## Mark Sheehan

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5837517/publications.pdf

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76 papers

1,378 citations

331670 21 h-index 395702 33 g-index

79 all docs

79 docs citations

79 times ranked 1879 citing authors

#	Article	IF	CITATIONS
1	Public involvement in the governance of population-level biomedical research: unresolved questions and future directions. Journal of Medical Ethics, 2021, 47, 522-525.	1.8	20
2	Trust, trustworthiness and sharing patient data for research. Journal of Medical Ethics, 2021, 47, e26-e26.	1.8	24
3	Measuring the impact of participatory research in psychiatry: How the search for epistemic justifications obscures ethical considerations. Health Expectations, 2021, 24, 54-61.	2.6	24
4	The ethics of grandfather clauses in healthcare resource allocation. Bioethics, 2021, 35, 151-160.	1.4	2
5	â€There's more to life than money and health': Family caregivers' views on the role of Power of Attorney in proxy decisions about research participation for people living with dementia. Dementia, 2021, 20, 308-325.	2.0	8
6	Development of a decision support intervention for family members of adults who lack capacity to consent to trials. BMC Medical Informatics and Decision Making, 2021, 21, 30.	3.0	13
7	Ethical Issues in Consent for the Reuse of Data in Health Data Platforms. Science and Engineering Ethics, 2021, 27, 9.	2.9	13
8	Ethics review of big data research: What should stay and what should be reformed?. BMC Medical Ethics, 2021, 22, 51.	2.4	39
9	Constructing authentic decisions: proxy decision making for research involving adults who lack capacity to consent. Journal of Medical Ethics, 2021, 47, e42-e42.	1.8	11
10	Tragic choices in intensive care during the COVID-19 pandemic: on fairness, consistency and community. Journal of Medical Ethics, 2020, 46, 646-651.	1.8	18
11	†Your country needs you': the ethics of allocating staff to high-risk clinical roles in the management of patients with COVID-19. Journal of Medical Ethics, 2020, 46, 436-440.	1.8	43
12	Experimental Philosophical Bioethics. AJOB Empirical Bioethics, 2020, 11, 30-33.	1.6	29
13	Should the Decisions of Institutional Review Boards Be Consistent?. Ethics & Thics & Thics & Ethics &	0.9	28
14	Protection by exclusion? The (lack of) inclusion of adults who lack capacity to consent to research in clinical trials in the UK. Trials, 2019, 20, 474.	1.6	54
15	â€~lt's a tough decision': a qualitative study of proxy decision-making for research involving adults who lack capacity to consent in UK. Age and Ageing, 2019, 48, 903-909.	1.6	28
16	Authority and the Future of Consent in Population-Level Biomedical Research. Public Health Ethics, 2019, 12, 225-236.	1.0	17
17	Understanding and using patient experiences as evidence in healthcare priority setting. Cost Effectiveness and Resource Allocation, 2019, 17, 20.	1.5	37
18	Research involving adults lacking capacity to consent: a content analysis of participant information sheets for consultees and legal representatives in England and Wales. Trials, 2019, 20, 233.	1.6	19

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19	Patient and public involvement: Two sides of the same coin or different coins altogether?. Bioethics, 2019, 33, 708-715.	1.4	29
20	Structural Transformation to Attain Responsible BIOSciences (STARBIOS2): Protocol for a Horizon 2020 Funded European Multicenter Project to Promote Responsible Research and Innovation. JMIR Research Protocols, 2019, 8, e11745.	1.0	11
21	The Family Context of Assent: Comparison of Child and Parent Perspectives on Familial Decisionâ€Making. Children and Society, 2018, 32, 266-278.	1.7	O
22	In defence of governance: ethics review and social research. Journal of Medical Ethics, 2018, 44, 710-716.	1.8	15
23	Exploring the ethics of global health research priority-setting. BMC Medical Ethics, 2018, 19, 94.	2.4	61
24	Healthcare professionals' understanding of the legislation governing research involving adults lacking mental capacity in England and Wales: a national survey. Journal of Medical Ethics, 2018, 44, 632-637.	1.8	21
25	¿De quién es el conocimiento? El papel del conocimiento en un currÃeulum de gran autonomÃa. Arbor, 2018, 194, 442.	0.3	0
26	Reasonable disagreement and the justification of pre-emptive ethics governance in social research: a response to Hammersley. Journal of Medical Ethics, 2018, 44, 719-720.	1.8	1
27	Ethical understandings of proxy decision making for research involving adults lacking capacity: A systematic review (framework synthesis) of empirical research. AJOB Empirical Bioethics, 2018, 9, 267-286.	1.6	18
28	National Standards for Public Involvement in Research: missing the forest for the trees. Journal of Medical Ethics, 2018, 44, 801-804.	1.8	31
29	Standards of practice in empirical bioethics research: towards a consensus. BMC Medical Ethics, 2018, 19, 68.	2.4	62
30	The new Health Research Authority framework: still work to do to facilitate high-quality public health research in English local authorities. Perspectives in Public Health, 2017, 137, 262-265.	1.6	1
31	Variation in university research ethics review: Reflections following an inter-university study in England. Research Ethics, 2016, 12, 217-233.	1.7	24
32	The role of emotion in ethics and bioethics: dealing with repugnance and disgust. Journal of Medical Ethics, 2016, 42, 1-2.	1.8	13
33	Gene editing of human embryos and designing descendants. Maturitas, 2016, 94, 20-21.	2.4	1
34	Just Wars and doctors' strikes. Journal of Medical Ethics, 2016, 42, 693-694.	1.8	2
35	Research led by participants: a new social contract for a new kind of research. Journal of Medical Ethics, 2016, 42, 216-219.	1.8	67
36	Expertise, Ethics Expertise, and Clinical Ethics Consultation: Achieving Terminological Clarity. Journal of Medicine and Philosophy, 2016, 41, 416-433.	0.8	27

