

Sally A Norton

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

2,794
citations

218662
26
h-index

182417
51
g-index

62
all docs

62
docs citations

62
times ranked

2422
citing authors

#	ARTICLE	IF	CITATIONS
1	Completion of Patient-Reported Outcome Questionnaires Among Older Adults with Advanced Cancer. <i>Journal of Pain and Symptom Management</i> , 2022, 63, 301-310.	1.2	10
2	The shared uncertainty experience of older adults with advanced cancer and their caregivers. <i>Psycho-Oncology</i> , 2022, 31, 1041-1049.	2.3	4
3	Protocol paper: Multi-site, cluster-randomized clinical trial for optimizing functional outcomes of older cancer survivors after chemotherapy. <i>Journal of Geriatric Oncology</i> , 2022, , .	1.0	0
4	Conceptualizing and Counting Discretionary Utilization in the Final 100 Days of Life: A Scoping Review. <i>Journal of Pain and Symptom Management</i> , 2020, 59, 894-915.e14.	1.2	15
5	Family caregiver descriptions of stopping chemotherapy and end-of-life transitions. <i>Supportive Care in Cancer</i> , 2019, 27, 669-675.	2.2	35
6	Effects of the Values and Options in Cancer Care Communication Intervention on Personal Caregiver Experiences of Cancer Care and Bereavement Outcomes. <i>Journal of Palliative Medicine</i> , 2019, 22, 1394-1400.	1.1	17
7	Physician and Patient Characteristics Associated With More Intensive End-of-Life Care. <i>Journal of Pain and Symptom Management</i> , 2019, 58, 208-215.e1.	1.2	13
8	The contagion of optimism: The relationship between patient optimism and palliative care clinician overestimation of survival among hospitalized patients with advanced cancer. <i>Psycho-Oncology</i> , 2019, 28, 1286-1292.	2.3	12
9	End-of-Life Preferences, Length-of-Life Conversations, and Hospice Enrollment in Palliative Care: A Direct Observation Cohort Study among People with Advanced Cancer. <i>Journal of Palliative Medicine</i> , 2019, 22, 152-156.	1.1	15
10	When chemotherapy fails: Emotionally charged experiences faced by family caregivers of patients with advanced cancer. <i>Patient Education and Counseling</i> , 2019, 102, 909-915.	2.2	22
11	“Treat us with dignity”: a qualitative study of the experiences and recommendations of lesbian, gay, bisexual, transgender, and queer (LGBTQ) patients with cancer. <i>Supportive Care in Cancer</i> , 2019, 27, 2525-2532.	2.2	80
12	Racial/ethnic differences in prognosis communication during initial inpatient palliative care consultations among people with advanced cancer. <i>Patient Education and Counseling</i> , 2019, 102, 1098-1103.	2.2	31
13	Palliative Care Clinician Overestimation of Survival in Advanced Cancer: Disparities and Association With End-of-Life Care. <i>Journal of Pain and Symptom Management</i> , 2019, 57, 233-240.	1.2	31
14	Is Annual Income a Predictor of Completion of Advance Directives (ADs) in Patients With Cancer. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 402-407.	1.4	15
15	An integrated operational definition and conceptual model of asthma self-management in teens. <i>Journal of Asthma</i> , 2018, 55, 1315-1327.	1.7	18
16	Staff Experiences Forming and Sustaining Palliative Care Teams in Nursing Homes. <i>Gerontologist</i> , The, 2018, 58, e218-e225.	3.9	17
17	Feeling Heard and Understood in the Hospital Environment: Benchmarking Communication Quality Among Patients With Advanced Cancer Before and After Palliative Care Consultation. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 239-244.	1.2	27
18	Preference for Palliative Care in Cancer Patients: Are Men and Women Alike?. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 1-6.e1.	1.2	75

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19	Parental coping in the context of having a child who is facing death: A theoretical framework. Palliative and Supportive Care, 2018, 16, 432-441.	1.0	15
20	Impact of Nursing Home Palliative Care Teams on End-of-Life Outcomes. Medical Care, 2018, 56, 11-18.	2.4	28
21	Distress Due to Prognostic Uncertainty in Palliative Care: Frequency, Distribution, and Outcomes among Hospitalized Patients with Advanced Cancer. Journal of Palliative Medicine, 2018, 21, 315-321.	1.1	42
22	Perceptions and experiences underlying self-management and reporting of symptoms in teens with asthma. Journal of Asthma, 2017, 54, 143-152.	1.7	36
23	Improving palliative care through teamwork (IMPACTT) in nursing homes: Study design and baseline findings. Contemporary Clinical Trials, 2017, 56, 1-8.	1.8	10
24	The Role of Hospice and Palliative Care Nurses in Quality Improvement. Journal of Hospice and Palliative Nursing, 2017, 19, 160-165.	0.9	9
25	Feeling Heard and Understood: A Patient-Reported Quality Measure for the Inpatient Palliative Care Setting. Journal of Pain and Symptom Management, 2016, 51, 150-154.	1.2	48
26	New approaches to qualitative interviewing: Development of a card sort technique to understand subjective patterns of symptoms and responses. International Journal of Nursing Studies, 2016, 58, 90-96.	5.6	19
27	Methodological Research Priorities in Palliative Care and Hospice Quality Measurement. Journal of Pain and Symptom Management, 2016, 51, 155-162.	1.2	30
28	Measuring What Matters: Top-Ranked Quality Indicators for Hospice and Palliative Care From the American Academy of Hospice and Palliative Medicine and Hospice and Palliative Nurses Association. Journal of Pain and Symptom Management, 2015, 49, 773-781.	1.2	185
29	Design of, and enrollment in, the palliative care communication research initiative: a direct-observation cohort study. BMC Palliative Care, 2015, 14, 40.	1.8	25
30	Goal Communication in Palliative Care Decision-Making Consultations. Journal of Pain and Symptom Management, 2015, 50, 701-706.	1.2	17
31	Using music[al] knowledge to represent expressions of emotions. Patient Education and Counseling, 2015, 98, 1339-1345.	2.2	8
32	Developing Palliative Care Practice Guidelines and Standards for Nursing Home-Based Palliative Care Teams: A Delphi Study. Journal of the American Medical Directors Association, 2015, 16, 86.e1-86.e7.	2.5	30
33	Emotional Distress and Compassionate Responses in Palliative Care Decision-Making Consultations. Journal of Palliative Medicine, 2014, 17, 579-584.	1.1	24
34	A Place to Get Worse. Journal of Hospice and Palliative Nursing, 2014, 16, 338-345.	0.9	12
35	Direct Observation of Prognosis Communication in Palliative Care: A Descriptive Study. Journal of Pain and Symptom Management, 2013, 45, 202-212.	1.2	29
36	Patient and family members' perceptions of palliative care in heart failure. Heart and Lung: Journal of Acute and Critical Care, 2013, 42, 112-119.	1.6	50

