## John Wilbanks

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4554697/publications.pdf

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

#	Article	IF	CITATIONS
1	Remote smartphone monitoring of Parkinson's disease and individual response to therapy. Nature Biotechnology, 2022, 40, 480-487.	9.4	73
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	Assessment of the <i>All of Us</i> research program's informed consent process. AJOB Empirical Bioethics, 2021, 12, 72-83.	0.8	17
4	Endoscopic features for early decision to evaluate superior mesenteric artery syndrome in children. BMC Pediatrics, 2021, 21, 392.	0.7	1
5	Empirical validation of an automated approach to data use oversight. Cell Genomics, 2021, 1, 100031.	3.0	17
6	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
7	Bringing Code to Data: Do Not Forget Governance. Journal of Medical Internet Research, 2020, 22, e18087.	2.1	11
8	The Rise of Citizen Science in Health and Biomedical Research. American Journal of Bioethics, 2019, 19, 3-14.	0.5	170
9	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
10	Ethical Issues in Consumer Informatics and Online Content., 2019,, 327-336.		3
11	Implementing a universal informed consent process for the Research Program. Pacific Symposium on Biocomputing, 2019, 24, 427-438.	0.7	5
12	Design Issues in E-Consent. Journal of Law, Medicine and Ethics, 2018, 46, 110-118.	0.4	53
13	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
14	An open source pharma roadmap. PLoS Medicine, 2017, 14, e1002276.	3.9	26
15	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
16	The mPower study, Parkinson disease mobile data collected using ResearchKit. Scientific Data, 2016, 3, 160011.	2.4	439
17	Stop the privatization of health data. Nature, 2016, 535, 345-348.	13.7	51
18	First, design for data sharing. Nature Biotechnology, 2016, 34, 377-379.	9.4	51

#	Article	IF	CITATIONS
19	Portable Approaches to Informed Consent and Open Data., 2014,, 234-252.		6
20	A fool's errand. Nature, 2013, 495, 440-441.	13.7	5
21	Why Open Drug Discovery Needs Four Simple Rules for Licensing Data and Models. PLoS Computational Biology, 2012, 8, e1002706.	1.5	17
22	Openness as infrastructure. Journal of Cheminformatics, 2011, 3, 36.	2.8	5
23	We need a Web for data. Learned Publishing, 2010, 23, 333-335.	0.8	1
24	'Omics Data Sharing. Science, 2009, 326, 234-236.	6.0	136
25	Another reason for opening access to research. BMJ: British Medical Journal, 2006, 333, 1306-1308.	2.4	24
26	What the semantic web could do for the life sciences. Drug Discovery Today Biosilico, 2004, 2, 228-236.	0.7	22
27	An Interoperability Principle for Knowledge Creation and Governance: The Role of Emerging Institutions. , 0, , 199-226.		2
28	Developing a Transparent, Participant-Navigated Electronic Informed Consent for Mobile-Mediated Research. SSRN Electronic Journal, 0, , .	0.4	18