Karla T Washington

List of Publications by Year in descending order

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Version: 2024-02-01

257450 361022 127 1,992 24 35 citations g-index h-index papers 127 127 127 1915 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Information needs of informal caregivers of older adults with chronic health conditions. Patient Education and Counseling, 2011, 83, 37-44.	2.2	126
2	Barriers to Hospice Use among African Americans: A Systematic Review. Health and Social Work, 2008, 33, 267-274.	1.0	62
3	Barriers to Pain Management: Caregiver Perceptions and Pain Talk by Hospice Interdisciplinary Teams. Journal of Pain and Symptom Management, 2008, 36, 374-382.	1.2	57
4	A Noninferiority Trial of a Problem-Solving Intervention for Hospice Caregivers: In Person versus Videophone. Journal of Palliative Medicine, 2012, 15, 653-660.	1.1	52
5	The Role of Human Factors in Telehealth. Telemedicine Journal and E-Health, 2010, 16, 446-453.	2.8	48
6	A Problem Solving Intervention for Hospice Caregivers: A Pilot Study. Journal of Palliative Medicine, 2010, 13, 1005-1011.	1.1	47
7	The Effect of Internet Group Support for Caregivers on Social Support, Self-Efficacy, and Caregiver Burden: A Meta-Analysis. Telemedicine Journal and E-Health, 2017, 23, 621-629.	2.8	44
8	A Systematic Review of the Evidence Base for Telehospice. Telemedicine Journal and E-Health, 2012, 18, 38-47.	2.8	41
9	Understanding Social Support Burden Among Family Caregivers. Health Communication, 2014, 29, 901-910.	3.1	41
10	Hospice Caregivers' Experiences With Pain Management: "l'm Not a Doctor, and I Don't Know if I Helped Her Go Faster or Slower― Journal of Pain and Symptom Management, 2013, 46, 846-858.	1.2	40
11	Online social support groups for informal caregivers of hospice patients with cancer. European Journal of Oncology Nursing, 2020, 44, 101698.	2.1	40
12	Pain Management Concerns From the Hospice Family Caregivers' Perspective. American Journal of Hospice and Palliative Medicine, 2018, 35, 601-611.	1.4	39
13	Stress Variances Among Informal Hospice Caregivers. Qualitative Health Research, 2012, 22, 1114-1125.	2.1	38
14	Hospice Caregiver Depression: The Evidence Surrounding the Greatest Pain of All. Journal of Social Work in End-of-Life and Palliative Care, 2013, 9, 256-271.	0.6	36
15	The Prevalence and Risks for Depression and Anxiety in Hospice Caregivers. Journal of Palliative Medicine, 2017, 20, 366-371.	1.1	36
16	Use of videophones to deliver a cognitive-behavioural therapy to hospice caregivers. Journal of Telemedicine and Telecare, 2011, 17, 142-145.	2.7	33
17	A multimethod analysis of shared decision-making in hospice interdisciplinary team meetings including family caregivers. Palliative Medicine, 2016, 30, 270-278.	3.1	33
18	Hospice Family Caregiver Involvement in Care Plan Meetings: A Mixed-Methods Randomized Controlled Trial. American Journal of Hospice and Palliative Medicine, 2017, 34, 849-859.	1.4	33

#	Article	IF	CITATIONS
19	Gender Differences in Caregiving at End of Life: Implications for Hospice Teams. Journal of Palliative Medicine, 2015, 18, 1048-1053.	1.1	32
20	Delivering problemâ€solving therapy to family caregivers of people with cancer: A feasibility study in outpatient palliative care. Psycho-Oncology, 2018, 27, 2494-2499.	2.3	31
21	A study of information flow in hospice interdisciplinary team meetings. Journal of Interprofessional Care, 2008, 22, 621-629.	1.7	29
22	â€~They're part of the team': participant evaluation of the ACTIVE intervention. Palliative Medicine, 2009, 23, 549-555.	3.1	29
23	Sleep Problems, Anxiety, and Global Self-Rated Health Among Hospice Family Caregivers. American Journal of Hospice and Palliative Medicine, 2018, 35, 244-249.	1.4	29
24	A Problemâ€Solving Intervention for Hospice Family Caregivers: A Randomized Clinical Trial. Journal of the American Geriatrics Society, 2019, 67, 1345-1352.	2.6	28
25	The evidence supporting educational videos for patients and caregivers receiving hospice and palliative care: A systematic review. Patient Education and Counseling, 2020, 103, 1677-1691.	2.2	27
26	Social Work Role in Hospice Pain Management: A National Survey. Journal of Social Work in End-of-Life and Palliative Care, 2009, 5, 61-74.	0.6	26
27	Lessons Learned from a Secret Facebook Support Group. Health and Social Work, 2015, 40, 125-133.	1.0	24
28	Caregiver Evaluation of the ACTIVE Intervention. American Journal of Hospice and Palliative Medicine, 2014, 31, 444-453.	1.4	23
29	Unique characteristics of informal hospice cancer caregiving. Supportive Care in Cancer, 2015, 23, 2121-2128.	2.2	23
30	Informal Hospice Caregiving: The Toll on Quality of Life. Journal of Social Work in End-of-Life and Palliative Care, 2008, 4, 312-332.	0.6	22
31	Telehospice Acceptance Among Providers: A Multidisciplinary Comparison. American Journal of Hospice and Palliative Medicine, 2009, 25, 452-457.	1.4	22
32	Challenges and Strategies for Hospice Caregivers: A Qualitative Analysis. Gerontologist, The, 2017, 57, gnw054.	3.9	22
33	Pain in Hospice Patients With Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2016, 31, 524-529.	1.9	22
34	Trends in Engagement in Advance Care Planning Behaviors and the Role of Socioeconomic Status. American Journal of Hospice and Palliative Medicine, 2016, 33, 651-657.	1.4	22
35	The Social Convoy for Family Caregivers Over the Course of Hospice. Journal of Pain and Symptom Management, 2016, 51, 213-219.	1.2	21
36	Perspectives of Health Care Providers on US South Asians' Attitudes Toward Pain Management at End of Life. American Journal of Hospice and Palliative Medicine, 2016, 33, 849-857.	1.4	21

