Lucy Frith

List of Publications by Year in descending order

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

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79	1,845	23	39
papers	citations	h-index	g-index
82	82	82	1806
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	ESHRE guideline: female fertility preservationâ€. Human Reproduction Open, 2020, 2020, hoaa052.	5.4	282
2	Gamete donation and anonymity. Human Reproduction, 2001, 16, 818-824.	0.9	128
3	Variations in use of childbirth interventions in 13 high-income countries: A multinational cross-sectional study. PLoS Medicine, 2020, 17, e1003103.	8.4	92
4	How parents and practitioners experience research without prior consent (deferred consent) for emergency research involving children with life threatening conditions: a mixed method study. BMJ Open, 2015, 5, e008522.	1.9	84
5	SYMBIOTIC EMPIRICAL ETHICS: A PRACTICAL METHODOLOGY. Bioethics, 2012, 26, 198-206.	1.4	73
6	Donor-conceived people's views and experiences of their genetic origins: a critical analysis of the research evidence. Journal of Law & Medicine, 2012, 19, 769-89.	0.0	64
7	Standards of practice in empirical bioethics research: towards a consensus. BMC Medical Ethics, 2018, 19, 68.	2.4	62
8	UK gamete donors' reflections on the removal of anonymity: implications for recruitment. Human Reproduction, 2007, 22, 1675-1680.	0.9	61
9	Beneath the Rhetoric: The Role of Rights in the Practice of Non-Anonymous Gamete Donation. Bioethics, 2001, 15, 473-484.	1.4	50
10	How experience makes a difference: practitioners' views on the use of deferred consent in paediatric and neonatal emergency care trials. BMC Medical Ethics, 2013, 14, 45.	2.4	45
11	HIV self-testing: a time to revise current policy. Lancet, The, 2007, 369, 243-245.	13.7	41
12	Conscientious objection to participation in abortion by midwives and nurses: a systematic review of reasons. BMC Medical Ethics, 2018, 19, 31.	2.4	40
13	Expectations and experiences of gamete donors and donor-conceived adults searching for genetic relatives using DNA linking through a voluntary register. Human Reproduction, 2015, 30, 111-121.	0.9	38
14	Fifteen-minute consultation: an evidence-based approach to research without prior consent (deferred) Tj ETQq0 and Practice Edition, 2016, 101, 49-53.	0 0 rgBT / 0.5	Overlock 10 T 38
15	Secrets and disclosure in donor conception. Sociology of Health and Illness, 2018, 40, 188-203.	2.1	34
16	Assessing the performance of maternity care in Europe: a critical exploration of tools and indicators. BMC Health Services Research, 2015, 15, 491.	2.2	33
17	Assisted reproductive technology in the USA: is more regulation needed?. Reproductive BioMedicine Online, 2014, 29, 516-523.	2.4	30
18	Conditional embryo relinquishment: choosing to relinquish embryos for family-building through a Christian embryo 'adoption' programme. Human Reproduction, 2011, 26, 3327-3338.	0.9	29

