

Jody E Platt

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9528179/publications.pdf>

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	Family Caregiversâ€™ Experiences With Telehealth During COVID-19: Insights From Michigan. <i>Annals of Family Medicine</i> , 2022, 20, 69-71.	1.9	13
2	Discrimination, trust, and withholding information from providers: Implications for missing data and inequity. <i>SSM - Population Health</i> , 2022, 18, 101092.	2.7	14
3	Summary of fourth annual <scp>MCBK</scp> public meeting: Mobilizing computable biomedical knowledgeâ€™ metadata and trust. <i>Learning Health Systems</i> , 2022, 6, e10301.	2.0	2
4	Dynamics of Physiciansâ€™ Trust in Fellow Health Care Providers and the Role of Health Information Technology. <i>Medical Care Research and Review</i> , 2021, 78, 338-349.	2.1	7
5	What Does â€œPatient-Centeredâ€ Mean? Qualitative Perspectives from Older Adults and Family Caregivers. <i>Gerontology and Geriatric Medicine</i> , 2021, 7, 233372142110176.	1.5	3
6	Understanding racial differences in attitudes about public health efforts during COVID-19 using an explanatory mixed methods design. <i>Social Science and Medicine</i> , 2021, 287, 114379.	3.8	8
7	â€œIf It Needs to be Done, It Needs to be Doneâ€: National Survey of Youth Experiences and Perspectives on Caregiving. <i>Journal of Adolescent Health</i> , 2021, 69, 664-667.	2.5	1
8	Summary of third annual MCBK public meeting: Mobilizing computable biomedical knowledgeâ€™ Accelerating the second knowledge revolution. <i>Learning Health Systems</i> , 2021, 5, e10255.	2.0	1
9	Exploring How Personal, Social, and Institutional Characteristics Contribute to Geriatric Medicine Subspecialty Decisions: A Qualitative Study of Traineesâ€™ Perceptions. <i>Academic Medicine</i> , 2021, 96, 425-432.	1.6	7
10	Rethinking ethical oversight in the era of the learning health system. <i>Healthcare</i> , 2020, 8, 100462.	1.3	7
11	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
12	Building and maintaining trust in clinical decision support: Recommendations from the Patientâ€™Centered CDS Learning Network. <i>Learning Health Systems</i> , 2020, 4, e10208.	2.0	19
13	Summary of second annual MCBK public meeting: Mobilizing Computable Biomedical Knowledgeâ€™ A movement to accelerate translation of knowledge into action. <i>Learning Health Systems</i> , 2020, 4, e10222.	2.0	4
14	Patient-Reported Experiences of Discrimination in the US Health Care System. <i>JAMA Network Open</i> , 2020, 3, e2029650.	5.9	101
15	The publicâ€™s comfort with sharing health data with third-party commercial companies. <i>Humanities and Social Sciences Communications</i> , 2020, 7, .	2.9	30
16	Do people have an ethical obligation to share their health information? Comparing narratives of altruism and health information sharing in a nationally representative sample. <i>PLoS ONE</i> , 2020, 15, e0244767.	2.5	9
17	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
18	Trust in provider care teams and health information technologyâ€™ mediated communication. <i>American Journal of Managed Care</i> , 2020, 26, 23-25.	1.1	2

#	ARTICLE	IF	CITATIONS
19	Title is missing!. , 2020, 15, e0244767.		0
20	Title is missing!. , 2020, 15, e0244767.		0
21	Title is missing!. , 2020, 15, e0244767.		0
22	Title is missing!. , 2020, 15, e0244767.		0
23	The public's trust and information brokers in health care, public health and research. Journal of Health Organization and Management, 2019, 33, 929-948.	1.3	18
24	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
25	Willingness to Participate in Health Information Networks with Diverse Data Use: Evaluating Public Perspectives. EGEMS (Washington, DC), 2019, 7, 33.	2.0	11
26	Ethical, legal, and social implications of learning health systems. Learning Health Systems, 2018, 2, e10051.	2.0	5
27	Public Trust in Health Information Sharing: A Measure of System Trust. Health Services Research, 2018, 53, 824-845.	2.0	52
28	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
29	Encouraging Participation And Transparency In Biobank Research. Health Affairs, 2018, 37, 1313-1320.	5.2	30
30	Engaging a state: Facebook comments on a large population biobank. Journal of Community Genetics, 2017, 8, 183-197.	1.2	6
31	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
32	Measuring physicians' trust: A scoping review with implications for public policy. Social Science and Medicine, 2016, 165, 75-81.	3.8	49
33	Innovating consent for pediatric HCT patients. Bone Marrow Transplantation, 2016, 51, 885-888.	2.4	1
34	Facebook Advertising Across an Engagement Spectrum: A Case Example for Public Health Communication. JMIR Public Health and Surveillance, 2016, 2, e27.	2.6	53
35	Public Trust in Health Information Sharing: Implications for Biobanking and Electronic Health Record Systems. Journal of Personalized Medicine, 2015, 5, 3-21.	2.5	65
36	Tell it Like it Seems: Challenges Identifying Potential Requirements of a Learning Health System. , 2015, , .		1

#	ARTICLE	IF	CITATIONS
37	Testing an Online, Dynamic Consent Portal for Large Population Biobank Research. Public Health Genomics, 2015, 18, 26-39.	1.0	41
38	Public preferences regarding informed consent models for participation in population-based genomic research. Genetics in Medicine, 2014, 16, 11-18.	2.4	74
39	Community perspectives on public health biobanking: an analysis of community meetings on the Michigan BioTrust for Health. Journal of Community Genetics, 2014, 5, 125-138.	1.2	33
40	â€˜Cool! and creepyâ€™: engaging with college student stakeholders in Michiganâ€™s biobank. Journal of Community Genetics, 2014, 5, 349-362.	1.2	16
41	â€˜Born in Michigan? You're in the Biobank': Engaging Population Biobank Participants through Facebook Advertisements. Public Health Genomics, 2013, 16, 145-158.	1.0	18
42	Analysis of Pioneering Computable Biomedical Knowledge Repositories and their Emerging Governance Structures. Data Intelligence, 0, , 1-26.	1.5	0
43	A survey of computable biomedical knowledge repositories. Learning Health Systems, 0, , .	2.0	0