

Robert L Klitzman

List of Publications by Year in descending order

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Version: 2024-02-01

128
papers

3,334
citations

147801

31
h-index

182427

51
g-index

130
all docs

130
docs citations

130
times ranked

3345
citing authors

#	ARTICLE	IF	CITATIONS
1	Understanding Ethical Challenges in Medical Education Research. <i>Academic Medicine</i> , 2022, 97, 18-21.	1.6	6
2	Henrietta Lacks's family's lawsuits: ethical questions and solutions. <i>Trends in Biotechnology</i> , 2022, 40, 769-772.	9.3	1
3	In Reply to Jain and Kassam:. <i>Academic Medicine</i> , 2022, 97, 8-8.	1.6	0
4	How hospital chaplains develop and use rituals to address medical staff distress. <i>SSM Qualitative Research in Health</i> , 2022, 2, 100087.	1.5	5
5	Hospital chaplains' communication with patients: Characteristics, functions and potential benefits. <i>Patient Education and Counseling</i> , 2022, 105, 2905-2912.	2.2	8
6	Reducing the Single IRB Burden: Streamlining Electronic IRB Systems. <i>AJOB Empirical Bioethics</i> , 2021, 12, 33-40.	1.6	4
7	Typologies and Meanings of Prayer Among Patients. <i>Journal of Religion and Health</i> , 2021, , 1.	1.7	3
8	How Artistic Representation Can Inform Current Debates About Chimeras. <i>Journal of Medical Humanities</i> , 2021, 42, 337-343.	0.7	0
9	Preparing for the Next Generation of Ethical Challenges Concerning Heritable Human Genome Editing. <i>American Journal of Bioethics</i> , 2021, 21, 1-4.	0.9	0
10	Experiment on identical siblings separated at birth: ethical implications for researchers, universities, and archives today. <i>Journal of Medical Ethics</i> , 2021, 47, 657-661.	1.8	0
11	Roles of genetics and blood type in clinical responses to COVID-19: ethical and policy concerns. <i>Journal of Medical Ethics</i> , 2021, 47, 149-151.	1.8	0
12	Doctor, Will You Pray for Me? Responding to Patients' Religious and Spiritual Concerns. <i>Academic Medicine</i> , 2021, 96, 349-354.	1.6	5
13	Impact of patient education videos on genetic counseling outcomes after exome sequencing. <i>Patient Education and Counseling</i> , 2020, 103, 127-135.	2.2	18
14	Bioethics of Translating Limited Evidence into Clinical Practice: Case Study of the Cerebroplacental Ratio. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2020, 42, 1154-1157.	0.7	0
15	Needs to Prepare for "Post-COVID-19 Syndrome". <i>American Journal of Bioethics</i> , 2020, 20, 4-6.	0.9	6
16	Legal Immunity for Physicians During the COVID-19 Pandemic. <i>Chest</i> , 2020, 158, 1343-1345.	0.8	8
17	Local Knowledge and Single IRBs for Multisite Studies: Challenges and Solutions. <i>Ethics & Human Research</i> , 2019, 41, 22-31.	0.9	12
18	Evaluation of the cost and effectiveness of diverse recruitment methods for a genetic screening study. <i>Genetics in Medicine</i> , 2019, 21, 2371-2380.	2.4	10

