

Daniel Strech

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

116
papers

1,881
citations

25
h-index

38
g-index

167
ext. papers

2,509
ext. citations

3.5
avg, IF

5.33
L-index

#	Paper	IF	Citations
116	Results publications are inadequately linked to trial registrations: An automated pipeline and evaluation of German university medical centers.. <i>Clinical Trials</i> , 2022 , 17407745221087456	2.2	0
115	Do German university medical centres promote robust and transparent research? A cross-sectional study of institutional policies.. <i>Health Research Policy and Systems</i> , 2022 , 20, 39	3.7	1
114	Data protection-compliant broad consent for secondary use of health care data and human biosamples for (bio)medical research: towards a new German national standard. <i>Journal of Biomedical Informatics</i> , 2022 , 104096	10.2	1
113	Results availability and timeliness of registered COVID-19 clinical trials: interim cross-sectional results from the DIRECCT study. <i>BMJ Open</i> , 2021 , 11, e053096	3	0
112	Results dissemination from completed clinical trials conducted at German university medical centers remained delayed and incomplete. The 2014-2017 cohort.. <i>Journal of Clinical Epidemiology</i> , 2021 ,	5.7	3
111	Pandemien und Forschungsethik. Eine Übersicht zu zentralen Herausforderungen 2021 , 163-178		
110	Investigator brochures for phase I/II trials lack information on the robustness of preclinical safety studies. <i>British Journal of Clinical Pharmacology</i> , 2021 , 87, 2723-2731	3.8	3
109	The full spectrum of ethical issues in dementia research: findings of a systematic qualitative review. <i>BMC Medical Ethics</i> , 2021 , 22, 32	2.9	2
108	Web-based survey among animal researchers on publication practices and incentives for increasing publication rates. <i>PLoS ONE</i> , 2021 , 16, e0250362	3.7	1
107	Robust preclinical evidence in somatic cell genome editing: A key driver of responsible and efficient therapeutic innovations. <i>Drug Discovery Today</i> , 2021 , 26, 2238-2243	8.8	1
106	Details of risk-benefit communication in informed consent documents for phase I/II trials. <i>Clinical Trials</i> , 2021 , 18, 71-80	2.2	2
105	Reporting of prior clinical studies in Investigator Brochures did not adhere to the basic principles of evidence synthesis: a cross-sectional study. <i>Journal of Clinical Epidemiology</i> , 2021 , 130, 87-95	5.7	0
104	Improving the trustworthiness, usefulness, and ethics of biomedical research through an innovative and comprehensive institutional initiative. <i>PLoS Biology</i> , 2020 , 18, e3000576	9.7	12
103	Attitudes towards animal study registries and their characteristics: An online survey of three cohorts of animal researchers. <i>PLoS ONE</i> , 2020 , 15, e0226443	3.7	3
102	Umgang mit Mittelbegrenzung im Gesundheitswesen 2020 , 751-770		
101	Implementation of data access and use procedures in clinical data warehouses. A systematic review of literature and publicly available policies. <i>BMC Medical Informatics and Decision Making</i> , 2020 , 20, 157	3.6	2
100	Results dissemination of registered clinical trials across Polish academic institutions: a cross-sectional analysis. <i>BMJ Open</i> , 2020 , 10, e034666	3	4

