

Eric M Meslin

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

2,227
citations

201674

27
h-index

233421

45
g-index

63
all docs

63
docs citations

63
times ranked

2721
citing authors

#	ARTICLE	IF	CITATIONS
1	Toward better governance of human genomic data. <i>Nature Genetics</i> , 2021, 53, 2-8.	21.4	31
2	Don Chalmers: His Contributions to Legal Research and Education, Health Law, and Research Ethics, Locally and Globally. <i>Journal of Law & Medicine</i> , 2020, 28, 289-297.	0.0	0
3	Sharing with Strangers: Governance Models for Borderless Genomic Research in a Territorial World. <i>Kennedy Institute of Ethics Journal</i> , 2019, 29, 67-95.	0.5	9
4	Women's opinions of legal requirements for drug testing in prenatal care. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2017, 30, 1693-1698.	1.5	4
5	Ethics and Childhood Vaccination Policy in the United States. <i>American Journal of Public Health</i> , 2016, 106, 273-278.	2.7	59
6	To research (or not) that is the question: ethical issues in research when medical care is disrupted by political action: a case study from Eldoret, Kenya. <i>Journal of Medical Ethics</i> , 2016, 42, 61-65.	1.8	6
7	Ethics review for international data-intensive research. <i>Science</i> , 2016, 351, 1399-1400.	12.6	44
8	Industry Support of Medical Research: Important Opportunity or Treacherous Pitfall?. <i>Journal of General Internal Medicine</i> , 2016, 31, 228-233.	2.6	31
9	Benchmarks for ethically credible partnerships between industry and academic health centers: beyond disclosure of financial conflicts of interest. <i>Clinical and Translational Medicine</i> , 2015, 4, 36.	4.0	15
10	How Bioethics Principles Can Aid Design of Electronic Health Records to Accommodate Patient Granular Control. <i>Journal of General Internal Medicine</i> , 2015, 30, 3-6.	2.6	19
11	Designing a Patient-Centered User Interface for Access Decisions about EHR Data: Implications from Patient Interviews. <i>Journal of General Internal Medicine</i> , 2015, 30, 7-16.	2.6	48
12	Provider Responses to Patients Controlling Access to their Electronic Health Records: A Prospective Cohort Study in Primary Care. <i>Journal of General Internal Medicine</i> , 2015, 30, 31-37.	2.6	35
13	Patient Preferences in Controlling Access to Their Electronic Health Records: a Prospective Cohort Study in Primary Care. <i>Journal of General Internal Medicine</i> , 2015, 30, 25-30.	2.6	55
14	"Because It Was Hard" Some Lessons Developing a Joint IRB Between Moi University (Kenya) and Indiana University (USA). <i>American Journal of Bioethics</i> , 2014, 14, 17-19.	0.9	4
15	Research Ethics Capacity Building in Sub-Saharan Africa: A Review of NIH Fogarty-Funded Programs 2000-2012. <i>Journal of Empirical Research on Human Research Ethics</i> , 2014, 9, 24-40.	1.3	57
16	Building a data sharing model for global genomic research. <i>Genome Biology</i> , 2014, 15, 430.	8.8	37
17	Ethics, Information Technology, and Public Health: Duties and Challenges in Computational Epidemiology. <i>Computers in Health Care</i> , 2014, , 191-209.	0.3	11
18	Mapping the translational science policy "valley of death". <i>Clinical and Translational Medicine</i> , 2013, 2, 14.	4.0	58

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19	Giving patients granular control of personal health information: Using an ethics "Points to Consider"™ to inform informatics system designers. <i>International Journal of Medical Informatics</i> , 2013, 82, 1136-1143.	3.3	34
20	Attitudes About the Use of Newborn Dried Blood Spots for Research: A Survey of Underrepresented Parents. <i>Academic Pediatrics</i> , 2013, 13, 451-457.	2.0	17
21	Taking Stock of the Ethical Foundations of International Health Research: Pragmatic Lessons from the IUAE™/Moi Academic Research Ethics Partnership. <i>Journal of General Internal Medicine</i> , 2013, 28, 639-645.	2.6	7
22	A Bibliometric Analysis of an International Research Ethics Trainee Program. <i>Journal of Empirical Research on Human Research Ethics</i> , 2013, 8, 75-81.	1.3	7
23	ELSI 2.0 for Genomics and Society. <i>Science</i> , 2012, 336, 673-674.	12.6	39
24	A Lifecycle Approach to the Evaluation of FDA Approval Methods and Regulatory Actions. <i>JAMA - Journal of the American Medical Association</i> , 2012, 307, 2491-2.	7.4	34
25	Perspective. <i>Academic Medicine</i> , 2012, 87, 1165-1170.	1.6	58
26	Ethical Issues in a Stage 1 Cognitive-Behavioral Therapy Feasibility Study and Trial to Reduce Alcohol Use Among HIV-Infected Outpatients in Western Kenya. <i>Journal of Empirical Research on Human Research Ethics</i> , 2012, 7, 29-37.	1.3	21
27	Deflating the Genomic Bubble. <i>Science</i> , 2011, 331, 861-862.	12.6	161
28	Biobanking and public health: is a human rights approach the tie that binds?. <i>Human Genetics</i> , 2011, 130, 451-63.	3.8	19
29	Ethics and Privacy Considerations for Systems Biology Applications in Predictive and Personalized Medicine. , 2011, , 1-27.		2
30	Ethics of implementing Electronic Health Records in developing countries: points to consider. <i>AMIA ... Annual Symposium proceedings</i> , 2011, 2011, 1499-505.	0.2	12
31	The Value of Using Top-Down and Bottom-Up Approaches for Building Trust and Transparency in Biobanking. <i>Public Health Genomics</i> , 2010, 13, 207-214.	1.0	33
32	Research Ethics in the Era of Personalized Medicine: Updating Science's Contract with Society. <i>Public Health Genomics</i> , 2010, 13, 378-384.	1.0	34
33	Extending the reach of public health genomics: What should be the agenda for public health in an era of genome-based and "personalized" medicine?. <i>Genetics in Medicine</i> , 2010, 12, 785-791.	2.4	95
34	Pediatric Assent for a Study of Antiretroviral Therapy Dosing for Children in Western Kenya: A Case Study in International Research Collaboration. <i>Journal of Empirical Research on Human Research Ethics</i> , 2009, 4, 3-16.	1.3	15
35	Ethical Issues in Measuring Biomarkers in Children's Environmental Health. <i>Environmental Health Perspectives</i> , 2009, 117, 1185-1190.	6.0	17
36	Pharmacotherapy and Pregnancy. <i>Clinical and Translational Science</i> , 2009, 2, 11-14.	3.1	6

