

Mark Yarborough

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

447
citations

758635

12
h-index

752256

20
g-index

37
all docs

37
docs citations

37
times ranked

454
citing authors

#	ARTICLE	IF	CITATIONS
1	Using the concept of “deserved trust” to strengthen the value and integrity of biomedical research. <i>Accountability in Research</i> , 2021, 28, 456-469.	1.6	4
2	Do we really know how many clinical trials are conducted ethically? Why research ethics committee review practices need to be strengthened and initial steps we could take to strengthen them. <i>Journal of Medical Ethics</i> , 2021, 47, 572-579.	1.0	9
3	Moving towards less biased research. <i>BMJ Open Science</i> , 2021, 5, e100116.	0.8	4
4	Rescuing Informed Consent: How the new “Key Information” and “Reasonable Person” Provisions in the Revised U.S. Common Rule open the door to long Overdue Informed Consent Disclosure Improvements and why we need to walk Through that door. <i>Science and Engineering Ethics</i> , 2020, 26, 1423-1443.	1.7	6
5	Bioethics Emergencies Can Be Used to Perform a Real-World Test of Utilitarian Policies. <i>American Journal of Bioethics</i> , 2020, 20, 101-103.	0.5	1
6	Marketing the Research Missions of Academic Medical Centers: Why Messages Blurring Lines Between Clinical Care and Research Are Bad for both Business and Ethics. <i>Cambridge Quarterly of Healthcare Ethics</i> , 2019, 28, 468-475.	0.5	3
7	Twentieth-century science education and 21st-century genetic engineering technologies: A toxic mix. <i>Accountability in Research</i> , 2019, 26, 271-275.	1.6	2
8	Four erroneous beliefs thwarting more trustworthy research. <i>ELife</i> , 2019, 8, .	2.8	10
9	The bench is closer to the bedside than we think: Uncovering the ethical ties between preclinical researchers in translational neuroscience and patients in clinical trials. <i>PLoS Biology</i> , 2018, 16, e2006343.	2.6	20
10	Why There Is No Obligation to Participate in Clinical Research. <i>Journal of Law, Medicine and Ethics</i> , 2017, 45, 327-332.	0.4	4
11	Ethical issues related to clinical research and rare diseases. <i>Translational Science of Rare Diseases</i> , 2017, 2, 175-194.	1.6	5
12	Building Trust Between Institutional Review Boards and Researchers. <i>Journal of General Internal Medicine</i> , 2016, 31, 987-989.	1.3	5
13	When There Are Only Two Who Can Tango: Ethical Concerns at the Juncture of Highly Novel Interventions and Precisely Targeted Research Populations. <i>American Journal of Bioethics</i> , 2015, 15, 85-86.	0.5	0
14	Inconsistent Approaches to Research Involving Cognitively Impaired Adults: Why the Broad View of Substituted Judgment Is Our Best Guide. <i>American Journal of Bioethics</i> , 2015, 15, 66-67.	0.5	4
15	Openness in science is key to keeping public trust. <i>Nature</i> , 2014, 515, 313-313.	13.7	19
16	Taking steps to increase the trustworthiness of scientific research. <i>FASEB Journal</i> , 2014, 28, 3841-3846.	0.2	13
17	Teaching Research Ethics Better: Focus on Excellent Science, Not Bad Scientists. <i>Clinical and Translational Science</i> , 2013, 6, 201-203.	1.5	16
18	Newborn screening and cascade testing for <i>FMR1</i> mutations. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 59-69.	0.7	24

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19	Relationships Hold the Key to Trustworthy and Productive Translational Science: Recommendations for Expanding Community Engagement in Biomedical Research. <i>Clinical and Translational Science</i> , 2013, 6, 310-313.	1.5	33
20	The Complex Ethics of First in Human Stem Cell Clinical Trials. <i>AJOB Neuroscience</i> , 2012, 3, 14-16.	0.6	6
21	Public trust and research a decade later: What have we learned since Jesse Gelsinger's death?. <i>Molecular Genetics and Metabolism</i> , 2009, 97, 4-5.	0.5	38
22	Transforming the Culture of Biomedical Research From Compliance to Trustworthiness: Insights From Nonmedical Sectors. <i>Academic Medicine</i> , 2009, 84, 472-477.	0.8	31
23	Responsible patient advocacy: Perspectives from the Alpha-1 Foundation. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 2845-2850.	0.7	6
24	Bioethics Consultation and Patient Advocacy Organizations: Expanding the Dialogue about Professional Conflicts of Interest. <i>Cambridge Quarterly of Healthcare Ethics</i> , 2007, 16, 74-81.	0.5	4
25	Currents in Contemporary Ethics: Informed Trust and the Financing of Biomedical Research. <i>Journal of Law, Medicine and Ethics</i> , 2006, 34, 460-464.	0.4	9
26	Additional Thoughts on Rethinking Research Ethics. <i>American Journal of Bioethics</i> , 2005, 5, 40-42.	0.5	11
27	Deciding for others at the end of life: storytelling and moral agency. <i>Journal of Clinical Ethics</i> , 2005, 16, 127-43.	0.1	6
28	Restoring and Preserving Trust in Biomedical Research. <i>Academic Medicine</i> , 2002, 77, 8-14.	0.8	31
29	Adults Are Not Big Children: Examining Surrogate Consent to Research Using Adults with Dementia. <i>Cambridge Quarterly of Healthcare Ethics</i> , 2002, 11, 160-168.	0.5	15
30	STRANGE BEDFELLOWS: HOW MEDICAL JURISPRUDENCE HAS INFLUENCED MEDICAL ETHICS AND MEDICAL PRACTICE. <i>Journal of Legal Medicine</i> , 2002, 23, 617-624.	0.4	0
31	Interprofessional Education in Ethics at an Academic Health Sciences Center. <i>Academic Medicine</i> , 2000, 75, 793-800.	0.8	48
32	The Private Health Insurance Industry: The Real Barrier to Healthcare Access?. <i>Cambridge Quarterly of Healthcare Ethics</i> , 1994, 3, 99-107.	0.5	1
33	Physician Advertising. <i>Southern Medical Journal</i> , 1989, 82, 1538-1544.	0.3	7
34	The role of beneficence in clinical genetics: Non-directive counseling reconsidered. <i>Theoretical Medicine and Bioethics</i> , 1989, 10, 139-149.	0.4	37
35	Continued Treatment of the Fatally Ill for the Benefit of Others. <i>Journal of the American Geriatrics Society</i> , 1988, 36, 63-67.	1.3	14