Thomas May

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3454488/publications.pdf

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51 papers	575 citations	687363 13 h-index	713466 21 g-index
52	52	52	536
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Ethical Issues in Clinical Genetics. , 2022, , 183-190.		O
2	A state-based approach to genomics for rare disease and population screening. Genetics in Medicine, 2021, 23, 777-781.	2.4	19
3	Ethical Considerations in the Use of Direct-to-Consumer Genetic Testing for Adopted Persons. Adoption Quarterly, 2021, 24, 89-100.	1.0	6
4	The Role of Race in Pandemic Vaccine Allocation. American Journal of Bioethics, 2021, 21, 89-91.	0.9	2
5	Hare's Archangel, Human Fallibility, and Utilitarian Justification(?) of Deception. American Journal of Bioethics, 2021, 21, 17-19.	0.9	2
6	Addressing perceived economic obstacles to genetic testing as a way to mitigate disparities in family health history for adoptees. Health Economics, Policy and Law, 2020, 15, 277-287.	1.8	4
7	Return of raw data in genomic testing and research: ownership, partnership, and risk–benefit. Genetics in Medicine, 2020, 22, 12-14.	2.4	2
8	Unintended Effects, latrogenic Harms, and the Challenge of Population-Wide Vaccination Compliance. American Journal of Bioethics, 2020, 20, 60-62.	0.9	1
9	Lockdown-type measures look effective against covid-19. BMJ, The, 2020, 370, m2809.	6.0	30
10	Are Public Repository Requirements Exacerbating Lack of Diversity?. Trends in Genetics, 2020, 36, 390-394.	6.7	4
11	Age, "Life-Cycles,―and the Allocation of Scarce Medical Resources. Chest, 2020, 158, 1837-1838.	0.8	4
12	You Kids Get off My Ethics Lawn!: An Admitted Curmudgeonly Critique of Credentialing Individual Clinical Ethics Consultants. American Journal of Bioethics, 2020, 20, 32-34.	0.9	8
13	Recruiting diversity where it exists: The Alabama Genomic Health Initiative. Journal of Genetic Counseling, 2020, 29, 471-478.	1.6	11
14	Anti-Vaxxers, Politicization of Science, and the Need for Trust in Pandemic Response. Journal of Health Communication, 2020, 25, 761-763.	2.4	18
15	Geographic variation and risk factors for teenage pregnancy in Uganda. African Health Sciences, 2020, 20, 1898-907.	0.7	4
16	N-of-1 Precision Medicine and Research Oversight. American Journal of Bioethics, 2019, 19, 36-37.	0.9	0
17	The value of genetic testing for family health history of adopted persons. Nature Reviews Genetics, 2019, 20, 65-66.	16.3	7
18	How the Atacama Skeleton Might Advance Discussion of Responsible Conduct of Research Responsibilities. Human Biology, 2019, 91, 5.	0.2	0