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37	“That Don’t Work for Me” Journal of Hospice and Palliative Nursing, 2013, 15, 177-182.	0.9	8
38	Latent Classes of Prognosis Conversations in Palliative Care: A Mixed-Methods Study. Journal of Palliative Medicine, 2013, 16, 653-660.	1.1	21
39	Living with advanced heart failure or COPD: Experiences and goals of individuals nearing the end of life. Research in Nursing and Health, 2013, 36, 349-358.	1.6	35
40	Palliative Care Communication: Linking Patients' Prognoses, Values, and Goals of Care. Research in Nursing and Health, 2013, 36, 582-590.	1.6	35
41	End-of-Life Quality-of-Care Measures for Nursing Homes: Place of Death and Hospice. Journal of Palliative Medicine, 2012, 15, 438-446.	1.1	25
42	Who Is Attending? End-of-Life Decision Making in the Intensive Care Unit. Journal of Palliative Medicine, 2012, 15, 56-62.	1.1	35
43	Family Members’ Informal Roles in End-of-Life Decision Making in Adult Intensive Care Units. American Journal of Critical Care, 2012, 21, 43-51.	1.6	62
44	Internet use by parents of infants with positive newborn screens. Journal of Inherited Metabolic Disease, 2012, 35, 879-884.	3.6	28
45	Navigating Tensions: Integrating Palliative Care Consultation Services Into an Academic Medical Center Setting. Journal of Pain and Symptom Management, 2011, 42, 680-690.	1.2	21
46	Parents' Experiences of Expanded Newborn Screening Evaluations. Pediatrics, 2011, 128, 53-61.	2.1	62
47	Meaning and Practice of Palliative Care for Hospitalized Older Adults with Life Limiting Illnesses. Journal of Aging Research, 2011, 2011, 1-8.	0.9	5
48	Measuring End-of-Life Care Processes in Nursing Homes. Gerontologist, The, 2009, 49, 803-815.	3.9	33
49	“What Bothers You the Most?” Initial Responses From Patients Receiving Palliative Care Consultation. American Journal of Hospice and Palliative Medicine, 2008, 25, 88-92.	1.4	15
50	Challenges in Conducting End-of-Life Research in Critical Care. AACN Advanced Critical Care, 2008, 19, 170-177.	1.1	7
51	Challenges in Conducting End-of-Life Research in Critical Care. AACN Advanced Critical Care, 2008, 19, 170-177.	1.1	21
52	Proactive palliative care in the medical intensive care unit: Effects on length of stay for selected high-risk patients. Critical Care Medicine, 2007, 35, 1530-1535.	0.9	915
53	Intensive care unit cultures and end-of-life decision making. Journal of Critical Care, 2007, 22, 159-168.	2.2	71
54	What is Most Important for You to Achieve?: An Analysis of Patient Responses When Receiving Palliative Care Consultation. Journal of Palliative Medicine, 2006, 9, 382-388.	1.1	34

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55	Striving for Congruence: The Interconnection Between Values, Practice, and Political Action. Policy, Politics, and Nursing Practice, 2005, 6, 20-29.	1.8	12
56	The dying patient in the ICU: role of the interdisciplinary team. Critical Care Clinics, 2004, 20, 525-540.	2.6	55
57	Life Support Withdrawal: Communication and Conflict. American Journal of Critical Care, 2003, 12, 548-555.	1.6	71
58	Life support withdrawal: communication and conflict. American Journal of Critical Care, 2003, 12, 548-55.	1.6	22
59	Working toward consensus: Providers' strategies to shift patients from curative to palliative treatment choices. Research in Nursing and Health, 2001, 24, 258-269.	1.6	63
60	Moral reasoning among graduate students in nursing. Nursing Outlook, 2000, 48, 73-80.	2.6	12
61	Facilitating End-of-Life Decision-Making: Strategies for Communicating and Assessing. Journal of Gerontological Nursing, 2000, 26, 6-9.	0.6	52
62	An exploration of methodological pluralism in nursing research. Research in Nursing and Health, 1998, 21, 545-555.	1.6	20