

# Lucy Frith

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6466023/publications.pdf>

Version: 2024-02-01

79  
papers

1,845  
citations

279798

23  
h-index

302126

39  
g-index

82  
all docs

82  
docs citations

82  
times ranked

1806  
citing authors

#	ARTICLE	IF	CITATIONS
1	Community engagement in genomic research: Proposing a strategic model for effective participation of indigenous communities. <i>Developing World Bioethics</i> , 2022, 22, 189-202.	0.9	4
2	Good practice recommendations for information provision for those involved in reproductive donation. <i>Human Reproduction Open</i> , 2022, 2022, hoac001.	5.4	16
3	The concise argument "choice, choices and the choice agenda. <i>Journal of Medical Ethics</i> , 2022, 48, 1-2.	1.8	2
4	Editorial "The ethical implications of treating a pregnant woman to benefit the fetus. <i>Journal of Cystic Fibrosis</i> , 2022, 21, 558-559.	0.7	1
5	Tensions Between Ethics and the Law: Examination of a Legal Case by Two Midwives Invoking a Conscientious Objection to Abortion in Scotland. <i>HEC Forum</i> , 2021, 33, 189-213.	0.8	3
6	Public involvement in the governance of population-level biomedical research: unresolved questions and future directions. <i>Journal of Medical Ethics</i> , 2021, 47, 522-525.	1.8	20
7	The ethics of preconception expanded carrier screening in patients seeking assisted reproduction. <i>Human Reproduction Open</i> , 2021, 2021, hoaa063.	5.4	14
8	Which ethical values underpin England's National Health Service reset of paediatric and maternity services following COVID-19: a rapid review. <i>BMJ Open</i> , 2021, 11, e049214.	1.9	6
9	Neither "Crisis Light" nor "Business as Usual": Considering the Distinctive Ethical Issues Raised by the Contingency and Reset Phases of a Pandemic. <i>American Journal of Bioethics</i> , 2021, 21, 34-37.	0.9	3
10	Public involvement and engagement in big data research: protocol for a scoping review and a systematic review of delivery and effectiveness of strategies for involvement and engagement. <i>BMJ Open</i> , 2021, 11, e050167.	1.9	2
11	Strategies to enhance recruitment and consent to intensive care studies: a qualitative study with researchers and patient "public involvement contributors. <i>BMJ Open</i> , 2021, 11, e048193.	1.9	7
12	COVID-19 and beyond: the ethical challenges of resetting health services during and after public health emergencies. <i>Journal of Medical Ethics</i> , 2020, 46, 715-716.	1.8	17
13	ESHRE guideline: female fertility preservation. <i>Human Reproduction Open</i> , 2020, 2020, hoaa052.	5.4	282
14	Understanding childbirth as a complex salutogenic phenomenon: The EU COST BIRTH Action Special Collection. <i>PLoS ONE</i> , 2020, 15, e0236722.	2.5	4
15	Proposing a sociology of donation: The donation of body parts and products for art, education, research, or treatment. <i>Sociology Compass</i> , 2020, 14, 1-16.	2.5	25
16	Pandemic medical ethics. <i>Journal of Medical Ethics</i> , 2020, 46, 353-354.	1.8	8
17	Constructing a Bioethical Framework to Evaluate and Optimise Newborn Bloodspot Screening for Cystic Fibrosis. <i>International Journal of Neonatal Screening</i> , 2020, 6, 40.	3.2	2
18	The concise argument. <i>Journal of Medical Ethics</i> , 2020, 46, 1-2.	1.8	3