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19	Ethical standards for mental health and psychosocial support research in emergencies: review of literature and current debates. Globalization and Health, 2017, 13, 8.	4.9	27
20	The Role of Birth Certificates in Relation to Access to Biographical and Genetic History in Donor Conception. International Journal of Children's Rights, 2009, 17, 207-233.	0.6	26
21	The UK's gamete donor`crisis' - a critical analysis. Critical Social Policy, 2008, 28, 74-95.	1.9	25
22	Forming a family with sperm donation: a survey of 244 non-biological parents. Reproductive BioMedicine Online, 2012, 24, 709-718.	2.4	25
23	A survey of 1700 women who formed their families using donor spermatozoa. Reproductive BioMedicine Online, 2013, 27, 436-447.	2.4	25
24	Proposing a sociology of donation: The donation of body parts and products for art, education, research, or treatment. Sociology Compass, 2020, 14, 1-16.	2.5	25
25	Ethical boundaryâ€work in the infertility clinic. Sociology of Health and Illness, 2011, 33, 570-585.	2.1	23
26	The NHS and market forces in healthcare: the need for organisational ethics. Journal of Medical Ethics, 2013, 39, 17-21.	1.8	21
27	Online interviewing with interpreters in humanitarian contexts. International Journal of Qualitative Studies on Health and Well-being, 2018, 13, 1444887.	1.6	20
28	Searching for â€relations' using a DNA linking register by adults conceived following sperm donation. BioSocieties, 2018, 13, 170-189.	1.3	20
29	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
30	Managing Ethical Challenges to Mental Health Research in Postâ€Conflict Settings. Developing World Bioethics, 2016, 16, 15-28.	0.9	18
31	Evolving minimum standards in responsible international sperm donor offspring quota. Reproductive BioMedicine Online, 2015, 30, 568-580.	2.4	17
32	Gamete donors' reasons for, and expectations and experiences of, registration with a voluntary donor linking register. Human Fertility, 2017, 20, 268-278.	1.7	17
33	COVID-19 and beyond: the ethical challenges of resetting health services during and after public health emergencies. Journal of Medical Ethics, 2020, 46, 715-716.	1.8	17
34	Good practice recommendations for information provision for those involved in reproductive donationâ€. Human Reproduction Open, 2022, 2022, hoac001.	5.4	16
35	Voluntary DNA-based information exchange and contact services following donor conception: an analysis of service users' needs. New Genetics and Society, 2016, 35, 372-392.	1.2	15
36	Children's views on research without prior consent in emergency situations: a UK qualitative study. BMJ Open, 2018, 8, e022894.	1.9	15

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37	Debt Counselling for Depression in Primary Care: an adaptive randomised controlled pilot trial (DeCoDer study). Health Technology Assessment, 2017, 21, 1-164.	2.8	15
38	The ethics of preconception expanded carrier screening in patients seeking assisted reproduction. Human Reproduction Open, 2021, 2021, hoaa063.	5.4	14
39	Relative solidarity: Conceptualising communal participation in genomic research among potential research participants in a developing Sub-Saharan African setting. PLoS ONE, 2018, 13, e0195171.	2.5	13
40	Donor Conception and Mandatory Paternity Testing: The Right to Know and the Right to Be Told. American Journal of Bioethics, 2013, 13, 50-52.	0.9	12
41	Family building using embryo adoption: relationships and contact arrangements between provider and recipient families—a mixed-methods study. Human Reproduction, 2017, 32, 1092-1099.	0.9	12
42	They Can't HaveMyEmbryo: The Ethics of Conditional Embryo Donation. Bioethics, 2013, 27, 317-324.	1.4	10
43	The limits of evidence: evidence based policy and the removal of gamete donor anonymity in the UK. Monash Bioethics Review, 2015, 33, 29-44.	0.8	10
44	Variations in childbirth interventions in high-income countries: protocol for a multinational cross-sectional study. BMJ Open, 2018, 8, e017993.	1.9	10
45	Access to genetic and biographical history in donor conception: An analysis of recent trends and future possibilities., 2015,, 136-152.		9
46	Priority setting and evidence based purchasing. Health Care Analysis, 1999, 7, 139-151.	2.2	8
47	Social enterprises, health-care provision and ethical capital. Social Enterprise Journal, 2014, 10, 105-120.	1.8	8
48	Pandemic medical ethics. Journal of Medical Ethics, 2020, 46, 353-354.	1.8	8
49	Process and consensus: ethical decision-making in the infertility clinic-a qualitative study. Journal of Medical Ethics, 2009, 35, 662-667.	1.8	7
50	Publishing Research in Empirical Bioethics: Quality, Disciplines and Expertise., 0,, 235-255.		7
51	Strategies to enhance recruitment and consent to intensive care studies: a qualitative study with researchers and patient–public involvement contributors. BMJ Open, 2021, 11, e048193.	1.9	7
52	HIV testing and informed consent. Journal of Medical Ethics, 2005, 31, 699-700.	1.8	6
53	The use of clinical ethics committees in infertility clinics. Human Fertility, 2009, 12, 89-94.	1.7	6
54	Role of public involvement in the Royal College of Physicians' Future Hospitals healthcare improvement programme: an evaluation. BMJ Open, 2019, 9, e027680.	1.9	6