99	Broad donor consent for human biobanks in Germany and Europe: a strategy to facilitate cross-border sharing and exchange of human biological materials and related data. <i>Journal of Laboratory Medicine</i> , 2019 , 43, 291-299	0.9	2
98	Result dissemination from clinical trials conducted at German university medical centers was delayed and incomplete. <i>Journal of Clinical Epidemiology</i> , 2019 , 115, 37-45	5.7	19
97	3Rs missing: animal research without scientific value is unethical.. <i>BMJ Open Science</i> , 2019 , 3,	4.6	20
96	Publication rates in animal research. Extent and characteristics of published and non-published animal studies followed up at two German university medical centres. <i>PLoS ONE</i> , 2019 , 14, e0223758	3.7	14
95	Practice evaluation of biobank ethics and governance: current needs and future perspectives. <i>Journal of Medical Genetics</i> , 2019 , 56, 176-185	5.8	7
94	The spectrum of ethical issues in a Learning Health Care System: a systematic qualitative review. <i>International Journal for Quality in Health Care</i> , 2018 , 30, 161-168	1.9	17
93	Preclinical efficacy studies in investigator brochures: Do they enable risk-benefit assessment?. <i>PLoS Biology</i> , 2018 , 16, e2004879	9.7	39
92	Standards of practice in empirical bioethics research: towards a consensus. <i>BMC Medical Ethics</i> , 2018 , 19, 68	2.9	37
91	The Public Awareness of and Attitude Toward Research Biobanks - A Regional German Survey. <i>Frontiers in Genetics</i> , 2018 , 9, 190	4.5	10
90	Routinely collected data for randomized trials: promises, barriers, and implications. <i>Trials</i> , 2018 , 19, 29	2.8	51
89	Current practices for access, compensation, and prioritization in biobanks. Results from an interview study. <i>European Journal of Human Genetics</i> , 2018 , 26, 1572-1581	5.3	9
88	Tracking the timely dissemination of clinical studies. Characteristics and impact of 10 tracking variables. <i>F1000Research</i> , 2018 , 7, 1863	3.6	2
87	5. Ethik und rechtliche Fragen 2018 , 157-186		
86	Why are so few patients rating their physicians on German physician rating websites? A qualitative study. <i>BMC Health Services Research</i> , 2018 , 18, 670	2.9	7
85	The bench is closer to the bedside than we think: Uncovering the ethical ties between preclinical researchers in translational neuroscience and patients in clinical trials. <i>PLoS Biology</i> , 2018 , 16, e2006343	9.7	15
84	Preclinical efficacy in therapeutic area guidelines from the U.S. Food and Drug Administration and the European Medicines Agency: a cross-sectional study. <i>British Journal of Pharmacology</i> , 2018 , 175, 4229-4238	8.6	11
83	Off-label use, compassionate use und individuelle Heilversuche: ethische Implikationen zulassungserschreitender Arzneimittelanwendungen 2017 , 95-105		3
82	Access policies in biobank research: what criteria do they include and how publicly available are they? A cross-sectional study. <i>European Journal of Human Genetics</i> , 2017 , 25, 293-300	5.3	21

81	Participatory improvement of a template for informed consent documents in biobank research - study results and methodological reflections. <i>BMC Medical Ethics</i> , 2017 , 18, 78	2.9	16
80	An integrated conceptual framework for evaluating and improving understanding of informed consent. <i>Trials</i> , 2017 , 18, 482	2.8	8
79	What methods do reviews of normative ethics literature use for search, selection, analysis, and synthesis? In-depth results from a systematic review of reviews. <i>Systematic Reviews</i> , 2017 , 6, 261	3	5
78	Ethical issues in obesity prevention for school children: a systematic qualitative review. <i>International Journal of Public Health</i> , 2017 , 62, 981-988	4	3
77	Ethical issues in public health surveillance: a systematic qualitative review. <i>BMC Public Health</i> , 2017 , 17, 295	4.1	32
76	Developments in the Frequency of Ratings and Evaluation Tendencies: A Review of German Physician Rating Websites. <i>Journal of Medical Internet Research</i> , 2017 , 19, e299	7.6	12
75	Public Awareness and Use of German Physician Ratings Websites: Cross-Sectional Survey of Four North German Cities. <i>Journal of Medical Internet Research</i> , 2017 , 19, e387	7.6	12
74	The full spectrum of ethical issues in the care of patients with ALS: a systematic qualitative review. <i>Journal of Neurology</i> , 2016 , 263, 201-209	5.5	16
73	Kommentar I zum Fall: Der Patientenwille als oberste Instanz – Schwierige Umsetzung in der klinischen Forschung – <i>Ethik in Der Medizin</i> , 2016 , 28, 153-155	0.5	
72	Current state of ethics literature synthesis: a systematic review of reviews. <i>BMC Medicine</i> , 2016 , 14, 152	11.4	28
71	Input analysis for two public consultations on the EU Clinical Trials Regulation. <i>Health Research Policy and Systems</i> , 2016 , 14, 69	3.7	
70	Collective agency and the concept of public involvement: A practice-oriented analysis. <i>BMC Medical Ethics</i> , 2016 , 17, 1	2.9	36
69	The contribution and attitudes of research ethics committees to complete registration and non-selective reporting of clinical trials: A European survey. <i>Research Ethics</i> , 2016 , 12, 123-136	2.8	1
68	Bias in dissemination of clinical research findings: structured OPEN framework of what, who and why, based on literature review and expert consensus. <i>BMJ Open</i> , 2016 , 6, e010024	3	11
67	Animal Study Registries: Results from a Stakeholder Analysis on Potential Strengths, Weaknesses, Facilitators, and Barriers. <i>PLoS Biology</i> , 2016 , 14, e2000391	9.7	16
66	The Full Spectrum of Clinical Ethical Issues in Kidney Failure. Findings of a Systematic Qualitative Review. <i>PLoS ONE</i> , 2016 , 11, e0149357	3.7	9
65	Umgang mit Mittelbegrenzung im Gesundheitswesen 2016 , 1-20		
64	Applying for, reviewing and funding public health research in Germany and beyond. <i>Health Research Policy and Systems</i> , 2016 , 14, 43	3.7	5

