

# Karen E Steinhauser

## List of Publications by Year in descending order

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Version: 2024-02-01

45  
papers

5,673  
citations

279798

23  
h-index

254184

43  
g-index

45  
all docs

45  
docs citations

45  
times ranked

4473  
citing authors

#	ARTICLE	IF	CITATIONS
1	Who Benefits Most? A Multisite Study of Coaching and Resident Well-being. <i>Journal of General Internal Medicine</i> , 2022, 37, 539-547.	2.6	12
2	Triadic agreement about advanced cancer treatment decisions: Perceptions among patients, families, and oncologists. <i>Patient Education and Counseling</i> , 2022, 105, 982-986.	2.2	12
3	Clinician End-of-Life Experiences With Pediatric Muslim Patients at a US Quaternary Care Center. <i>Journal of Pain and Symptom Management</i> , 2022, 63, 673-679.	1.2	2
4	Spirituality in Patients With Heart Failure. <i>JACC: Heart Failure</i> , 2022, 10, 217-226.	4.1	12
5	Spirituality in Serious Illness and Health. <i>JAMA - Journal of the American Medical Association</i> , 2022, 328, 184.	7.4	88
6	A novel decision aid for acute myeloid leukemia: a feasibility and preliminary efficacy trial. <i>Supportive Care in Cancer</i> , 2021, 29, 3563-3569.	2.2	9
7	Coordinating assessment of spiritual needs: a cross-walk of narrative and psychometric assessment tools used in palliative care. <i>Journal of Health Care Chaplaincy</i> , 2021, , 1-14.	1.1	1
8	Exploring Culturally Responsive Religious and Spirituality Health Care Communications among African Americans with Advanced Heart Failure, Their Family Caregivers, and Clinicians. <i>Journal of Palliative Medicine</i> , 2021, 24, 1798-1806.	1.1	1
9	Enhancing meaning in the face of advanced cancer and pain: Qualitative evaluation of a meaning-centered psychosocial pain management intervention. <i>Palliative and Supportive Care</i> , 2020, 18, 263-270.	1.0	21
10	Current measures of distress may not account for what's most important in existential care interventions: Results of the outlook trial. <i>Palliative and Supportive Care</i> , 2020, 18, 648-657.	1.0	4
11	Reducing Disparities in the Quality of Palliative Care for Older African Americans through Improved Advance Care Planning: Study Design and Protocol. <i>Journal of Palliative Medicine</i> , 2019, 22, S-90-S-100.	1.1	27
12	Comparing internal and external validation in the discovery of qualitative treatment-subgroup effects using two small clinical trials. <i>Contemporary Clinical Trials Communications</i> , 2019, 15, 100372.	1.1	1
13	African Americans With Advanced Heart Failure Are More Likely to Die in a Health Care Facility Than at Home or in Hospice: An Analysis From the PAL-HF Trial. <i>Journal of Cardiac Failure</i> , 2019, 25, 693-694.	1.7	9
14	Evaluating the Feasibility and Acceptability of a Telehealth Program in a Rural Palliative Care Population: TapCloud for Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 7-14.	1.2	75
15	Themes Addressed by Couples With Advanced Cancer During a Communication Skills Training Intervention. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 252-258.	1.2	9
16	Palliative care and hospital readmissions in patients with advanced heart failure: Insights from the PAL-HF trial. <i>American Heart Journal</i> , 2018, 204, 202-204.	2.7	14
17	Patient experiences of acute myeloid leukemia: A qualitative study about diagnosis, illness understanding, and treatment decision-making. <i>Psycho-Oncology</i> , 2017, 26, 2063-2068.	2.3	87
18	It Is Like Heart Failure. It Is Chronic and It Will Kill You: A Qualitative Analysis of Burnout Among Hospice and Palliative Care Clinicians. <i>Journal of Pain and Symptom Management</i> , 2017, 53, 901-910.e1.	1.2	66