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19	How Single Institutional Review Boards Manage Their Own Conflicts of Interest. <i>Academic Medicine</i> , 2019, 94, 1554-1560.	1.6	3
20	Infertility providers'™ and patients'™ views and experiences concerning doctor shopping in the USA. <i>Human Fertility</i> , 2019, 22, 238-245.	1.7	8
21	Designing Babies. , 2019, , .		15
22	Challenges, Dilemmas and Factors Involved in PGD Decision Making: Providers'™ and Patients'™ Views, Experiences and Decisions. <i>Journal of Genetic Counseling</i> , 2018, 27, 909-919.	1.6	19
23	How Infertility Patients and Providers View and Confront Religious and Spiritual Issues. <i>Journal of Religion and Health</i> , 2018, 57, 223-239.	1.7	7
24	Bringing science and advocacy together to address health needs of people who inject drugs. <i>Journal of Medical Ethics</i> , 2018, 44, 165-166.	1.8	1
25	Gatekeepers for infertility treatment? Views of ART providers concerning referrals by non-ART providers. <i>Reproductive Biomedicine and Society Online</i> , 2018, 5, 17-30.	1.8	7
26	Impact of Receiving Secondary Results from Genomic Research: A 12-Month Longitudinal Study. <i>Journal of Genetic Counseling</i> , 2018, 27, 709-722.	1.6	26
27	Reliance agreements and single IRB review of multisite research: Concerns of IRB members and staff. <i>AJOB Empirical Bioethics</i> , 2018, 9, 164-172.	1.6	17
28	Impediments to communication and relationships between infertility care providers and patients. <i>BMC Women's Health</i> , 2018, 18, 84.	2.0	21
29	Predictive testing and clinical trials in Huntington's disease: An ethical analysis. <i>Movement Disorders</i> , 2018, 33, 243-247.	3.9	3
30	Unconventional combinations of prospective parents: ethical challenges faced by IVF providers. <i>BMC Medical Ethics</i> , 2017, 18, 18.	2.4	16
31	“Will they be good enough parents?” Ethical dilemmas, views, and decisions among assisted reproductive technology (ART) providers. <i>AJOB Empirical Bioethics</i> , 2017, 8, 253-265.	1.6	7
32	How much is a child worth? Providers'™ and patients'™ views and responses concerning ethical and policy challenges in paying for ART. <i>PLoS ONE</i> , 2017, 12, e0171939.	2.5	28
33	Buying and selling human eggs: infertility providers'™ ethical and other concerns regarding egg donor agencies. <i>BMC Medical Ethics</i> , 2016, 17, 71.	2.4	18
34	Struggles in Defining and Addressing Requests for “Family Balancing”: Ethical Issues Faced by Providers and Patients. <i>Journal of Law, Medicine and Ethics</i> , 2016, 44, 616-629.	0.9	12
35	How old is too old? Challenges faced by clinicians concerning age cutoffs for patients undergoing in vitro fertilization. <i>Fertility and Sterility</i> , 2016, 106, 216-224.	1.0	31
36	Deciding how many embryos to transfer: ongoing challenges and dilemmas. <i>Reproductive Biomedicine and Society Online</i> , 2016, 3, 1-15.	1.8	30

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37	Reducing the number of fetuses in a pregnancy: providers' and patients' views of challenges. <i>Human Reproduction</i> , 2016, 31, 2570-2576.	0.9	18
38	Reviewing <scp>HIV</scp>-Related Research in Emerging Economies: The Role of Government Reviewing Agencies. <i>Developing World Bioethics</i> , 2016, 16, 4-14.	0.9	2
39	Ethical and Psychosocial Considerations in Informing HIV-Exposed Uninfected Children That They Were Exposed to HIV and Antiretroviral Medications In Utero. <i>American Journal of Public Health</i> , 2016, 106, 1390-1396.	2.7	2
40	Pain during embryo transfer is independently associated with clinical pregnancy in fresh/frozen assisted reproductive technology cycles. <i>Journal of Obstetrics and Gynaecology Research</i> , 2016, 42, 684-693.	1.3	13
41	How Agencies Market Egg Donation on the Internet: A Qualitative Study. <i>Journal of Law, Medicine and Ethics</i> , 2015, 43, 610-618.	0.9	18
42	Evolving Challenges and Research-Needs Concerning Ebola. <i>American Journal of Public Health</i> , 2015, 105, 1513-1515.	2.7	10
43	The Need for Vigilance in the Marketing of Genomic Tests in Psychiatry. <i>Journal of Nervous and Mental Disease</i> , 2015, 203, 809-810.	1.0	4
44	Linguistic and Cultural Challenges in Communication and Translation in US-Sponsored HIV Prevention Research in Emerging Economies. <i>PLoS ONE</i> , 2015, 10, e0133394.	2.5	11
45	New Challenges for Electronic Health Records. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 29.	7.4	59
46	Creating and selling embryos for "donation": ethical challenges. <i>American Journal of Obstetrics and Gynecology</i> , 2015, 212, 167-170.e1.	1.3	7
47	Controversies concerning mitochondrial replacement therapy. <i>Fertility and Sterility</i> , 2015, 103, 344-346.	1.0	13
48	Consenting for Molecular Diagnostics. <i>Clinical Chemistry</i> , 2015, 61, 139-141.	3.2	3
49	Association of Researcher Characteristics with Views on Return of Incidental Findings from Genomic Research. <i>Journal of Genetic Counseling</i> , 2015, 24, 833-841.	1.6	17
50	Electronic Health Records and Adolescent Privacy"Reply. <i>JAMA - Journal of the American Medical Association</i> , 2015, 313, 1373.	7.4	1
51	Kamakahi vs ASRM and the future of compensation for human eggs. <i>American Journal of Obstetrics and Gynecology</i> , 2015, 213, 186-187.e1.	1.3	9
52	Informed consent for return of incidental findings in genomic research. <i>Genetics in Medicine</i> , 2014, 16, 367-373.	2.4	58
53	How US institutional review boards decide when researchers need to translate studies. <i>Journal of Medical Ethics</i> , 2014, 40, 193-197.	1.8	7
54	Models of Consent to Return of Incidental Findings in Genomic Research. <i>Hastings Center Report</i> , 2014, 44, 22-32.	1.0	72

