

# Susan Bull

## List of Publications by Year in descending order

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33  
papers

1,030  
citations

430442

18  
h-index

454577

30  
g-index

34  
all docs

34  
docs citations

34  
times ranked

1042  
citing authors

#	ARTICLE	IF	CITATIONS
1	Consent and Community Engagement in Diverse Research Contexts: Reviewing and Developing Research and Practice. <i>Journal of Empirical Research on Human Research Ethics</i> , 2013, 8, 1-18.	0.6	123
2	Seeking consent to genetic and genomic research in a rural Ghanaian setting: A qualitative study of the MalariaGEN experience. <i>BMC Medical Ethics</i> , 2012, 13, 15.	1.0	97
3	Impact of social stigma on the process of obtaining informed consent for genetic research on podoconiosis: a qualitative study. <i>BMC Medical Ethics</i> , 2009, 10, 13.	1.0	85
4	Views of Ethical Best Practices in Sharing Individual-Level Data From Medical and Public Health Research. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 225-238.	0.6	60
5	Best Practices for Ethical Sharing of Individual-Level Health Research Data From Low- and Middle-Income Settings. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 302-313.	0.6	60
6	Involving Research Stakeholders in Developing Policy on Sharing Public Health Research Data in Kenya. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 264-277.	0.6	46
7	Tailoring Information Provision and Consent Processes to Research Contexts: The Value of Rapid Assessments. <i>Journal of Empirical Research on Human Research Ethics</i> , 2012, 7, 37-52.	0.6	45
8	Perceived Benefits, Harms, and Views About How to Share Data Responsibly. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 278-289.	0.6	45
9	Developing Ethical Practices for Public Health Research Data Sharing in South Africa. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 290-301.	0.6	44
10	"It is an entrustment": Broad consent for genomic research and biobanks in sub-Saharan Africa. <i>Developing World Bioethics</i> , 2019, 19, 9-17.	0.6	42
11	Key criteria for the ethical acceptability of COVID-19 human challenge studies: Report of a WHO Working Group. <i>Vaccine</i> , 2021, 39, 633-640.	1.7	42
12	Research Stakeholders' Views on Benefits and Challenges for Public Health Research Data Sharing in Kenya: The Importance of Trust and Social Relations. <i>PLoS ONE</i> , 2015, 10, e0135545.	1.1	39
13	Ethical issues in the export, storage and reuse of human biological samples in biomedical research: perspectives of key stakeholders in Ghana and Kenya. <i>BMC Medical Ethics</i> , 2014, 15, 76.	1.0	37
14	Sweat, Skepticism, and Uncharted Territory. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 239-250.	0.6	37
15	Trust, Respect, and Reciprocity. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 251-263.	0.6	35
16	Sharing Public Health Research Data. <i>Journal of Empirical Research on Human Research Ethics</i> , 2015, 10, 217-224.	0.6	28
17	Understandings of genomic research in developing countries: a qualitative study of the views of MalariaGEN participants in Mali. <i>BMC Medical Ethics</i> , 2015, 16, 42.	1.0	23
18	Ensuring Consent to Research is Voluntary: How Far Do We Need to Go?. <i>American Journal of Bioethics</i> , 2011, 11, 27-29.	0.5	22

#	ARTICLE	IF	CITATIONS
19	Challenges arising when seeking broad consent for health research data sharing: a qualitative study of perspectives in Thailand. BMC Medical Ethics, 2018, 19, 86.	1.0	18
20	Ethics in collaborative global health research networks. Clinical Ethics, 2009, 4, 165-168.	0.5	17
21	SARS-CoV-2 challenge studies: ethics and risk minimisation. Journal of Medical Ethics, 2020, , medethics-2020-106504.	1.0	15
22	Equitable data sharing in epidemics and pandemics. BMC Medical Ethics, 2021, 22, 136.	1.0	10
23	Sharing Individual-Level Health Research Data: Experiences, Challenges and a Research Agenda. Asian Bioethics Review, 2017, 9, 393-400.	0.9	7
24	How should assent to research be sought in low income settings? Perspectives from parents and children in Southern Malawi. BMC Medical Ethics, 2019, 20, 32.	1.0	7
25	The ethics of data sharing and biobanking in health research. Wellcome Open Research, 2020, 5, 270.	0.9	7
26	Consensus standards for introductory e-learning courses in human participants research ethics. Journal of Medical Ethics, 2014, 40, 426-428.	1.0	6
27	Researcher and study participants' perspectives of consent in clinical studies in four referral hospitals in Vietnam. BMC Medical Ethics, 2020, 21, 4.	1.0	5
28	Vaccine-enhanced disease: case studies and ethical implications for research and public health. Wellcome Open Research, 2021, 6, 154.	0.9	4
29	"Are we getting the biometric bioethics right?" the use of biometrics within the healthcare system in Malawi. Global Bioethics, 2020, 31, 67-80.	0.5	3
30	Engaging publics in biobanking and genetic research governance - a literature review towards informing practice in India. Wellcome Open Research, 0, 6, 5.	0.9	2
31	Engaging publics in biobanking and genetic research governance - a literature review towards informing practice in India. Wellcome Open Research, 0, 6, 5.	0.9	1
32	Respecting values and perspectives in biobanking and genetic research governance: Outcomes of a qualitative study in Bengaluru, India. Wellcome Open Research, 0, 7, 78.	0.9	1
33	Ethics and Tropical Diseases. , 2014, , 31-39.e1.		0