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37	Placebo-controlled study in neuromyelitis optica—Ethical and design considerations. Multiple Sclerosis Journal, 2016, 22, 862-872.	3.0	63
38	Developing a new justification for assent. BMC Medical Ethics, 2016, 17, 2.	2.4	30
39	Metaethics, Relativism and Empirical Bioethics. , 2016, , 33-50.		5
40	Refining the Enrolment Process in Emergency Medicine Research. The European Journal of Cardiovascular Medicine, 2016, 4, 506-510.	1.0	6
41	Caring About the Social Determinants of Health. American Journal of Bioethics, 2015, 15, 48-50.	0.9	1
42	The right to know and genetic testing. Journal of Medical Ethics, 2015, 41, 287-288.	1.8	7
43	Position statement on ethics, equipoise and research on charged particle radiation therapy. Journal of Medical Ethics, 2014, 40, 572-575.	1.8	20
44	Reining in patient and individual choice. Journal of Medical Ethics, 2014, 40, 291-292.	1.8	6
45	Ethical Review of Research on Human Subjects at Unilever: Reflections on Governance. Bioethics, 2014, 28, 284-292.	1.4	1
46	Placebo use in vaccine trials: Recommendations of a WHO expert panel. Vaccine, 2014, 32, 4708-4712.	3.8	45
47	Emergency medicine research: rites, rituals and consent. Emergency Medicine Journal, 2014, 31, 90-91.	1.0	2
48	"A degree of latitudeâ€Â• Thinking historically and making holistic judgements about internally assessed NCEA course work. Set Research Information for Teachers, 2014, , 18-23.	0.2	1
49	On the Nature and Sociology of Bioethics. Health Care Analysis, 2013, 21, 54-69.	2.2	15
50	Randomization Should Be Disclosed to Potential Research Subjects. American Journal of Bioethics, 2013, 13, 35-37.	0.9	1
51	Do we need research ethics committees?. Journal of Medical Ethics, 2013, 39, 485-485.	1.8	4
52	Religious red herrings. Journal of Medical Ethics, 2013, 39, 585-586.	1.8	0
53	Commissioning ethically and legally: the more things change, the more they stay the same. British Journal of General Practice, 2013, 63, 496-497.	1.4	3
54	New European Union regulation of clinical trials is not conflicting on deferred consent in emergency situations. BMJ, The, 2013, 346, f1163-f1163.	6.0	1

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55	Assent is not consent. Journal of Medical Ethics, 2012, 38, 3-3.	1.8	28
56	Toward Methodological Innovation in Empirical Ethics Research. Cambridge Quarterly of Healthcare Ethics, 2012, 21, 466-480.	0.8	76
57	Allocating Health Care Resources in the UK. , 2012, , 219-230.		5
58	Can Broad Consent be Informed Consent?. Public Health Ethics, 2011, 4, 226-235.	1.0	125
59	Delia Smith and the ethics committee. BMJ: British Medical Journal, 2011, 343, d6511-d6511.	2.3	0
60	Broad consent is informed consent. BMJ: British Medical Journal, 2011, 343, d6900-d6900.	2.3	34
61	Minimal Risk Remains an Open Question. American Journal of Bioethics, 2011, 11, 25-27.	0.9	2
62	It's unethical for general practitioners to be commissioners. BMJ, The, 2011, 342, d1430-d1430.	6.0	6
63	No Sex Please, We're Social Scientists?. American Journal of Bioethics, 2010, 10, 39-41.	0.9	4
64	Making Sense of the Immorality of Unnaturalness. Cambridge Quarterly of Healthcare Ethics, 2009, 18, 177-188.	0.8	10
65	The Duty to Disclose Adverse Clinical Trial Results. American Journal of Bioethics, 2009, 9, 24-32.	0.9	12
66	Disclosing Clinical Trial Results: Publicity, Significance and Independence. American Journal of Bioethics, 2009, 9, W3-W5.	0.9	0
67	Should research ethics committees meet in public?. Journal of Medical Ethics, 2008, 34, 631-635.	1.8	8
68	Is gene therapy for the treatment of male infertility ethical?. Nature Reviews Urology, 2008, 5, 596-597.	1.4	2
69	Ethics and policy: dealing with public attitudes. Radiation Protection Dosimetry, 2008, 129, 295-298.	0.8	4
70	Resources and the Rule of Rescue. Journal of Applied Philosophy, 2007, 24, 352-366.	1.0	22
71	Is the Community Consultation Requirement Necessary?. American Journal of Bioethics, 2006, 6, 38-40.	0.9	4
72	When should patients be held responsible for their lifestyle choices? BMJ: British Medical Journal, 2006, 332, 279.1.	2.3	2

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73	ethical review and ethical behaviour. BMJ: British Medical Journal, 2005, 330, 473.1.	2.3	3
74	Orphan drugs and the NHS. BMJ: British Medical Journal, 2005, 331, 1144.4-1145.	2.3	8
75	Ethical aspects of saviour siblings: procreative reasons and the treatment of children., 0,, 59-70.		0
76	Moral Relativism. , 0, , 93-98.		1