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55	Psychiatrists's Views of the Genetic Bases of Mental Disorders and Behavioral Traits and Their Use of Genetic Tests. <i>Journal of Nervous and Mental Disease</i> , 2014, 202, 530-538.	1.0	18
56	Should Life Insurers Have Access to Genetic Test Results?. <i>JAMA - Journal of the American Medical Association</i> , 2014, 312, 1855.	7.4	11
57	Processes and factors involved in decisions regarding return of incidental genomic findings in research. <i>Genetics in Medicine</i> , 2014, 16, 311-317.	2.4	28
58	Advocating for longitudinal follow-up of the health and welfare of egg donors. <i>Fertility and Sterility</i> , 2014, 102, 662-666.	1.0	23
59	Use of Genetic Tests among Neurologists and Psychiatrists: Knowledge, Attitudes, Behaviors, and Needs for Training. <i>Journal of Genetic Counseling</i> , 2014, 23, 156-163.	1.6	87
60	Views of preimplantation genetic diagnosis among psychiatrists and neurologists. <i>Journal of reproductive medicine, The</i> , 2014, 59, 385-92.	0.2	10
61	Egg donation brokers: an analysis of agency versus in vitro fertilization clinic websites. <i>Journal of reproductive medicine, The</i> , 2014, 59, 534-41.	0.2	6
62	Return of Secondary Genomic Findings vs Patient Autonomy. <i>JAMA - Journal of the American Medical Association</i> , 2013, 310, 369.	7.4	68
63	Attitudes and Practices Among Internists Concerning Genetic Testing. <i>Journal of Genetic Counseling</i> , 2013, 22, 90-100.	1.6	147
64	Views of internists towards uses of PGD. <i>Reproductive BioMedicine Online</i> , 2013, 26, 142-147.	2.4	21
65	Researchers's views on return of incidental genomic research results: qualitative and quantitative findings. <i>Genetics in Medicine</i> , 2013, 15, 888-895.	2.4	103
66	How good does the science have to be in proposals submitted to Institutional Review Boards? An Interview Study of Institutional Review Board personnel. <i>Clinical Trials</i> , 2013, 10, 761-766.	1.6	18
67	How IRBs View and Make Decisions about Social Risks. <i>Journal of Empirical Research on Human Research Ethics</i> , 2013, 8, 58-65.	1.3	25
68	How IRBs view and make decisions about coercion and undue influence: Table 1. <i>Journal of Medical Ethics</i> , 2013, 39, 224-229.	1.8	50
69	Views of IRBs Concerning Their Local Ecologies: Perceptions of Relationships, Systems, and Tensions Between IRBs and Their Institutions. <i>American Journal of Bioethics Primary Research</i> , 2013, 4, 31-43.	1.5	2
70	How IRBs View and Make Decisions about Consent Forms. <i>Journal of Empirical Research on Human Research Ethics</i> , 2013, 8, 8-19.	1.3	21
71	How IRB leaders view and approach challenges raised by industry-funded research. <i>IRB: Ethics & Human Research</i> , 2013, 35, 9-17.	0.8	3
72	Local IRBs vs. Federal Agencies: Shifting Dynamics, Systems, and Relationships. <i>Journal of Empirical Research on Human Research Ethics</i> , 2012, 7, 50-62.	1.3	6

