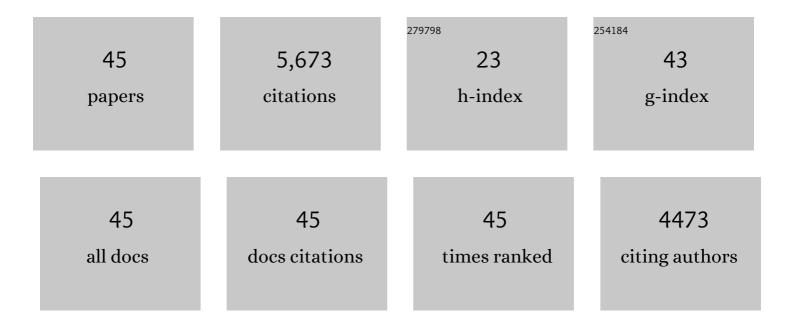
## Karen E Steinhauser

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Factors Considered Important at the End of Life by Patients, Family, Physicians, and Other Care Providers. JAMA - Journal of the American Medical Association, 2000, 284, 2476.	7.4	2,077
2	In Search of a Good Death: Observations of Patients, Families, and Providers. Annals of Internal Medicine, 2000, 132, 825.	3.9	898
3	Palliative Care in Heart Failure. Journal of the American College of Cardiology, 2017, 70, 331-341.	2.8	425
4	Provision of Spiritual Support to Patients With Advanced Cancer by Religious Communities and Associations With Medical Care at the End of Life. JAMA Internal Medicine, 2013, 173, 1109.	5.1	242
5	State of the Science of Spirituality and Palliative Care Research Part I: Definitions, Measurement, and Outcomes. Journal of Pain and Symptom Management, 2017, 54, 428-440.	1.2	192
6	"Are You at Peace?― Archives of Internal Medicine, 2006, 166, 101.	3.8	179
7	Identifying, recruiting, and retaining seriously-ill patients and their caregivers in longitudinal research. Palliative Medicine, 2006, 20, 745-754.	3.1	177
8	Measuring quality of life at the end of life: Validation of the QUAL-E. Palliative and Supportive Care, 2004, 2, 3-14.	1.0	157
9	Do Preparation and Life Completion Discussions Improve Functioning and Quality of Life in Seriously Ill Patients? Pilot Randomized Control Trial. Journal of Palliative Medicine, 2008, 11, 1234-1240.	1.1	142
10	Initial Assessment of a New Instrument to Measure Quality of Life at the End of Life. Journal of Palliative Medicine, 2002, 5, 829-841.	1.1	139
11	State of the Science of Spirituality and Palliative Care Research Part II: Screening, Assessment, and Interventions. Journal of Pain and Symptom Management, 2017, 54, 441-453.	1.2	125
12	Building Resilience for Palliative Care Clinicians: An Approach to Burnout Prevention Based on Individual Skills and Workplace Factors. Journal of Pain and Symptom Management, 2016, 52, 284-291.	1.2	107
13	Spirituality in Serious Illness and Health. JAMA - Journal of the American Medical Association, 2022, 328, 184.	7.4	88
14	Patient experiences of acute myeloid leukemia: A qualitative study about diagnosis, illness understanding, and treatment decisionâ€making. Psycho-Oncology, 2017, 26, 2063-2068.	2.3	87
15	Evaluating the Feasibility and Acceptability of a Telehealth Program in a Rural Palliative Care Population: TapCloud for Palliative Care. Journal of Pain and Symptom Management, 2018, 56, 7-14.	1.2	75
16	Evolution in Measuring the Quality of Dying. Journal of Palliative Medicine, 2002, 5, 407-414.	1.1	72
17	"lt Is Like Heart Failure. It Is Chronic … and It Will Kill You― A Qualitative Analysis of Burnout Among Hospice and Palliative Care Clinicians. Journal of Pain and Symptom Management, 2017, 53, 901-910.e1.	1.2	66
18	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. BMC Palliative Care, 2017, 16, 45.	1.8	51