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19	Clarifying a Clinical Ethics Service's Value, the Visible and the Hidden. Journal of Clinical Ethics, 2019, 30, 251-261.	0.3	2
20	Healthcare Challenges Faced by Adopted Persons Lacking Family Health History Information. Narrative Inquiry in Bioethics, 2018, 8, 103-106.	0.1	12
21	Autonomy, Well-Being, and the Value of Genetic Testing for Adopted Persons. HEC Forum, 2018, 30, 283-295.	0.8	6
22	Sociogenetic Risks â€" Ancestry DNA Testing, Third-Party Identity, and Protection of Privacy. New England Journal of Medicine, 2018, 379, 410-412.	27.0	23
23	The Importance of Trust in the Vaccine Safety Enterprise. American Journal of Bioethics, 2017, 17, 48-50.	0.9	4
24	Adult adoptees' attitudes regarding the potential use of genetic information to fill the gap in their family health history. Adoption & Fostering, 2017, 41, 159-169.	0.5	16
25	Does Lack of "Genetic-Relative Family Health History―Represent a Potentially Avoidable Health Disparity for Adoptees?. American Journal of Bioethics, 2016, 16, 33-38.	0.9	27
26	In sickness and in health: context matters when considering potential benefits and risks of genome-wide sequencing. Genetics in Medicine, 2015, 17, 681-682.	2.4	1
27	Can targeted genetic testing offer useful health information to adoptees?. Genetics in Medicine, 2015, 17, 533-535.	2.4	21
28	An Adoptive Parental Perspective on Personal Genomic Screening. Pediatrics, 2015, 135, e811-e814.	2.1	6
29	On the Justifiability of ACMG Recommendations for Reporting of Incidental Findings in Clinical Exome and Genome Sequencing. Journal of Law, Medicine and Ethics, 2015, 43, 134-142.	0.9	19
30	The Limits of Traditional Approaches to Informed Consent for Genomic Medicine. HEC Forum, 2014, 26, 185-202.	0.8	8
31	Dual use opportunity and public health infrastructure. Journal of Medical Ethics, 2013, 39, 206-207.	1.8	0
32	On the Ethics of Clinical Whole Genome Sequencing of Children. Pediatrics, 2013, 132, 207-209.	2.1	14
33	Rethinking Clinical Risk for DNA Sequencing. American Journal of Bioethics, 2012, 12, 24-26.	0.9	9
34	Personal Morality and Professional Obligations: Rights of Conscience and Informed Consent. Perspectives in Biology and Medicine, 2009, 52, 30-38.	0.5	9
35	Children and Parents as Members of the Research Team: Fair Employment Practices Without a Union Contract. Ethics and Behavior, 2008, 18, 199-214.	1.8	1
36	Viewpoint: IRBs, Hospital Ethics Committees, and the Need for ???Translational Informed Consent???. Academic Medicine, 2007, 82, 670-674.	1.6	20

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37	Expanding Bioshield: A Call for Caution. American Journal of Public Health, 2007, 97, S23-S25.	2.7	2
38	Access to Hospitals in the Wake of Terrorism: Challenges and Needs for Maintaining Public Confidence. Disaster Management and Response, 2006, 4, 67-71.	1.4	8
39	PUBLIC COMMUNICATION, RISK PERCEPTION, AND THE VIABILITY OF PREVENTIVE VACCINATION AGAINST COMMUNICABLE DISEASES. Bioethics, 2005, 19, 407-421.	1.4	69
40	Free-Riding, Fairness, and the Rights of Minority Groups in Exemption from Mandatory Childhood Vaccination. Hum Vaccin, 2005, 1, 12-15.	2.4	28
41	Funding Agendas: Has Bioterror Defense Been Over-Prioritized?. American Journal of Bioethics, 2005, 5, 34-44.	0.9	13
42	Quality of life, justice, and the demands of hospital-based nursing. Public Affairs Quarterly, 2005, 19, 213-25.	0.2	0
43	Political Authority in a Bioterror Emergency. Journal of Law, Medicine and Ethics, 2004, 32, 159-163.	0.9	4
44	Isolation is not the answer. Nature, 2004, 429, 603-603.	27.8	10
45	Social Restrictions on Informed Consent: Research Ethics and Medical Decision Making. HEC Forum, 2004, 16, 38-44.	0.8	2
46	â€~Clustering of exemptions' as a collective action threat to herd immunity. Vaccine, 2003, 21, 1048-1051.	3.8	96
47	Vaccines as community-focused therapy. Expert Review of Vaccines, 2003, 2, 341-343.	4.4	12
48	Should Smallpox Vaccine be Made Available to the General Public?. Kennedy Institute of Ethics Journal, 2003, 13, 67-82.	0.5	5
49	The smallpox vaccination of health care workers: professional obligations and defense against bioterrorism. Hastings Center Report, 2003, 33, 26-33.	1.0	3
50	Do healthcare professionals have an obligation to be vaccinated against smallpox?., 2003, 2, 209-12.		0
51	Terror and triage: prioritizing access to mass smallpox vaccination. Creighton Law Review, 2003, 36, 359-74.	0.0	3