John Wilbanks

List of Publications by Year
in descending order

		623574	610775
28 papers	1,836 citations	14 h-index	24 g-index
30 all docs	30 docs citations	30 times ranked	3461 citing authors

#	Article	IF	Citations
1	The mPower study, Parkinson disease mobile data collected using ResearchKit. Scientific Data, 2016, 3, 160011.	2.4	439
2	The National COVID Cohort Collaborative (N3C): Rationale, design, infrastructure, and deployment. Journal of the American Medical Informatics Association: JAMIA, 2021, 28, 427-443.	2.2	342
3	Indicators of retention in remote digital health studies: a cross-study evaluation of 100,000 participants. Npj Digital Medicine, 2020, 3, 21.	5.7	238
4	The Rise of Citizen Science in Health and Biomedical Research. American Journal of Bioethics, 2019, 19, 3-14.	0.5	170
5	'Omics Data Sharing. Science, 2009, 326, 234-236.	6.0	136
6	Remote smartphone monitoring of Parkinson's disease and individual response to therapy. Nature Biotechnology, 2022, 40, 480-487.	9.4	73
7	Formative Evaluation of Participant Experience With Mobile eConsent in the App-Mediated Parkinson mPower Study: A Mixed Methods Study. JMIR MHealth and UHealth, 2017, 5, e14.	1.8	56
8	Design Issues in E-Consent. Journal of Law, Medicine and Ethics, 2018, 46, 110-118.	0.4	53
9	Stop the privatization of health data. Nature, 2016, 535, 345-348.	13.7	51
10	First, design for data sharing. Nature Biotechnology, 2016, 34, 377-379.	9.4	51
11	Responsible sharing of biomedical data and biospecimens via the "Automatable Discovery and Access Matrix"(ADA-M). Npj Genomic Medicine, 2018, 3, 17.	1.7	38
12	An open source pharma roadmap. PLoS Medicine, 2017, 14, e1002276.	3.9	26
13	Another reason for opening access to research. BMJ: British Medical Journal, 2006, 333, 1306-1308.	2.4	24
14	What the semantic web could do for the life sciences. Drug Discovery Today Biosilico, 2004, 2, 228-236.	0.7	22
15	Developing a Transparent, Participant-Navigated Electronic Informed Consent for Mobile-Mediated Research. SSRN Electronic Journal, 0, , .	0.4	18
16	Why Open Drug Discovery Needs Four Simple Rules for Licensing Data and Models. PLoS Computational Biology, 2012, 8, e1002706.	1.5	17
17	Assessment of the *All of Us* research program's informed consent process. AJOB Empirical Bioethics, 2021, 12, 72-83.	0.8	17
18	Empirical validation of an automated approach to data use oversight. Cell Genomics, 2021, 1, 100031.	3.0	17

#	Article	IF	Citations
19	Bringing Code to Data: Do Not Forget Governance. Journal of Medical Internet Research, 2020, 22, e18087.	2.1	11
20	The Role of Government in Precision Medicine, Precision Public Health and the Intersection With Healthy Living. Progress in Cardiovascular Diseases, 2019, 62, 50-54.	1.6	9
21	Portable Approaches to Informed Consent and Open Data. , 2014, , 234-252.		6
22	Openness as infrastructure. Journal of Cheminformatics, 2011, 3, 36.	2.8	5
23	A fool's errand. Nature, 2013, 495, 440-441.	13.7	5
24	Implementing a universal informed consent process for the Research Program. Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing, 2019, 24, 427-438.	0.7	5
25	Ethical Issues in Consumer Informatics and Online Content. , 2019, , 327-336.		3
26	An Interoperability Principle for Knowledge Creation and Governance: The Role of Emerging Institutions. , 0, , 199-226.		2
27	We need a Web for data. Learned Publishing, 2010, 23, 333-335.	0.8	1
28	Endoscopic features for early decision to evaluate superior mesenteric artery syndrome in children. BMC Pediatrics, 2021, 21, 392.	0.7	1