 441-443.	2.4	19

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37	Vulnerability to Influence: A Two-Way Street. American Journal of Bioethics, 2004, 4, 50-52.	0.9	18
38	'I can coexist with HIV': a qualitative study of perceptions of HIV cure among people living with HIV in Guangzhou, China. Journal of Virus Eradication, 2016, 2, 170-4.	0.5	17
39	Online Education and e-Consent for GeneScreen, a Preventive Genomic Screening Study. Public Health Genomics, 2017, 20, 235-246.	1.0	15
40	Affected by HIV Stigma: Interpreting Results from a Population Survey of an Urban Center in Guangxi, China. AIDS and Behavior, 2014, 18, 192-201.	2.7	13
41	The Organization of Sex Work in Low- and High-Priced Venues with a Focus on the Experiences of Ethnic Minority Women Working in These Venues. AIDS and Behavior, 2014, 18, 172-180.	2.7	13
42	Population-Based Sexual Behavior Surveys in China: Liuzhou Compared with Other Prefectural Cities. AIDS and Behavior, 2014, 18, 118-125.	2.7	13
43	"Forward-Thinking―in U.S. Biobanking. Genetic Testing and Molecular Biomarkers, 2017, 21, 148-154.	0.7	12
44	Is Enhancement the Price of Prevention in Human Gene Editing?. CRISPR Journal, 2018, 1, 351-354.	2.9	12
45	How Biomedical HIV Prevention Trials Incorporate Behavioral and Social Sciences Research: A Typology of Approaches. AIDS and Behavior, 2019, 23, 2146-2154.	2.7	11
46	The Rise of Population Genomic Screening: Characteristics of Current Programs and the Need for Evidence Regarding Optimal Implementation. Journal of Personalized Medicine, 2022, 12, 692.	2.5	11
47	Why we should continue to worry about the therapeutic misconception. Journal of Clinical Ethics, 2013, 24, 381-6.	0.3	10
48	Is there evidence that we should screen the general population for Lynch syndrome with genetic testing? A systematic review. Pharmacogenomics and Personalized Medicine, 2017, Volume10, 49-60.	0.7	9
49	The Rise of Technology in Chinese Hospitals. International Journal of Technology Assessment in Health Care, 1987, 3, 253-263.	0.5	8
50	Examining the Cascade of Participant Attrition in a Genomic Medicine Research Study: Barriers and Facilitators to Achieving Diversity. Public Health Genomics, 2017, 20, 332-342.	1.0	8
51	Age and perceived risks and benefits of preventive genomic screening. Genetics in Medicine, 2018, 20, 1038-1044.	2.4	8
52	Traditional, complementary, and alternative medical cures for HIV: rationale and implications for HIV cure research. Global Public Health, 2019, 14, 152-160.	2.0	8
53	Psychological adaptation to diagnostic genomic sequencing results: The role of hope fulfillment Health Psychology, 2019, 38, 527-535.	1.6	8
54	Cohorts as collections of bodies and communities of persons: insights from the SEARCH010/RV254 research cohort. International Health, 2020, 12, 584-590.	2.0	6

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#	Article	IF	CITATIONS
55	Assessing the implications of positive genomic screening results. Personalized Medicine, 2020, 17, 101-109.	1.5	6
56	The View from the Benches: Scientists' Perspectives on the Uses and Governance of Human Gene-Editing Research. CRISPR Journal, 2021, 4, 609-615.	2.9	5
57	Public Comments on Proposed Regulatory Reforms That Would Impact Biospecimen Research: The Good, the Bad, and the Puzzling. IRB: Ethics & Human Research, 2015, 37, 1-10.	0.8	5
58	Recommendations from Thai stakeholders about protecting HIV remission (â€~cure') trial participants: report from a participatory workshop. International Health, 2020, 12, 567-574.	2.0	4
59	Attitudes About Analytic Treatment Interruption (ATI) in HIV Remission Trials with Different Antiretroviral Therapy (ART) Resumption Criteria. AIDS and Behavior, 2022, 26, 1504-1516.	2.7	4
60	Standardization as performative accountability in biobanking. BioSocieties, 2016, 11, 67-81.	1.3	2
61	Grudging Trust and the Limits of Trustworthy Biorepository Curation. American Journal of Bioethics, 2018, 18, 23-25.	0.9	2
62	Parallel but connected: Nuances of conducting behavioral and social science research alongside ethically challenging HIV remission trials. Contemporary Clinical Trials Communications, 2020, 19, 100594.	1.1	2
63	Beyond the Medical Model: Retooling Bioethics for the Work Ahead. American Journal of Bioethics, 2021, 21, 53-55.	0.9	2
64	Perceptions of HIV cure and willingness to participate in HIV cure-related trials among people enrolled in the Netherlands cohort study on acute HIV infection. Journal of Virus Eradication, 2022, 8, 100072.	0.5	2
65	Development and validation of a measure of comprehension of genomic screening—negative results (CoG-NR). European Journal of Human Genetics, 2020, 28, 1394-1402.	2.8	1
66	Ethical, legal and social implications of human genome studies in radiation research: a workshop report for studies on atomic bomb survivors at the Radiation Effects Research Foundation. Journal of Radiation Research, 2021, 62, 656-661.	1.6	1
67	Scientists' Views on Scientific Self-Governance for Human Genome Editing Research. Human Gene Therapy, 0, , .	2.7	1
68	With great (participant) rights comes great (researcher) responsibility. Genetics in Medicine, 2016, 18, 124-125.	2.4	0
69	Burden or benefit? Effects of providing education about and the option to request additional genomic findings from diagnostic exome sequencing: A randomized controlled trial. Patient Education and Counseling, 2021, 104, 2989-2998.	2.2	0