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37	Innovative Tools to Support Family Caregivers of Persons with Cancer: The Role of Information Technology. Seminars in Oncology Nursing, 2019, 35, 384-388.	1.5	21
38	A systematic review of interventions for older adults living in jails and prisons. Aging and Mental Health, 2020, 24, 1019-1027.	2.8	21
39	Research Review: Sibling placement in foster care: a review of the evidence. Child and Family Social Work, 2007, 12, 426-433.	1.4	20
40	Anxiety among informal hospice caregivers: An exploratory study. Palliative and Supportive Care, 2015, 13, 567-573.	1.0	20
41	Family caregivers' perspectives on communication with cancer care providers. Journal of Psychosocial Oncology, 2019, 37, 777-790.	1.2	20
42	Communication Challenges and Strategies of U.S. Health Professionals Caring for Seriously III South Asian Patients and Their Families. Journal of Palliative Medicine, 2017, 20, 611-617.	1.1	19
43	Hospice Family Members' Perceptions of and Experiences With End-of-Life Care in the Nursing Home. Journal of the American Medical Directors Association, 2014, 15, 744-750.	2.5	18
44	Experiences of Sexual and Gender Minorities Caring for Adults with Non-AIDS-Related Chronic Illnesses. Social Work Research, 2015, 39, 71-81.	0.6	18
45	Stakeholder Perspectives on the Biopsychosocial and Spiritual Realities of Living With ALS: Implications for Palliative Care Teams. American Journal of Hospice and Palliative Medicine, 2019, 36, 851-857.	1.4	18
46	"lt is the â€~starting over' part that is so hard― Using an online group to support hospice bereavement. Palliative and Supportive Care, 2015, 13, 351-357.	1.0	17
47	Social Workers as Behavioral Health Consultants in the Primary Care Clinic. Health and Social Work, 2016, 41, 196-200.	1.0	16
48	Digital Storytelling: Families' Search for Meaning after Child Death. Journal of Social Work in End-of-Life and Palliative Care, 2017, 13, 239-250.	0.6	15
49	Accounts of Family Conflict in Home Hospice Care: The Central Role of Autonomy for Informal Caregiver Resilience. Journal of Family Nursing, 2019, 25, 190-218.	1.9	15
50	Challenges in Implementing Hospice Clinical Trials: Preserving Scientific Integrity While Facing Change. Journal of Pain and Symptom Management, 2020, 59, 365-371.	1.2	15
51	Qualitative evaluation of a problem-solving intervention for informal hospice caregivers. Palliative Medicine, 2012, 26, 1018-1024.	3.1	14
52	Family Members' Experience With Hospice in Nursing Homes. American Journal of Hospice and Palliative Medicine, 2016, 33, 354-362.	1.4	14
53	Relationships among Symptom Management Burden, Coping Responses, and Caregiver Psychological Distress at End of Life. Journal of Palliative Medicine, 2018, 21, 1234-1241.	1.1	14
54	Digital Storytelling as an Intervention for Bereaved Family Members. Omega: Journal of Death and Dying, 2021, 82, 570-586.	1.0	14