63	Do Public Involvement Activities in Biomedical Research and Innovation Recruit Representatively? A Systematic Qualitative Review. <i>Public Health Genomics</i> , 2016 , 19, 193-202	1.9	6
62	Ethics Reporting in Biospecimen and Genetic Research: Current Practice and Suggestions for Changes. <i>PLoS Biology</i> , 2016 , 14, e1002521	9.7	2
61	A template for broad consent in biobank research. Results and explanation of an evidence and consensus-based development process. <i>European Journal of Medical Genetics</i> , 2016 , 59, 295-309	2.6	35
60	Ethical review of biobank research: Should RECs review each release of material from biobanks operating under an already-approved broad consent and data protection model?. <i>European Journal of Medical Genetics</i> , 2015 , 58, 545-9	2.6	9
59	Research Guideline Recommendations for Broad Consent Forms in Biobank Research and How They Are Currently Addressed in Practice. <i>American Journal of Bioethics</i> , 2015 , 15, 60-3	1.1	5
58	Clinical Ethics in Gabon: The Spectrum of Clinical Ethical Issues Based on Findings from In-Depth Interviews at Three Public Hospitals. <i>PLoS ONE</i> , 2015 , 10, e0132374	3.7	8
57	Putting public health ethics into practice: a systematic framework. <i>Frontiers in Public Health</i> , 2015 , 3, 23	6	40
56	Evidence-informed recommendations to reduce dissemination bias in clinical research: conclusions from the OPEN (Overcome failure to Publish nEgative fiNDings) project based on an international consensus meeting. <i>BMJ Open</i> , 2015 , 5, e006666	3	35
55	Ethics Literacy and "Ethics University": Two Intertwined Models for Public Involvement and Empowerment in Bioethics. <i>Frontiers in Public Health</i> , 2015 , 3, 287	6	0
54	Do editorial policies support ethical research? A thematic text analysis of author instructions in psychiatry journals. <i>PLoS ONE</i> , 2014 , 9, e97492	3.7	7
53	Current practice of public involvement activities in biomedical research and innovation: a systematic qualitative review. <i>PLoS ONE</i> , 2014 , 9, e113274	3.7	17
52	When Choosing Wisely meets clinical practice guidelines. <i>Zeitschrift Fur Evidenz, Fortbildung Und Qualitat Im Gesundheitswesen</i> , 2014 , 108, 601-3	1.4	12
51	Did we describe what you meant? Findings and methodological discussion of an empirical validation study for a systematic review of reasons. <i>BMC Medical Ethics</i> , 2014 , 15, 69	2.9	5
50	International requirements for consent in biobank research: qualitative review of research guidelines. <i>Journal of Medical Genetics</i> , 2014 , 51, 773-81	5.8	14
49	Systematic and transparent inclusion of ethical issues and recommendations in clinical practice guidelines: a six-step approach. <i>Implementation Science</i> , 2014 , 9, 184	8.4	8
48	How can bedside rationing be justified despite coexisting inefficiency? The need for benchmarks of efficiency. <i>Journal of Medical Ethics</i> , 2014 , 40, 89-93	2.5	10
47	Which public to involve? More reflection on collective agency and sufficient representativeness is needed. <i>American Journal of Bioethics</i> , 2014 , 14, 31-3	1.1	
46	Participation rate or informed choice? Rethinking the European key performance indicators for mammography screening. <i>Health Policy</i> , 2014 , 115, 100-3	3.2	11

