

# Eva C Winkler

## List of Publications by Year in descending order

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Version: 2024-02-01

43  
papers

1,183  
citations

567281

15  
h-index

414414

32  
g-index

59  
all docs

59  
docs citations

59  
times ranked

1766  
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluating requests for physician-assisted suicide. A survey among German oncologists. <i>Cancer Medicine</i> , 2023, 12, 1813-1820.	2.8	3
2	Sekundärnutzung klinischer Daten aus der Patientenversorgung für Forschungszwecke – Eine qualitative Interviewstudie zu Nutzen- und Risikopotenzialen aus Sicht von Expertinnen und Experten für den deutschen Forschungskontext. , 2022, , 185-210.		8
3	Sekundärnutzung klinischer Daten in datensammelnden, nicht-interventionellen Forschungs- oder Lernaktivitäten – Begriff, Studientypen und ethische Herausforderungen. , 2022, , 71-98.		2
4	Patients' Willingness to Provide Their Clinical Data for Research Purposes and Acceptance of Different Consent Models: Findings From a Representative Survey of Patients With Cancer. <i>Journal of Medical Internet Research</i> , 2022, 24, e37665.	4.3	12
5	Often Relatives are the Key – Family Involvement in Treatment Decision Making in Patients with Advanced Cancer Near the End of Life. <i>Oncologist</i> , 2021, 26, e831-e837.	3.7	12
6	Nipping Diseases in the Bud? Ethical and Social Considerations of the Concept of "Disease Interception". <i>Public Health Ethics</i> , 2021, 14, 100-108.	1.0	0
7	Secondary Use of Clinical Data in Data-Gathering, Non-Interventional Research or Learning Activities: Definition, Types, and a Framework for Risk Assessment. <i>Journal of Medical Internet Research</i> , 2021, 23, e26631.	4.3	16
8	Development and Evaluation of a Decision Aid to Support Patients' Participatory Decision-Making for Tumor-Specific and Palliative Therapy for Advanced Cancer: Protocol for a Pre-Post Study. <i>JMIR Research Protocols</i> , 2021, 10, e24954.	1.0	0
9	Patient-Reported Financial Distress in Cancer: A Systematic Review of Risk Factors in Universal Healthcare Systems. <i>Cancers</i> , 2021, 13, 5015.	3.7	18
10	Is Dupras and Bunnik's Framework for Assessing Privacy Risks in Multi-Omic Research and Databases Still Too Exceptionalist?. <i>American Journal of Bioethics</i> , 2021, 21, 80-82.	0.9	2
11	GA4GH: International policies and standards for data sharing across genomic research and healthcare. <i>Cell Genomics</i> , 2021, 1, 100029.	6.5	94
12	The COVID-19 Pandemic and Cancer Patients in Germany: Impact on Treatment, Follow-Up Care and Psychological Burden. <i>Frontiers in Public Health</i> , 2021, 9, 788598.	2.7	14
13	Caregivers' role in using a personal electronic health record: a qualitative study of cancer patients and caregivers in Germany. <i>BMC Medical Informatics and Decision Making</i> , 2020, 20, 158.	3.0	9
14	Late decisions about treatment limitation in patients with cancer: empirical analysis of end-of-life practices in a haematology and oncology unit at a German university hospital. <i>ESMO Open</i> , 2020, 5, e000950.	4.5	6
15	The patient-level effect of the cost of Cancer care – financial burden in German Cancer patients. <i>BMC Cancer</i> , 2020, 20, 529.	2.6	30
16	Do patients and research subjects have a right to receive their genomic raw data? An ethical and legal analysis. <i>BMC Medical Ethics</i> , 2020, 21, 7.	2.4	24
17	Digitalizing Health Services by Implementing a Personal Electronic Health Record in Germany: Qualitative Analysis of Fundamental Prerequisites From the Perspective of Selected Experts. <i>Journal of Medical Internet Research</i> , 2020, 22, e15102.	4.3	25
18	Cancer Patients' Preferences for either Quality of Life or a Longer Life Determine Their Willingness to Talk about Forgoing Cancer-Specific Treatment. <i>Oncology Research and Treatment</i> , 2019, 42, 458-465.	1.2	5