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73	To Protect Human Subjects, Review <i>What Was Done</i> , Not Proposed. <i>Science</i> , 2012, 335, 1576-1577.	12.6	11
74	Vermeer and Leeuwenhoek, <i>Figments of the Imagination?</i> . <i>FASEB Journal</i> , 2012, 26, 2238-2238.	0.5	0
75	Institutional Review Board Community Members. <i>Academic Medicine</i> , 2012, 87, 975-981.	1.6	56
76	US IRBS CONFRONTING RESEARCH IN THE DEVELOPING WORLD. <i>Developing World Bioethics</i> , 2012, 12, 63-73.	0.9	36
77	Recruiting egg donors online: an analysis of in vitro fertilization clinic and agency websites' adherence to American Society for Reproductive Medicine guidelines. <i>Fertility and Sterility</i> , 2012, 98, 995-1000.	1.0	61
78	From anonymity to "open doors": IRB responses to tensions with researchers. <i>BMC Research Notes</i> , 2012, 5, 347.	1.4	18
79	The Myth of Community Differences as the Cause of Variations Among IRBs. <i>American Journal of Bioethics Primary Research</i> , 2011, 2, 24-33.	1.5	31
80	THE REPORTING OF IRB REVIEW IN JOURNAL ARTICLES PRESENTING HIV RESEARCH CONDUCTED IN THE DEVELOPING WORLD. <i>Developing World Bioethics</i> , 2011, 11, 161-169.	0.9	8
81	Views and Experiences of IRBs concerning Research Integrity. <i>Journal of Law, Medicine and Ethics</i> , 2011, 39, 513-528.	0.9	27
82	"In Sickness and in Health": Disclosures of Genetic Risks in Dating. <i>Journal of Genetic Counseling</i> , 2011, 20, 98-112.	1.6	21
83	How local IRBs view central IRBs in the US. <i>BMC Medical Ethics</i> , 2011, 12, 13.	2.4	39
84	HIV/AIDS Research Conducted in the Developing World and Sponsored by the Developed World: Reporting of Research Ethics Committee Review in Two Countries. <i>Journal of Empirical Research on Human Research Ethics</i> , 2011, 6, 83-91.	1.3	8
85	"Members of the Same Club": Challenges and Decisions Faced by US IRBs in Identifying and Managing Conflicts of Interest. <i>PLoS ONE</i> , 2011, 6, e22796.	2.5	27
86	The Ethics Police?: IRBs' Views Concerning Their Power. <i>PLoS ONE</i> , 2011, 6, e28773.	2.5	43
87	Views of Discrimination among Individuals Confronting Genetic Disease. <i>Journal of Genetic Counseling</i> , 2010, 19, 68-83.	1.6	57
88	Misunderstandings Concerning Genetics Among Patients Confronting Genetic Disease. <i>Journal of Genetic Counseling</i> , 2010, 19, 430-446.	1.6	57
89	The process of deciding about prophylactic surgery for breast and ovarian cancer: Patient questions, uncertainties, and communication. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 52-66.	1.2	41
90	Exclusion of Genetic Information From the Medical Record. <i>JAMA - Journal of the American Medical Association</i> , 2010, 304, 1120.	7.4	21

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91	Disclosures of funding sources and conflicts of interest in published HIV/AIDS research conducted in developing countries. <i>Journal of Medical Ethics</i> , 2010, 36, 505-510.	1.8	10
92	The Use of Eggs and Embryos in Stem Cell Research. <i>Seminars in Reproductive Medicine</i> , 2010, 28, 336-344.	1.1	9
93	Defining Neuromarketing: Practices and Professional Challenges. <i>Harvard Review of Psychiatry</i> , 2010, 18, 230-237.	2.1	150
94	The Impact of Social Contexts in Testing for Alpha-1 Antitrypsin Deficiency: The Roles of Physicians and Others. <i>Genetic Testing and Molecular Biomarkers</i> , 2009, 13, 269-276.	0.7	12
95	“Am I my genes?” Questions of identity among individuals confronting genetic disease. <i>Genetics in Medicine</i> , 2009, 11, 880-889.	2.4	38
96	Voluntariness of Consent to Research: A Conceptual Model. <i>Hastings Center Report</i> , 2009, 39, 30-39.	1.0	83
97	Preimplantation genetic diagnosis on in vitro fertilization clinic websites: presentations of risks, benefits and other information. <i>Fertility and Sterility</i> , 2009, 92, 1276-1283.	1.0	17
98	Payment of egg donors in stem cell research in the USA. <i>Reproductive BioMedicine Online</i> , 2009, 18, 603-608.	2.4	34
99	Voluntariness of consent to research: a preliminary empirical investigation. <i>IRB: Ethics & Human Research</i> , 2009, 31, 10-4.	0.8	21
100	VIEWES OF THE PROCESS AND CONTENT OF ETHICAL REVIEWS OF HIV VACCINE TRIALS AMONG MEMBERS OF US INSTITUTIONAL REVIEW BOARDS AND SOUTH AFRICAN RESEARCH ETHICS COMMITTEES. <i>Developing World Bioethics</i> , 2008, 8, 207-218.	0.9	23
101	Anticipating issues related to increasing preimplantation genetic diagnosis use: a research agenda. <i>Reproductive BioMedicine Online</i> , 2008, 17, 33-42.	2.4	23
102	Kuru fieldwork in 1981 and beyond. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2008, 363, 3646-3647.	4.0	1
103	Ethical issues concerning disclosures of HIV diagnoses to perinatally infected children and adolescents. <i>Journal of Clinical Ethics</i> , 2008, 19, 31-42.	0.3	18
104	Disclosure of information to potential subjects on research recruitment web sites. <i>IRB: Ethics & Human Research</i> , 2008, 30, 15-20.	0.8	10
105	The roles of family members, health care workers, and others in decision-making processes about genetic testing among individuals at risk for Huntington disease. <i>Genetics in Medicine</i> , 2007, 9, 358-371.	2.4	33
106	Additional Implications of a National Survey on Ethics Consultation in United States Hospitals. <i>American Journal of Bioethics</i> , 2007, 7, 47-48.	0.9	2
107	The Reporting of Monetary Compensation in Research Articles. <i>Journal of Empirical Research on Human Research Ethics</i> , 2007, 2, 61-67.	1.3	14
108	Disclosures of Huntington disease risk within families: Patterns of decision-making and implications. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 1835-1849.	1.2	47