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37	Helsinki discords: FDA, ethics, and international drug trials. <i>Lancet</i> , The, 2009, 373, 13-14.	13.7	55
38	The Ethics of Information: Absolute Risk Reduction and Patient Understanding of Screening. <i>Journal of General Internal Medicine</i> , 2008, 23, 867-870.	2.6	30
39	Health-Related Philanthropy: Toward Understanding the Relationship Between the Donation of the Body (and Its Parts) and Traditional Forms of Philanthropic Giving. <i>Nonprofit and Voluntary Sector Quarterly</i> , 2008, 37, 44S-62S.	1.9	12
40	Research Ethics Recommendations for Whole-Genome Research: Consensus Statement. <i>PLoS Biology</i> , 2008, 6, e73.	5.6	212
41	Patient Attitudes Toward Genotyping in an Urban Women's Health Clinic. <i>Obstetrics and Gynecology</i> , 2008, 112, 1023-1028.	2.4	6
42	Cancer Patients' Attitudes toward Future Research Uses of Stored Human Biological Materials. <i>Journal of Empirical Research on Human Research Ethics</i> , 2007, 2, 15-22.	1.3	46
43	Challenging a Well Established Consent Norm?: One Time Consent for Biobank Research. <i>Journal of International Biotechnology Law</i> , 2007, 4, .	0.1	13
44	Australia needs a better system for health care evaluation. <i>Medical Journal of Australia</i> , 2007, 186, 220-221.	1.7	12
45	Encouraging Translational Research Through Harmonization of FDA and Common Rule Informed Consent Requirements for Research with Banked Specimens. <i>Journal of Legal Medicine</i> , 2006, 27, 119-166.	0.5	22
46	Shifting paradigms in health services research ethics consent, privacy, and the challenges for IRBs. <i>Journal of General Internal Medicine</i> , 2006, 21, 279-280.	2.6	14
47	A Needs Assessment to Build International Research Ethics Capacity. <i>Journal of Empirical Research on Human Research Ethics</i> , 2006, 1, 23-38.	1.3	20
48	Medical Ethics Education: Where Are We? Where Should We Be Going? A Review. <i>Academic Medicine</i> , 2005, 80, 1143-1152.	1.6	264
49	Disclosure of Genetic Information Obtained Through Research. <i>Genetic Testing and Molecular Biomarkers</i> , 2004, 8, 347-355.	1.7	27
50	Creating incentives for genomic research to improve targeting of therapies. <i>Nature Medicine</i> , 2004, 10, 1289-1291.	30.7	14
51	Ethical issues in the collection, storage, and research use of human biological materials. <i>Translational Research</i> , 2004, 144, 229-234.	2.3	48
52	Faculty Ownership of Medical Facilities: Inappropriate Conflict or an Opportunity that Benefits Physicians and Patients?. <i>Academic Medicine</i> , 2004, 79, 1051-1055.	1.6	5
53	When Policy Analysis Is Carried Out in Public. , 2003, , 87-111.		0
54	Raising the bar in research ethics. <i>Postgraduate Medicine</i> , 2002, 112, 5-112.	2.0	2

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55	Some Initial Reflections on NBAC. Kennedy Institute of Ethics Journal, 2002, 12, 95-102.	0.5	8
56	Ethical Issues in the Design and Conduct of Clinical Trials in Developing Countries. New England Journal of Medicine, 2001, 345, 139-142.	27.0	172
57	Of Clones, Stem Cells, and Children: Issues and Challenges in Human Research Ethics. Journal of Women's Health and Gender-Based Medicine, 2000, 9, 831-841.	1.5	6
58	Engaging the public in policy development: The national bioethics advisory commission report on <i>research involving persons with mental disorders that may affect decisionmaking capacity</i>. Accountability in Research, 1999, 7, 227-239.	2.4	21
59	The Ethical, Legal, and Social Implications Research Program at the National Human Genome Research Institute. Kennedy Institute of Ethics Journal, 1997, 7, 291-298.	0.5	57
60	United States and Canadian approaches to justice in health care: A comparative analysis of health care systems and values. Theoretical Medicine and Bioethics, 1994, 15, 181-200.	0.5	14
61	The Moral Costs of the Ontario Physicians' Strike. Hastings Center Report, 1987, 17, 11.	1.0	12
62	Ethics and Privacy Considerations for Systems Biology Applications in Predictive and Personalized Medicine. , 0, , 1378-1404.		1