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19	Randomized trial of an uncertainty self-management telephone intervention for patients awaiting liver transplant. <i>Patient Education and Counseling</i> , 2017, 100, 509-517.	2.2	27
20	State of the Science of Spirituality and Palliative Care Research Part I: Definitions, Measurement, and Outcomes. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 428-440.	1.2	192
21	State of the Science of Spirituality and Palliative Care Research: Research Landscape and Future Directions. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 426-427.	1.2	18
22	State of the Science of Spirituality and Palliative Care Research Part II: Screening, Assessment, and Interventions. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 441-453.	1.2	125
23	Addressing Patient Emotional and Existential Needs During Serious Illness: Results of the Outlook Randomized Controlled Trial. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 898-908.	1.2	36
24	Palliative Care in Heart Failure. <i>Journal of the American College of Cardiology</i> , 2017, 70, 331-341.	2.8	425
25	Engaging patients and families to create a feasible clinical trial integrating palliative and heart failure care: results of the ENABLE CHF-PC pilot clinical trial. <i>BMC Palliative Care</i> , 2017, 16, 45.	1.8	51
26	Why Burnout Is So Hard to Fix. <i>Journal of Oncology Practice</i> , 2017, 13, 348-351.	2.5	14
27	The feasibility and acceptability of a chaplain-led intervention for caregivers of seriously ill patients: A Caregiver Outlook pilot study. <i>Palliative and Supportive Care</i> , 2016, 14, 456-467.	1.0	21
28	Building Resilience for Palliative Care Clinicians: An Approach to Burnout Prevention Based on Individual Skills and Workplace Factors. <i>Journal of Pain and Symptom Management</i> , 2016, 52, 284-291.	1.2	107
29	The elephant in the room: Facilitating conversations about advanced cancer between patients and their spouses.. <i>Journal of Clinical Oncology</i> , 2016, 34, 233-233.	1.6	3
30	Transcending differences to study the transcendent: an exploratory study of researchers' and chaplains' reflections on interdisciplinary spiritual care research collaboration. <i>BMC Palliative Care</i> , 2015, 14, 12.	1.8	4
31	Symptom burden, quality of life, and distress in acute myeloid leukemia patients receiving induction chemotherapy: A prospective electronic patient-reported outcomes study.. <i>Journal of Clinical Oncology</i> , 2015, 33, e20702-e20702.	1.6	1
32	The Palliative Care in Heart Failure Trial: Rationale and design. <i>American Heart Journal</i> , 2014, 168, 645-651.e1.	2.7	45
33	Validation of a Measure of Family Experience of Patients with Serious Illness: The QUAL-E (Fam). <i>Journal of Pain and Symptom Management</i> , 2014, 48, 1168-1181.	1.2	15
34	Prognostic Disclosure in Acute Myeloid Leukemia (AML): A Qualitative Study of Patient Preferences and Physician Practices. <i>Blood</i> , 2014, 124, 1280-1280.	1.4	0
35	Provision of Spiritual Support to Patients With Advanced Cancer by Religious Communities and Associations With Medical Care at the End of Life. <i>JAMA Internal Medicine</i> , 2013, 173, 1109.	5.1	242
36	Editorial: Pairing self-management with Palliative Care: intervening in life-limiting illness. <i>Journal of Nursing and Healthcare of Chronic Illness</i> , 2011, 3, 1-3.	0.5	14

#	ARTICLE	IF	CITATIONS
37	Do Preparation and Life Completion Discussions Improve Functioning and Quality of Life in Seriously Ill Patients? Pilot Randomized Control Trial. <i>Journal of Palliative Medicine</i> , 2008, 11, 1234-1240.	1.1	142
38	“Are You at Peace?” <i>Archives of Internal Medicine</i> , 2006, 166, 101.	3.8	179
39	Identifying, recruiting, and retaining seriously-ill patients and their caregivers in longitudinal research. <i>Palliative Medicine</i> , 2006, 20, 745-754.	3.1	177
40	Measuring End-of-Life Care Outcomes Prospectively. <i>Journal of Palliative Medicine</i> , 2005, 8, s-30-s-41.	1.1	42
41	Measuring quality of life at the end of life: Validation of the QUAL-E. <i>Palliative and Supportive Care</i> , 2004, 2, 3-14.	1.0	157
42	Evolution in Measuring the Quality of Dying. <i>Journal of Palliative Medicine</i> , 2002, 5, 407-414.	1.1	72
43	Initial Assessment of a New Instrument to Measure Quality of Life at the End of Life. <i>Journal of Palliative Medicine</i> , 2002, 5, 829-841.	1.1	139
44	In Search of a Good Death: Observations of Patients, Families, and Providers. <i>Annals of Internal Medicine</i> , 2000, 132, 825.	3.9	898
45	Factors Considered Important at the End of Life by Patients, Family, Physicians, and Other Care Providers. <i>JAMA - Journal of the American Medical Association</i> , 2000, 284, 2476.	7.4	2,077