45	Extent of non-publication in cohorts of studies approved by research ethics committees or included in trial registries. <i>PLoS ONE</i> , 2014 , 9, e114023	3.7	115
44	Dissemination bias in systematic reviews of animal research: a systematic review. <i>PLoS ONE</i> , 2014 , 9, e116016	3.7	16
43	Wissenschaftlicher und sozialer Wert 2014 , 299-303		
42	Evidence-based health information from the users perspective--a qualitative analysis. <i>BMC Health Services Research</i> , 2013 , 13, 405	2.9	16
41	Inclusion of ethical issues in dementia guidelines: a thematic text analysis. <i>PLoS Medicine</i> , 2013 , 10, e1001408	10.8	18
40	The full spectrum of ethical issues in dementia care: systematic qualitative review. <i>British Journal of Psychiatry</i> , 2013 , 202, 400-6	5.4	40
39	Practice variation across consent templates for biobank research. a survey of German biobanks. <i>Frontiers in Genetics</i> , 2013 , 4, 240	4.5	13
38	How psychiatry journals support the unbiased translation of clinical research. A cross-sectional study of editorial policies. <i>PLoS ONE</i> , 2013 , 8, e75995	3.7	25
37	The impact of social media on medical professionalism: a systematic qualitative review of challenges and opportunities. <i>Journal of Medical Internet Research</i> , 2013 , 15, e184	7.6	77
36	Systematic Reviews of Reasons in Public Health Ethics Literature: A Roadmap. <i>Public Health Ethics Analysis</i> , 2013 , 119-138	0.2	
35	Where Public Health Meets Ethics. Conceptual Foundations and Practical Challenges of Public Health. <i>Public Health Ethics Analysis</i> , 2013 , 5-23	0.2	
34	The need for systematic reviews of reasons. <i>Bioethics</i> , 2012 , 26, 315-28	2	35
33	Normative arguments and new solutions for the unbiased registration and publication of clinical trials. <i>Journal of Clinical Epidemiology</i> , 2012 , 65, 276-81	5.7	25
32	How to write a systematic review of reasons. <i>Journal of Medical Ethics</i> , 2012 , 38, 121-6	2.5	78
31	Why the "appraisal of guidelines for research and evaluation" instrument can and should further inform ethics policy work. <i>American Journal of Bioethics</i> , 2012 , 12, 25-7	1.1	5
30	Public-Health-Ethik 2012 , 137-142		0
29	Zur Ethik einer restriktiven Regulierung der Studienregistrierung. <i>Ethik in Der Medizin</i> , 2011 , 23, 177-189	0.5	2
28	Improved evidence will support sound decision making but will not solve the problem of cost containment in oncology. <i>Journal of Clinical Oncology</i> , 2011 , 29, 3943	2.2	0

