

Jodyn E Platt

List of Publications by Year in descending order

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Version: 2024-02-01

43
papers

800
citations

623734

14
h-index

552781

26
g-index

44
all docs

44
docs citations

44
times ranked

983
citing authors

#	ARTICLE	IF	CITATIONS
1	Patient-Reported Experiences of Discrimination in the US Health Care System. <i>JAMA Network Open</i> , 2020, 3, e2029650.	5.9	101
2	Public preferences regarding informed consent models for participation in population-based genomic research. <i>Genetics in Medicine</i> , 2014, 16, 11-18.	2.4	74
3	An Analysis of the Learning Health System in Its First Decade in Practice: Scoping Review. <i>Journal of Medical Internet Research</i> , 2020, 22, e17026.	4.3	74
4	Public Trust in Health Information Sharing: Implications for Biobanking and Electronic Health Record Systems. <i>Journal of Personalized Medicine</i> , 2015, 5, 3-21.	2.5	65
5	Facebook Advertising Across an Engagement Spectrum: A Case Example for Public Health Communication. <i>JMIR Public Health and Surveillance</i> , 2016, 2, e27.	2.6	53
6	Public Trust in Health Information Sharing: A Measure of System Trust. <i>Health Services Research</i> , 2018, 53, 824-845.	2.0	52
7	Measuring physicians' trust: A scoping review with implications for public policy. <i>Social Science and Medicine</i> , 2016, 165, 75-81.	3.8	49
8	Testing an Online, Dynamic Consent Portal for Large Population Biobank Research. <i>Public Health Genomics</i> , 2015, 18, 26-39.	1.0	41
9	Community perspectives on public health biobanking: an analysis of community meetings on the Michigan BioTrust for Health. <i>Journal of Community Genetics</i> , 2014, 5, 125-138.	1.2	33
10	Encouraging Participation And Transparency In Biobank Research. <i>Health Affairs</i> , 2018, 37, 1313-1320.	5.2	30
11	The public's comfort with sharing health data with third-party commercial companies. <i>Humanities and Social Sciences Communications</i> , 2020, 7, .	2.9	30
12	Building and maintaining trust in clinical decision support: Recommendations from the Patient-Centered CDS Learning Network. <i>Learning Health Systems</i> , 2020, 4, e10208.	2.0	19
13	'Born in Michigan? You're in the Biobank': Engaging Population Biobank Participants through Facebook Advertisements. <i>Public Health Genomics</i> , 2013, 16, 145-158.	1.0	18
14	The public's trust and information brokers in health care, public health and research. <i>Journal of Health Organization and Management</i> , 2019, 33, 929-948.	1.3	18
15	'Cool! and creepy': engaging with college student stakeholders in Michigan's biobank. <i>Journal of Community Genetics</i> , 2014, 5, 349-362.	1.2	16
16	Discrimination, trust, and withholding information from providers: Implications for missing data and inequity. <i>SSM - Population Health</i> , 2022, 18, 101092.	2.7	14
17	Family Caregiver Access of Online Medical Records: Findings from the Health Information National Trends Survey. <i>Journal of General Internal Medicine</i> , 2020, 36, 3267-3269.	2.6	13
18	Family Caregivers' Experiences With Telehealth During COVID-19: Insights From Michigan. <i>Annals of Family Medicine</i> , 2022, 20, 69-71.	1.9	13

#	ARTICLE	IF	CITATIONS
19	Willingness to Participate in Health Information Networks with Diverse Data Use: Evaluating Public Perspectives. EGEMS (Washington, DC), 2019, 7, 33.	2.0	11
20	Do people have an ethical obligation to share their health information? Comparing narratives of altruism and health information sharing in a nationally representative sample. PLoS ONE, 2020, 15, e0244767.	2.5	9
21	Understanding racial differences in attitudes about public health efforts during COVID-19 using an explanatory mixed methods design. Social Science and Medicine, 2021, 287, 114379.	3.8	8
22	Dynamics of Physicians' Trust in Fellow Health Care Providers and the Role of Health Information Technology. Medical Care Research and Review, 2021, 78, 338-349.	2.1	7
23	Rethinking ethical oversight in the era of the learning health system. Healthcare, 2020, 8, 100462.	1.3	7
24	Exploring How Personal, Social, and Institutional Characteristics Contribute to Geriatric Medicine Subspecialty Decisions: A Qualitative Study of Trainees' Perceptions. Academic Medicine, 2021, 96, 425-432.	1.6	7
25	Engaging a state: Facebook comments on a large population biobank. Journal of Community Genetics, 2017, 8, 183-197.	1.2	6
26	A qualitative exploration of the informed consent process in hematopoietic cell transplantation clinical research and opportunities for improvement. Bone Marrow Transplantation, 2017, 52, 292-298.	2.4	6
27	Improving the Informed Consent Process in Hematopoietic Cell Transplantation: Patient, Caregiver, and Provider Perspectives. Biology of Blood and Marrow Transplantation, 2018, 24, 156-162.	2.0	6
28	Ethical, legal, and social implications of learning health systems. Learning Health Systems, 2018, 2, e10051.	2.0	5
29	Summary of second annual MCBK public meeting: Mobilizing Computable Biomedical Knowledge—A movement to accelerate translation of knowledge into action. Learning Health Systems, 2020, 4, e10222.	2.0	4
30	What Does "Patient-Centered" Mean? Qualitative Perspectives from Older Adults and Family Caregivers. Gerontology and Geriatric Medicine, 2021, 7, 233372142110176.	1.5	3
31	Trust in provider care teams and health information technology-mediated communication. American Journal of Managed Care, 2020, 26, 23-25.	1.1	2
32	Summary of fourth annual MCBK public meeting: Mobilizing computable biomedical knowledge—metadata and trust. Learning Health Systems, 2022, 6, e10301.	2.0	2
33	Tell it Like it Seems: Challenges Identifying Potential Requirements of a Learning Health System. , 2015, , .		1
34	Innovating consent for pediatric HCT patients. Bone Marrow Transplantation, 2016, 51, 885-888.	2.4	1
35	"If It Needs to be Done, It Needs to be Done": National Survey of Youth Experiences and Perspectives on Caregiving. Journal of Adolescent Health, 2021, 69, 664-667.	2.5	1
36	Summary of third annual MCBK public meeting: Mobilizing computable biomedical knowledge—Accelerating the second knowledge revolution. Learning Health Systems, 2021, 5, e10255.	2.0	1

#	ARTICLE	IF	CITATIONS
37	IDENTITY AND RESPONSIBILITY: WHAT IT MEANS TO BE A CAREGIVER AND ITS IMPLICATIONS FOR POLICY AND HEALTH CARE. Innovation in Aging, 2019, 3, S137-S137.	0.1	0
38	Analysis of Pioneering Computable Biomedical Knowledge Repositories and their Emerging Governance Structures. Data Intelligence, 0, , 1-26.	1.5	0
39	Title is missing!. , 2020, 15, e0244767.		0
40	Title is missing!. , 2020, 15, e0244767.		0
41	Title is missing!. , 2020, 15, e0244767.		0
42	Title is missing!. , 2020, 15, e0244767.		0
43	A survey of computable biomedical knowledge repositories. Learning Health Systems, 0, , .	2.0	0