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19	Non-Electoral Representation and Promoting Welfare” Beyond Descriptive Representation. <i>American Journal of Bioethics</i> , 2020, 20, 56-58.	0.9	0
20	Contacting gamete donors to facilitate diagnostic genetic testing for the donor-conceived child: what are the rights and obligations of gamete donors in these cases? A response to Horton et al. <i>Journal of Medical Ethics</i> , 2020, 46, 220-222.	1.8	5
21	Variations in use of childbirth interventions in 13 high-income countries: A multinational cross-sectional study. <i>PLoS Medicine</i> , 2020, 17, e1003103.	8.4	92
22	Ethics education and moral decision-making in clinical commissioning: an interview study. <i>British Journal of General Practice</i> , 2020, 70, e45-e54.	1.4	3
23	Lockdown, public good and equality during COVID-19. <i>Journal of Medical Ethics</i> , 2020, 46, 713-714.	1.8	5
24	The point of no return: Up to what point should we be allowed to withdraw consent to the storage and use of embryos and gametes?. <i>Bioethics</i> , 2019, 33, 637-643.	1.4	2
25	Editorial: Towards a sociology of donation. <i>Sociology of Health and Illness</i> , 2019, 41, 549-552.	2.1	4
26	Role of public involvement in the Royal College of Physicians’s Future Hospitals healthcare improvement programme: an evaluation. <i>BMJ Open</i> , 2019, 9, e027680.	1.9	6
27	Online interviewing with interpreters in humanitarian contexts. <i>International Journal of Qualitative Studies on Health and Well-being</i> , 2018, 13, 1444887.	1.6	20
28	Institute of Medical Ethics Guidelines for confirmation of appointment, promotion and recognition of UK bioethics and medical ethics researchers. <i>Journal of Medical Ethics</i> , 2018, 44, 289-291.	1.8	0
29	Variations in childbirth interventions in high-income countries: protocol for a multinational cross-sectional study. <i>BMJ Open</i> , 2018, 8, e017993.	1.9	10
30	Conscientious objection to participation in abortion by midwives and nurses: a systematic review of reasons. <i>BMC Medical Ethics</i> , 2018, 19, 31.	2.4	40
31	Secrets and disclosure in donor conception. <i>Sociology of Health and Illness</i> , 2018, 40, 188-203.	2.1	34
32	Searching for “relations” using a DNA linking register by adults conceived following sperm donation. <i>BioSocieties</i> , 2018, 13, 170-189.	1.3	20
33	Relative solidarity: Conceptualising communal participation in genomic research among potential research participants in a developing Sub-Saharan African setting. <i>PLoS ONE</i> , 2018, 13, e0195171.	2.5	13
34	Standards of practice in empirical bioethics research: towards a consensus. <i>BMC Medical Ethics</i> , 2018, 19, 68.	2.4	62
35	Children’s views on research without prior consent in emergency situations: a UK qualitative study. <i>BMJ Open</i> , 2018, 8, e022894.	1.9	15
36	Gamete donors’ reasons for, and expectations and experiences of, registration with a voluntary donor linking register. <i>Human Fertility</i> , 2017, 20, 268-278.	1.7	17

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37	Ethical standards for mental health and psychosocial support research in emergencies: review of literature and current debates. <i>Globalization and Health</i> , 2017, 13, 8.	4.9	27
38	Family building using embryo adoption: relationships and contact arrangements between provider and recipient families—a mixed-methods study. <i>Human Reproduction</i> , 2017, 32, 1092-1099.	0.9	12
39	Measuring antenatal care use in Europe: is the content and timing of care in pregnancy tool applicable?. <i>International Journal of Public Health</i> , 2017, 62, 583-590.	2.3	5
40	Debt Counselling for Depression in Primary Care: an adaptive randomised controlled pilot trial (DeCoDer study). <i>Health Technology Assessment</i> , 2017, 21, 1-164.	2.8	15
41	The changing face of the English National Health Service: new providers, markets and morality. <i>British Medical Bulletin</i> , 2016, 119, 5-16.	6.9	5
42	Voluntary DNA-based information exchange and contact services following donor conception: an analysis of service users'™ needs. <i>New Genetics and Society</i> , 2016, 35, 372-392.	1.2	15
43	Managing Ethical Challenges to Mental Health Research in Post-Conflict Settings. <i>Developing World Bioethics</i> , 2016, 16, 15-28.	0.9	18
44	Fifteen-minute consultation: an evidence-based approach to research without prior consent (deferred) and Practice Edition, 2016, 101, 49-53.	0.5	38
45	Assessing the performance of maternity care in Europe: a critical exploration of tools and indicators. <i>BMC Health Services Research</i> , 2015, 15, 491.	2.2	33
46	Evolving minimum standards in responsible international sperm donor offspring quota. <i>Reproductive BioMedicine Online</i> , 2015, 30, 568-580.	2.4	17
47	The limits of evidence: evidence based policy and the removal of gamete donor anonymity in the UK. <i>Monash Bioethics Review</i> , 2015, 33, 29-44.	0.8	10
48	Expectations and experiences of gamete donors and donor-conceived adults searching for genetic relatives using DNA linking through a voluntary register. <i>Human Reproduction</i> , 2015, 30, 111-121.	0.9	38
49	How parents and practitioners experience research without prior consent (deferred consent) for emergency research involving children with life threatening conditions: a mixed method study. <i>BMJ Open</i> , 2015, 5, e008522.	1.9	84
50	Access to genetic and biographical history in donor conception: An analysis of recent trends and future possibilities. , 2015, , 136-152.		9
51	The changing National Health Service: market-based reform and morality Comment on "Morality and Markets in the NHS". <i>International Journal of Health Policy and Management</i> , 2015, 4, 253-255.	0.9	3
52	Social enterprises, health-care provision and ethical capital. <i>Social Enterprise Journal</i> , 2014, 10, 105-120.	1.8	8
53	Patient and Public Participation in Health Care: Can We Do It Better?. <i>American Journal of Bioethics</i> , 2014, 14, 17-18.	0.9	4
54	Assisted reproductive technology in the USA: is more regulation needed?. <i>Reproductive BioMedicine Online</i> , 2014, 29, 516-523.	2.4	30