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109	“Patient-time”, “doctor-time”, and “institution-time” Perceptions and definitions of time among doctors who become patients. <i>Patient Education and Counseling</i> , 2007, 66, 147-155.	2.2	34
110	Decision-Making About Reproductive Choices Among Individuals At-Risk for Huntington's Disease. <i>Journal of Genetic Counseling</i> , 2007, 16, 347-362.	1.6	63
111	From “Male Bonding Rituals” to “Suicide Tuesday”. <i>Journal of Homosexuality</i> , 2006, 51, 7-32.	2.0	12
112	“Post-Residency Disease” and the Medical Self: Identity, Work, and Health Care Among Doctors Who Become Patients. <i>Perspectives in Biology and Medicine</i> , 2006, 49, 542-552.	0.5	16
113	Improving Education on Doctor-Patient Relationships and Communication: Lessons from Doctors Who Become Patients. <i>Academic Medicine</i> , 2006, 81, 447-453.	1.6	62
114	Views and approaches toward risks and benefits among doctors who become patients. <i>Patient Education and Counseling</i> , 2006, 64, 61-68.	2.2	15
115	Disclosures of illness by doctors to their patients: A qualitative study of doctors with HIV and other serious disorders. <i>Patient Education and Counseling</i> , 2006, 64, 277-284.	2.2	17
116	Qualifying Confidentiality: Historical and Empirical Issues and Facts. <i>American Journal of Bioethics</i> , 2006, 6, 26-27.	0.9	5
117	Antoni Van Leeuwenhoek, FRS on Vermeer: a figment of the imagination. <i>FASEB Journal</i> , 2006, 20, 591-594.	0.5	0
118	Problems in comprehension of informed consent in rural and peri-urban Mali, West Africa. <i>Clinical Trials</i> , 2006, 3, 306-313.	1.6	91
119	Challenges and changes in spirituality among doctors who become patients. <i>Social Science and Medicine</i> , 2005, 61, 2396-2406.	3.8	27
120	Contexts, Anyone?: The Need for Contextualization in the Debate About the Moral Status of Embryos. <i>American Journal of Bioethics</i> , 2005, 5, 56-58.	0.9	9
121	Naming names: Perceptions of name-based HIV reporting, partner notification, and criminalization of non-disclosure among persons living with HIV. <i>Sexuality Research and Social Policy</i> , 2004, 1, 38-57.	2.3	19
122	Patterns of Communication Between Gay and Lesbian Patients and Their Health Care Providers. <i>Journal of Homosexuality</i> , 2002, 42, 65-75.	2.0	118
123	MDMA (“ecstasy”) use, and its association with high risk behaviors, mental health, and other factors among gay/bisexual men in New York City. <i>Drug and Alcohol Dependence</i> , 2002, 66, 115-125.	3.2	138
124	HIV and the Law: Integrating Law, Policy, and Social Epidemiology. <i>Journal of Law, Medicine and Ethics</i> , 2002, 30, 533-547.	0.9	16
125	MDMA (“Ecstasy”) Abuse and High-Risk Sexual Behaviors Among 169 Gay and Bisexual Men. <i>American Journal of Psychiatry</i> , 2000, 157, 1162-1164.	7.2	130
126	Self-Disclosure of HIV Status to Sexual Partners: A Qualitative Study of Issues Faced by Gay Men. <i>Journal of the Gay and Lesbian Medical Association</i> , 1999, 3, 39-49.	0.6	31

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127	Sexual orientation and associated characteristics among north american academic psychiatrists. Journal of Sex Research, 1998, 35, 282-287.	2.5	2
128	The Natural Incubation Period of Kuru and the Episodes of Transmission in Three Clusters of Patients. Neuroepidemiology, 1984, 3, 3-20.	2.3	96