## Juli M Bollinger

List of Publications by Year in descending order

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933264 839398 20 329 10 18 citations h-index g-index papers 21 21 21 613 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Public preferences regarding informed consent models for participation in population-based genomic research. Genetics in Medicine, $2014$ , $16$ , $11$ - $18$ .	1.1	74
2	Patients' Attitudes toward the Donation of Biological Materials for the Derivation of Induced Pluripotent Stem Cells. Cell Stem Cell, 2014, 14, 9-12.	5.2	33
3	Choices for return of primary and secondary genomic research results of 790 members of families with Mendelian disease. European Journal of Human Genetics, 2017, 25, 530-537.	1.4	31
4	Development of the clinical next-generation sequencing industry in a shifting policy climate. Nature Biotechnology, 2014, 32, 980-982.	9.4	25
5	Patients' views concerning research on medical practices: Implications for consent. AJOB Empirical Bioethics, 2016, 7, 76-91.	0.8	23
6	Patients' perspectives on the derivation and use of organoids. Stem Cell Reports, 2021, 16, 1874-1883.	2.3	23
7	Enrolling Genomics Research Participants through a Clinical Setting: the Impact of Existing Clinical Relationships on Informed Consent and Expectations for Return of Research Results. Journal of Genetic Counseling, 2018, 27, 263-273.	0.9	19
8	Ethics and Collateral Findings in Pragmatic Clinical Trials. American Journal of Bioethics, 2020, 20, 6-18.	0.5	16
9	Barriers to clinical adoption of next-generation sequencing: a policy Delphi panel's solutions. Personalized Medicine, 2017, 14, 339-354.	0.8	15
10	Physicians' perspectives regarding pragmatic clinical trials. Journal of Comparative Effectiveness Research, 2016, 5, 499-506.	0.6	12
11	Risk perception before and after presymptomatic genetic testing for Huntington's disease: Not always what one might expect. Molecular Genetics & Enomic Medicine, 2018, 6, 1140-1147.	0.6	10
12	Perceptions, motivations, and concerns about living organ donation among people living with HIV. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2018, 30, 1595-1599.	0.6	10
13	Patients' Views About the Disclosure of Collateral Findings in Pragmatic Clinical Trials: a Focus Group Study. Journal of General Internal Medicine, 2020, 35, 3436-3442.	1.3	7
14	Whether, when, and how to communicate genetic risk to minors: â€~I wanted more information but I think they were scared I couldn't handle it'. Journal of Genetic Counseling, 2021, 30, 237-245.	0.9	7
15	Family Communication Patterns and Challenges of Huntington's Disease Risk, the Decision to Pursue Presymptomatic Testing, and Test Results. Journal of Huntington's Disease, 2020, 9, 265-274.	0.9	6
16	Developing context-specific next-generation sequencing policy. Nature Biotechnology, 2016, 34, 466-470.	9.4	5
17	Stakeholder perspectives regarding pragmatic clinical trial collateral findings. Learning Health Systems, 2020, 5, e10245.	1.1	5
18	Patients' Reactions to Letters Communicating Collateral Findings of Pragmatic Clinical Trials: a National Web-Based Survey. Journal of General Internal Medicine, 2022, 37, 1658-1664.	1.3	4

#	Article	IF	CITATIONS
19	Early experiences of independent advocates for potential HIV+ recipients of HIV+ donor organ transplants. Clinical Transplantation, 2019, 33, e13617.	0.8	3
20	Identification and management of pragmatic clinical trial collateral findings: A current understanding and directions for future research. Healthcare, 2021, 9, 100586.	0.6	1