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55	They Can't Have My Embryo: The Ethics of Conditional Embryo Donation. <i>Bioethics</i> , 2013, 27, 317-324.	1.4	10
56	A survey of 1700 women who formed their families using donor spermatozoa. <i>Reproductive BioMedicine Online</i> , 2013, 27, 436-447.	2.4	25
57	The NHS and market forces in healthcare: the need for organisational ethics. <i>Journal of Medical Ethics</i> , 2013, 39, 17-21.	1.8	21
58	Donor Conception and Mandatory Paternity Testing: The Right to Know and the Right to Be Told. <i>American Journal of Bioethics</i> , 2013, 13, 50-52.	0.9	12
59	How experience makes a difference: practitioners' views on the use of deferred consent in paediatric and neonatal emergency care trials. <i>BMC Medical Ethics</i> , 2013, 14, 45.	2.4	45
60	Forming a family with sperm donation: a survey of 244 non-biological parents. <i>Reproductive BioMedicine Online</i> , 2012, 24, 709-718.	2.4	25
61	SYMBIOTIC EMPIRICAL ETHICS: A PRACTICAL METHODOLOGY. <i>Bioethics</i> , 2012, 26, 198-206.	1.4	73
62	Donor-conceived people's views and experiences of their genetic origins: a critical analysis of the research evidence. <i>Journal of Law &amp; Medicine</i> , 2012, 19, 769-89.	0.0	64
63	Ethical boundary work in the infertility clinic. <i>Sociology of Health and Illness</i> , 2011, 33, 570-585.	2.1	23
64	Conditional embryo relinquishment: choosing to relinquish embryos for family-building through a Christian embryo 'adoption' programme. <i>Human Reproduction</i> , 2011, 26, 3327-3338.	0.9	29
65	The use of clinical ethics committees in infertility clinics. <i>Human Fertility</i> , 2009, 12, 89-94.	1.7	6
66	Use or ornament? Clinical ethics committees in infertility units: a qualitative study. <i>Clinical Ethics</i> , 2009, 4, 91-97.	0.7	5
67	The Role of Birth Certificates in Relation to Access to Biographical and Genetic History in Donor Conception. <i>International Journal of Children's Rights</i> , 2009, 17, 207-233.	0.6	26
68	Process and consensus: ethical decision-making in the infertility clinic—a qualitative study. <i>Journal of Medical Ethics</i> , 2009, 35, 662-667.	1.8	7
69	The UK's gamete donor 'crisis' - a critical analysis. <i>Critical Social Policy</i> , 2008, 28, 74-95.	1.9	25
70	UK gamete donors' reflections on the removal of anonymity: implications for recruitment. <i>Human Reproduction</i> , 2007, 22, 1675-1680.	0.9	61
71	HIV self-testing: a time to revise current policy. <i>Lancet</i> , The, 2007, 369, 243-245.	13.7	41
72	Gamete Donation, Identity, and the Offspring's Right to Know. <i>AMA Journal of Ethics</i> , 2007, 9, 644-648.	0.7	3

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73	Researching chronic childhood illness: autonomy or beneficence?. <i>Chronic Illness</i> , 2006, 2, 178-180.	1.5	0
74	HIV testing and informed consent. <i>Journal of Medical Ethics</i> , 2005, 31, 699-700.	1.8	6
75	Gamete donation and anonymity. <i>Human Reproduction</i> , 2001, 16, 818-824.	0.9	128
76	Beneath the Rhetoric: The Role of Rights in the Practice of Non-Anonymous Gamete Donation. <i>Bioethics</i> , 2001, 15, 473-484.	1.4	50
77	Priority setting and evidence based purchasing. <i>Health Care Analysis</i> , 1999, 7, 139-151.	2.2	8
78	Relatedness in clinical practice. , 0, , 129-144.		1
79	Publishing Research in Empirical Bioethics: Quality, Disciplines and Expertise. , 0, , 235-255.		7