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55	Which ethical values underpin England's National Health Service reset of paediatric and maternity services following COVID-19: a rapid review. BMJ Open, 2021, 11, e049214.	1.9	6
56	Use or ornament? Clinical ethics committees in infertility units: a qualitative study. Clinical Ethics, 2009, 4, 91-97.	0.7	5
57	The changing face of the English National Health Service: new providers, markets and morality. British Medical Bulletin, 2016, 119, 5-16.	6.9	5
58	Measuring antenatal care use in Europe: is the content and timing of care in pregnancy tool applicable?. International Journal of Public Health, 2017, 62, 583-590.	2.3	5
59	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. Journal of Medical Ethics, 2020, 46, 220-222.	1.8	5
60	Lockdown, public good and equality during COVID-19. Journal of Medical Ethics, 2020, 46, 713-714.	1.8	5
61	Patient and Public Participation in Health Care: Can We Do It Better?. American Journal of Bioethics, 2014, 14, 17-18.	0.9	4
62	Editorial: Towards a sociology of donation. Sociology of Health and Illness, 2019, 41, 549-552.	2.1	4
63	Understanding childbirth as a complex salutogenic phenomenon: The EU COST BIRTH Action Special Collection. PLoS ONE, 2020, 15, e0236722.	2.5	4
64	Community engagement in genomic research: Proposing a strategic model for effective participation of indigenous communities. Developing World Bioethics, 2022, 22, 189-202.	0.9	4
65	Gamete Donation, Identity, and the Offspring's Right to Know. AMA Journal of Ethics, 2007, 9, 644-648.	0.7	3
66	Tensions Between Ethics and the Law: Examination of a Legal Case by Two Midwives Invoking a Conscientious Objection to Abortion in Scotland. HEC Forum, 2021, 33, 189-213.	0.8	3
67	The concise argument. Journal of Medical Ethics, 2020, 46, 1-2.	1.8	3
68	Neither â€~Crisis Light' nor â€~Business as Usual': Considering the Distinctive Ethical Issues Raised by the Contingency and Reset Phases of a Pandemic. American Journal of Bioethics, 2021, 21, 34-37.	0.9	3
69	Ethics education and moral decision-making in clinical commissioning: an interview study. British Journal of General Practice, 2020, 70, e45-e54.	1.4	3
70	The changing National Health Service: market-based reform and morality Comment on "Morality and Markets in the NHS". International Journal of Health Policy and Management, 2015, 4, 253-255.	0.9	3
71	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?. Bioethics, 2019, 33, 637-643.	1.4	2
72	Constructing a Bioethical Framework to Evaluate and Optimise Newborn Bloodspot Screening for Cystic Fibrosis. International Journal of Neonatal Screening, 2020, 6, 40.	3.2	2

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#	Article	lF	CITATIONS
73	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. BMJ Open, 2021, 11, e050167.	1.9	2
74	The concise argument – choice, choices and the choice agenda. Journal of Medical Ethics, 2022, 48, 1-2.	1.8	2
75	Relatedness in clinical practice. , 0, , 129-144.		1
76	Editorial $\hat{a}\in$ The ethical implications of treating a pregnant woman to benefit the fetus. Journal of Cystic Fibrosis, 2022, 21, 558-559.	0.7	1
77	Researching chronic childhood illness: autonomy or beneficence?. Chronic Illness, 2006, 2, 178-180.	1.5	O
78	Institute of Medical Ethics Guidelines for confirmation of appointment, promotion and recognition of UK bioethics and medical ethics researchers. Journal of Medical Ethics, 2018, 44, 289-291.	1.8	0
79	Non-Electoral Representation and Promoting Welfare—Beyond Descriptive Representation. American Journal of Bioethics, 2020, 20, 56-58.	0.9	0