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19	Patients' Preferences in Non-Curable Cancer Disease. <i>Oncology Research and Treatment</i> , 2019, 42, 31-34.	1.2	11
20	Do Researchers in Empirical Ethics Studies Have a Duty to Act Upon their Findings? Case Study in End-of-Life Decision Making. <i>Journal of Empirical Research on Human Research Ethics</i> , 2019, 14, 438-440.	1.3	3
21	Between Minimal and Greater Than Minimal Risk: How Research Participants and Oncologists Assess Data-Sharing and the Risk of Re-identification in Genomic Research. <i>Philosophy and Technology</i> , 2019, 32, 39-55.	4.3	1
22	The second patient? Family members of cancer patients and their role in end-of-life decision making. <i>BMC Palliative Care</i> , 2018, 17, 29.	1.8	39
23	“Rather one more chemo than one less” Oncologists and Oncology Nurses’ Reasons for Aggressive Treatment of Young Adults with Advanced Cancer. <i>Oncologist</i> , 2018, 23, 256-262.	3.7	15
24	High prevalence of moral distress reported by oncologists and oncology nurses in end-of-life decision making. <i>Psycho-Oncology</i> , 2018, 27, 2733-2739.	2.3	58
25	Trust and responsibility in molecular tumour boards. <i>Bioethics</i> , 2018, 32, 464-472.	1.4	5
26	Utilizing a Prototype Patient-Controlled Electronic Health Record in Germany: Qualitative Analysis of User-Reported Perceptions and Perspectives. <i>JMIR Formative Research</i> , 2018, 2, e10411.	1.4	16
27	Development and Evaluation of an Ethical Guideline for Decisions to Limit Life-Prolonging Treatment in Advanced Cancer: Protocol for a Monocentric Mixed-Method Interventional Study. <i>JMIR Research Protocols</i> , 2018, 7, e157.	1.0	2
28	Applying systems biology to biomedical research and health care: a redefining definition of systems medicine. <i>BMC Health Services Research</i> , 2017, 17, 761.	2.2	24
29	Complexity of care and strategies of self-management in patients with colorectal cancer. <i>Patient Preference and Adherence</i> , 2017, Volume 11, 731-742.	1.8	21
30	A Personal Electronic Health Record: Study Protocol of a Feasibility Study on Implementation in a Real-World Health Care Setting. <i>JMIR Research Protocols</i> , 2017, 6, e33.	1.0	22
31	Researchers’ Duty to Share Pre-publication Data: From the Prima Facie Duty to Practice. <i>Law, Governance and Technology Series</i> , 2016, , 309-337.	0.4	5
32	So rare we need to hunt for them: reframing the ethical debate on incidental findings. <i>Genome Medicine</i> , 2015, 7, 83.	8.2	19
33	Ethical challenges of whole genome sequencing in translational research and answers by the EURAT-project. <i>Laboratoriums Medizin</i> , 2015, 38, .	0.6	1
34	Stakeholders’ perspectives on biobank-based genomic research: systematic review of the literature. <i>European Journal of Human Genetics</i> , 2015, 23, 1607-1614.	2.8	61
35	What Keeps Oncologists From Addressing Palliative Care Early on With Incurable Cancer Patients? An Active Stance Seems Key. <i>Oncologist</i> , 2015, 20, 56-61.	3.7	84
36	When do we need to care about the caregiver? Supportive care needs, anxiety, and depression among informal caregivers of patients with cancer and cancer survivors. <i>Cancer</i> , 2015, 121, 1513-1519.	4.1	295

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37	Ethische Herausforderungen der Genomsequenzierung in der translationalen Forschung und Antworten aus dem EURAT-Projekt. <i>Laboratoriums Medizin</i> , 2014, 38, .	0.6	1
38	Assessing patientâ€“caregiver communication in cancerâ€“a psychometric validation of the Cancer Communication Assessment Tool (CCAT-PF) in a German sample. <i>Supportive Care in Cancer</i> , 2014, 22, 2473-2478.	2.2	9
39	Cancer Patients' Preferences for Quantity or Quality of Life: German Translation and Validation of the Quality and Quantity Questionnaire. <i>Oncology Research and Treatment</i> , 2014, 37, 472-478.	1.2	16
40	<i>Onkologische Ethik.</i> , 2013, , 479-495.		0
41	Evaluating a patient's request for life-prolonging treatment: an ethical framework. <i>Journal of Medical Ethics</i> , 2012, 38, 647-651.	1.8	27
42	Ethical assessment of life-prolonging treatment. <i>Lancet Oncology</i> , The, 2011, 12, 720-722.	10.7	26
43	Patient Involvement in Decisions to Limit Treatment: The Crucial Role of Agreement Between Physician and Patient. <i>Journal of Clinical Oncology</i> , 2009, 27, 2225-2230.	1.6	102