

# Benjamin Berkman

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6749099/publications.pdf>

Version: 2024-02-01

47  
papers

1,606  
citations

516681

16  
h-index

315719

38  
g-index

47  
all docs

47  
docs citations

47  
times ranked

2394  
citing authors

#	ARTICLE	IF	CITATIONS
1	The ethics of encouraging employees to get the COVID-19 vaccination. <i>Journal of Public Health Policy</i> , 2022, 43, 311-319.	2.0	3
2	Autonomy concerns with using contracts to enhance patient adherence. <i>Bioethics</i> , 2022, , .	1.4	0
3	Navigating Pandemic Moral Distress at Home and at Work: Frontline Workers's Experiences. <i>AJOB Empirical Bioethics</i> , 2022, 13, 215-225.	1.6	2
4	A new ethical framework to determine acceptable risks in fetal therapy trials. <i>Prenatal Diagnosis</i> , 2022, 42, 962-969.	2.3	1
5	Allocation of scarce biospecimens for use in research. <i>Journal of Medical Ethics</i> , 2021, 47, 740-743.	1.8	2
6	Women's Just Another Added Benefit? Women's Experiences with Employment-Based Egg Freezing Programs. <i>AJOB Empirical Bioethics</i> , 2021, 12, 41-52.	1.8	5
7	Should germline genome editing be allowed? The effect of treatment characteristics on public acceptability. <i>Human Reproduction</i> , 2021, 36, 465-478.	0.9	5
8	The Deep Structure of Deliberate Ignorance: Mapping the Terrain. , 2021, , 65-88.		1
9	Exploring the motivations of research participants who chose not to learn medically actionable secondary genetic findings about themselves. <i>Genetics in Medicine</i> , 2021, 23, 2281-2288.	2.4	17
10	Implementing Expanded Prenatal Genetic Testing: Should Parents Have Access to Any and All Fetal Genetic Information?. <i>American Journal of Bioethics</i> , 2021, , 1-26.	0.9	26
11	Response to Clayton et al.. <i>Genetics in Medicine</i> , 2021, , .	2.4	0
12	An ethical framework for genetic counseling in the genomic era. <i>Journal of Genetic Counseling</i> , 2020, 29, 718-727.	1.6	35
13	Covid-19 Vaccine Trials and Incarcerated People – The Ethics of Inclusion. <i>New England Journal of Medicine</i> , 2020, 383, 1897-1899.	27.0	14
14	Commentary on “The right not to know and the obligation not to know”. <i>Journal of Medical Ethics</i> , 2020, 46, 304-305.	1.8	3
15	Re-examining the Ethics of Genetic Counselling in the Genomic Era. <i>Journal of Bioethical Inquiry</i> , 2020, 17, 325-335.	1.5	11
16	Should Patient Groups Have the Power to Redirect How Their Samples Are Used?. <i>American Journal of Bioethics</i> , 2019, 19, 26-28.	0.9	0
17	Noninvasive Prenatal Whole Genome Sequencing. <i>Obstetrics and Gynecology</i> , 2019, 133, 525-532.	2.4	24
18	Reconceptualizing harms and benefits in the genomic age. <i>Personalized Medicine</i> , 2018, 15, 419-428.	1.5	13