27	Reasons Why Post-Trial Access to Trial Drugs Should, or Need not be Ensured to Research Participants: A Systematic Review. <i>Public Health Ethics</i> , 2011 , 4, 160-184	1.8	59
26	Quality of ethical guidelines and ethical content in clinical guidelines: the example of end-of-life decision-making. <i>Journal of Medical Ethics</i> , 2011 , 37, 390-6	2.5	23
25	How to evaluate conflict of interest policies. <i>American Journal of Bioethics</i> , 2011 , 11, 37-9	1.1	6
24	Ethical principles for physician rating sites. <i>Journal of Medical Internet Research</i> , 2011 , 13, e113	7.6	42
23	Quality of reporting of randomized controlled trials of pharmacologic treatment of bipolar disorders: a systematic review. <i>Journal of Clinical Psychiatry</i> , 2011 , 72, 1214-21	4.6	14
22	Priorisierung und Rationierung im Gesundheitswesen - Status quo und Perspektiven 2011 , 51-61		3
21	Kostensensible Leitlinien als Priorisierungsinstrument 2011 , 75-98		1
20	Zur Abgrenzung zwischen Interessenkonflikten, Fehlverhalten und professioneller Integrität 2011 , 81-87		
19	How factual do we want the facts? Criteria for a critical appraisal of empirical research for use in ethics. <i>Journal of Medical Ethics</i> , 2010 , 36, 222-5	2.5	6
18	The role of ethics committees and ethics consultation in allocation decisions: a 4-stage process. <i>Medical Care</i> , 2010 , 48, 821-6	3.1	10
17	The representation of patient experience and satisfaction in physician rating sites. A criteria-based analysis of English- and German-language sites. <i>BMC Health Services Research</i> , 2010 , 10, 332	2.9	53
16	Herausforderungen expliziter Leistungsbegrenzungen durch kostensensible Leitlinien. Ergebnisse einer qualitativen Interviewstudie mit leitenden Klinikärzten. <i>Gesundheitsökonomie Und Qualitätsmanagement</i> , 2009 , 14, 38-43	0.3	7
15	Zuclopenthixol dihydrochloride for schizophrenia. <i>Schizophrenia Bulletin</i> , 2009 , 35, 855-6	1.3	4
14	Are physicians willing to ration health care? Conflicting findings in a systematic review of survey research. <i>Health Policy</i> , 2009 , 90, 113-24	3.2	60
13	Value judgments in the analysis and synthesis of evidence. <i>Journal of Clinical Epidemiology</i> , 2008 , 61, 521-4	5.7	46
12	How physicians allocate scarce resources at the bedside: a systematic review of qualitative studies. <i>Journal of Medicine and Philosophy</i> , 2008 , 33, 80-99	1.1	81
11	Systematic reviews of empirical bioethics. <i>Journal of Medical Ethics</i> , 2008 , 34, 472-7	2.5	45
10	Evidenz-basierte Ethik. <i>Ethik in Der Medizin</i> , 2008 , 20, 274-286	0.5	5

9	Eztliches Handeln bei Mittelknappheit. <i>Ethik in Der Medizin</i> , 2008 , 20, 94-109	0.5	26
8	BMBF-Klausurwoche: Clinical Ethics Consultation: theories & methods [Implementation] evaluation. <i>Ethik in Der Medizin</i> , 2008 , 20, 148-150	0.5	1
7	Evidence-based ethics--what it should be and what it shouldn't. <i>BMC Medical Ethics</i> , 2008 , 9, 16	2.9	18
6	Implizite Rationierung als Rechtsproblem Ergebnisse einer qualitativen Interviewstudie zur Situation in deutschen Krankenhäusern. <i>Medizinrecht</i> , 2007 , 25, 703-706	0	3
5	Zuclopenthixol dihydrochloride for schizophrenia. <i>The Cochrane Library</i> , 2005 , CD005474	5.2	10
4	Der Umgang mit Wahrscheinlichkeiten und das Vertrauen in die Medizin. <i>Ethik in Der Medizin</i> , 2005 , 17, 103-113	0.5	3
3	Challenges and proposed solutions in making clinical research on COVID-19 ethical. A status quo analysis across German research ethics committees		1
2	Results Availability and Timeliness of Registered COVID-19 Clinical Trials: A Cross-Sectional Study		1
1	Results dissemination from completed clinical trials conducted at German university medical centers remains delayed and incomplete. The 2014-2017 cohort		2