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19	The Palliative Care in Heart Failure Trial: Rationale and design. American Heart Journal, 2014, 168, 645-651.e1.	2.7	45
20	Measuring End-of-Life Care Outcomes Prospectively. Journal of Palliative Medicine, 2005, 8, s-30-s-41.	1.1	42
21	Addressing Patient Emotional and Existential Needs During Serious Illness: Results of the Outlook Randomized Controlled Trial. Journal of Pain and Symptom Management, 2017, 54, 898-908.	1.2	36
22	Randomized trial of an uncertainty self-management telephone intervention for patients awaiting liver transplant. Patient Education and Counseling, 2017, 100, 509-517.	2.2	27
23	Reducing Disparities in the Quality of Palliative Care for Older African Americans through Improved Advance Care Planning: Study Design and Protocol. Journal of Palliative Medicine, 2019, 22, S-90-S-100.	1.1	27
24	The feasibility and acceptability of a chaplain-led intervention for caregivers of seriously ill patients: A Caregiver Outlook pilot study. Palliative and Supportive Care, 2016, 14, 456-467.	1.0	21
25	Enhancing meaning in the face of advanced cancer and pain: Qualitative evaluation of a meaning-centered psychosocial pain management intervention. Palliative and Supportive Care, 2020, 18, 263-270.	1.0	21
26	State of the Science of Spirituality and Palliative Care Research: Research Landscape and Future Directions. Journal of Pain and Symptom Management, 2017, 54, 426-427.	1.2	18
27	Validation of a Measure of Family Experience of Patients with Serious Illness: The QUAL-E (Fam). Journal of Pain and Symptom Management, 2014, 48, 1168-1181.	1.2	15
28	Editorial: Pairing selfâ€management with Palliative Care: intervening in lifeâ€limiting illness. Journal of Nursing and Healthcare of Chronic Illness, 2011, 3, 1-3.	0.5	14
29	Why Burnout Is So Hard to Fix. Journal of Oncology Practice, 2017, 13, 348-351.	2.5	14
30	Palliative care and hospital readmissions in patients with advanced heart failure: Insights from the PAL-HF trial. American Heart Journal, 2018, 204, 202-204.	2.7	14
31	Who Benefits Most? A Multisite Study of Coaching and Resident Well-being. Journal of General Internal Medicine, 2022, 37, 539-547.	2.6	12
32	Triadic agreement about advanced cancer treatment decisions: Perceptions among patients, families, and oncologists. Patient Education and Counseling, 2022, 105, 982-986.	2.2	12
33	Spirituality in Patients With HeartÂFailure. JACC: Heart Failure, 2022, 10, 217-226.	4.1	12
34	Themes Addressed by Couples With Advanced Cancer During a Communication Skills Training Intervention. Journal of Pain and Symptom Management, 2018, 56, 252-258.	1.2	9
35	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. Journal of Cardiac Failure, 2019, 25, 693-694.	1.7	9
36	A novel decision aid for acute myeloid leukemia: a feasibility and preliminary efficacy trial. Supportive Care in Cancer, 2021, 29, 3563-3569.	2.2	9

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37	Transcending differences to study the transcendent: an exploratory study of researchers' and chaplains' reflections on interdisciplinary spiritual care research collaboration. BMC Palliative Care, 2015, 14, 12.	1.8	4
38	Current measures of distress may not account for what's most important in existential care interventions: Results of the outlook trial. Palliative and Supportive Care, 2020, 18, 648-657.	1.0	4
39	The elephant in the room: Facilitating conversations about advanced cancer between patients and their spouses Journal of Clinical Oncology, 2016, 34, 233-233.	1.6	3
40	Clinician End-of-Life Experiences With Pediatric Muslim Patients at a US Quaternary Care Center. Journal of Pain and Symptom Management, 2022, 63, 673-679.	1.2	2
41	Comparing internal and external validation in the discovery of qualitative treatment-subgroup effects using two small clinical trials. Contemporary Clinical Trials Communications, 2019, 15, 100372.	1.1	1
42	Coordinating assessment of spiritual needs: a cross-walk of narrative and psychometric assessment tools used in palliative care. Journal of Health Care Chaplaincy, 2021, , 1-14.	1.1	1
43	Exploring Culturally Responsive Religious and Spirituality Health Care Communications among African Americans with Advanced Heart Failure, Their Family Caregivers, and Clinicians. Journal of Palliative Medicine, 2021, 24, 1798-1806.	1.1	1
44	Symptom burden, quality of life, and distress in acute myeloid leukemia patients receiving induction chemotherapy: A prospective electronic patient-reported outcomes study Journal of Clinical Oncology, 2015, 33, e20702-e20702.	1.6	1
45	Prognostic Disclosure in Acute Myeloid Leukemia (AML): A Qualitative Study of Patient Preferences and Physician Practices. Blood, 2014, 124, 1280-1280.	1.4	Ο