#	ARTICLE	IF	CITATIONS
19	Reconsidering the Need for Reconsent at 18. <i>Pediatrics</i> , 2018, 142, .	2.1	12
20	Development of a consensus approach for return of pathology incidental findings in the Genotype-Tissue Expression (GTEx) project. <i>Journal of Medical Ethics</i> , 2018, 44, 643-645.	1.8	3
21	Recontacting participants for expanded uses of existing samples and data: a case study. <i>Genetics in Medicine</i> , 2017, 19, 883-889.	2.4	3
22	Prenatal Whole Genome Sequencing: An Argument for Professional Self-Regulation. <i>American Journal of Bioethics</i> , 2017, 17, 26-28.	0.9	10
23	A Proposed Process for Reliably Updating the Common Rule. <i>American Journal of Bioethics</i> , 2017, 17, 8-14.	0.9	6
24	Views of American OB/GYNs on the ethics of prenatal whole-genome sequencing. <i>Prenatal Diagnosis</i> , 2016, 36, 1250-1256.	2.3	24
25	When Participants in Genomic Research Grow Up: Contact and Consent at the Age of Majority. <i>Journal of Pediatrics</i> , 2016, 168, 226-231.e1.	1.8	17
26	The Ethics of Large-Scale Genomic Research. , 2016, , 53-69.		3
27	Towards a more representative morphology: clinical and ethical considerations for including diverse populations in diagnostic genetic atlases. <i>Genetics in Medicine</i> , 2016, 18, 1069-1074.	2.4	27
28	Compensation for egg donation: a zero-sum game. <i>Fertility and Sterility</i> , 2016, 105, 1153-1154.	1.0	4
29	Institutional review board perspectives on obligations to disclose genetic incidental findings to research participants. <i>Genetics in Medicine</i> , 2016, 18, 705-711.	2.4	26
30	When Should Genome Researchers Disclose Misattributed Paternity?. <i>Hastings Center Report</i> , 2015, 45, 28-36.	1.0	14
31	Parental Consent for the Use of Residual Newborn Screening Bloodspots. <i>JAMA - Journal of the American Medical Association</i> , 2015, 314, 21.	7.4	9
32	Scrutinizing the Right Not to Know. <i>American Journal of Bioethics</i> , 2015, 15, 17-19.	0.9	5
33	Points to Consider: Ethical, Legal, and Psychosocial Implications of Genetic Testing in Children and Adolescents. <i>American Journal of Human Genetics</i> , 2015, 97, 6-21.	6.2	453
34	Ethics of genetic and biomarker test disclosures in neurodegenerative disease prevention trials. <i>Neurology</i> , 2015, 84, 1488-1494.	1.1	49
35	FDA's Proposed Guidance for Laboratory Developed Tests: How Should Regulators Balance the Risks and Promise of Innovation in Clinical Genetics?. <i>FDA's Food and Drug Policy Forum</i> , 2015, 5, .	0.0	1
36	The unintended implications of blurring the line between research and clinical care in a genomic age. <i>Personalized Medicine</i> , 2014, 11, 285-295.	1.5	38

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37	A Framework for Analyzing the Ethics of Disclosing Genetic Research Findings. <i>Journal of Law, Medicine and Ethics</i> , 2014, 42, 190-207.	0.9	38
38	Return of Genomic Results to Research Participants: The Floor, the Ceiling, and the Choices In Between. <i>American Journal of Human Genetics</i> , 2014, 94, 818-826.	6.2	342
39	Grapppling With Genomic Incidental Findings in the Clinical Realm. <i>Chest</i> , 2014, 145, 226-230.	0.8	4
40	Finding Fault? Exploring Legal Duties to Return Incidental Findings in Genomic Research. <i>Georgetown Law Journal</i> , 2014, 102, 795-843.	1.0	12
41	Do Researchers Have an Obligation to Actively Look for Genetic Incidental Findings?. <i>American Journal of Bioethics</i> , 2013, 13, 32-42.	0.9	81
42	Disclosure of Incidental Findings From Next-Generation Sequencing in Pediatric Genomic Research. <i>Pediatrics</i> , 2013, 131, 564-571.	2.1	76
43	Response to Open Peer Commentaries on "Do Researchers Have an Obligation to Actively Look for Genetic Incidental Findings?" <i>American Journal of Bioethics</i> , 2013, 13, W10-W11.	0.9	3
44	Prenatal Whole Genome Sequencing. <i>Hastings Center Report</i> , 2012, 42, 28-40.	1.0	57
45	Genomics really gets personal: How exome and whole genome sequencing challenge the ethical framework of human genetics research. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 2916-2924.	1.2	119
46	Incorporating explicit ethical reasoning into pandemic influenza policies. <i>The Journal of Contemporary Health Law and Policy</i> , 2009, 26, 1-19.	0.0	6
47	Should institutions fund the feedback of individual findings in genomic research?. <i>Journal of Medical Ethics</i> , 0, , medethics-2021-107992.	1.8	2