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55	Recruitment Challenges and Strategies in a Home-Based Telehealth Study. Telemedicine Journal and E-Health, 2010, 16, 839-843.	2.8	13
56	Ethical Dilemmas Faced by Hospice Social Workers. Social Work in Health Care, 2014, 53, 950-968.	1.6	13
57	The Stress of Sadness. American Journal of Hospice and Palliative Medicine, 2015, 32, 745-749.	1.4	13
58	Communicating Caregivers' Challenges With Cancer Pain Management: An Analysis of Home Hospice Visits. Journal of Pain and Symptom Management, 2018, 55, 1296-1303.	1.2	13
59	Adapting the Resilience Framework for Family Caregivers of Hospice Patients With Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2019, 34, 399-411.	1.9	13
60	Job Satisfaction, Intent To Stay, and Recommended Job Improvements: The Palliative Nursing Assistant Speaks. Journal of Palliative Medicine, 2013, 16, 1356-1361.	1.1	12
61	The Promise of Secret Facebook Groups for Active Family Caregivers of Hospice Patients. Journal of Palliative Medicine, 2014, 17, 1199-1200.	1.1	12
62	Question Asking by Family Caregivers in Hospice Interdisciplinary Team Meetings. Research in Gerontological Nursing, 2010, 3, 82-88.	0.6	12
63	Team functioning in hospice interprofessional meetings: An exploratory study of providers' perspectives. Journal of Interprofessional Care, 2017, 31, 455-462.	1.7	11
64	Factors influencing engagement in an online support group for family caregivers of individuals with advanced cancer. Journal of Psychosocial Oncology, 2020, 38, 235-250.	1.2	10
65	"Just Let Me Go― End-of-Life Planning Among Ojibwe Elders. Gerontologist, The, 2018, 58, gnw151.	3.9	9
66	Shared Decision Making in Home Hospice Nursing Visits: A Qualitative Study. Journal of Pain and Symptom Management, 2018, 55, 922-929.	1.2	9
67	Health-Care Providers' Perspectives on Decision-Making Among Seriously Ill Patients of South Asian Origin in the United States. Journal of Palliative Care, 2019, 34, 181-188.	1.0	9
68	Design and Preliminary Testing of the Caregiver-Centered Communication Questionnaire (CCCQ). Journal of Palliative Care, 2020, 35, 154-160.	1.0	9
69	Hospice and palliative social workers' experiences with clients at risk of suicide. Palliative and Supportive Care, 2016, 14, 664-671.	1.0	9
70	Home Internet Use among Hospice Service Recipients: Recommendations for Web-Based Interventions. Journal of Medical Systems, 2007, 31, 385-389.	3.6	8
71	Problem Solving Interventions: An Opportunity for Hospice Social Workers to Better Meet Caregiver Needs. Journal of Social Work in End-of-Life and Palliative Care, 2012, 8, 3-9.	0.6	8
72	Quality Hospice Care in Adult Family Homes: Barriers and Facilitators. Journal of the American Medical Directors Association, 2018, 19, 136-140.	2.5	8

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73	A Comparison of Rural and Urban Hospice Family Caregivers' Cancer Pain Knowledge and Experience. Journal of Pain and Symptom Management, 2019, 58, 685-689.	1.2	8
74	Job Satisfaction Among Child Welfare Workers in Public and Performance-Based Contracting Environments. Journal of Public Child Welfare, 2009, 3, 159-172.	0.9	7
75	Exploring the Challenges that Family Caregivers Faced When Caring for Hospice Patients with Heart Failure. Journal of Social Work in End-of-Life and Palliative Care, 2018, 14, 162-176.	0.6	7
76	Comfort Needs of Cancer Family Caregivers in Outpatient Palliative Care. Journal of Hospice and Palliative Nursing, 2021, 23, 221-228.	0.9	7
77	Spoken words as biomarkers: using machine learning to gain insight into communication as a predictor of anxiety. Journal of the American Medical Informatics Association: JAMIA, 2020, 27, 929-933.	4.4	7
78	ENVISION: A Tool to Improve Communication in Hospice Interdisciplinary Team Meetings. Journal of Gerontological Nursing, 2020, 46, 9-14.	0.6	7
79	Treating Perpetrators of Child Physical Abuse. Trauma, Violence, and Abuse, 2009, 10, 115-124.	6.2	6
80	Randomized clinical trials in US hospices: challenges and the current state of the art. Clinical Investigation, 2015, 5, 839-846.	0.0	6
81	Caregiver Speaks Study Protocol: A Technologically-Mediated Storytelling Intervention for Hospice Family Caregivers of Persons Living With Dementia. American Journal of Hospice and Palliative Medicine, 2021, 38, 376-382.	1.4	6
82	Perceived Benefits and Burdens of Participation for Caregivers of Cancer Patients in Hospice Clinical Trials: A Pilot Study. Journal of Pain and Symptom Management, 2021, 61, 1147-1154.	1.2	6
83	Patient and Provider Satisfaction With Pediatric Urology Telemedicine Clinic. Journal of Patient Experience, 2021, 8, 237437352097573.	0.9	6
84	Perceptions of Parenting Residents Among Family Medicine Residency Directors. Family Medicine, 2018, 50, 756-762.	0.5	6
85	Use of the Time, Interaction, and Performance Theory to Study Hospice Interdisciplinary Team Meetings. Journal of Hospice and Palliative Nursing, 2008, 10, 376-381.	0.9	5
86	Family Perspectives on the Hospice Experience in Adult Family Homes. Journal of Gerontological Social Work, 2011, 54, 159-174.	1.0	5
87	Application of the VALUE Communication Principles in ACTIVE Hospice Team Meetings. Journal of Palliative Medicine, 2013, 16, 60-66.	1.1	5
88	Differential effects of patient symptom subtypes on informal hospice caregiver depression. Palliative and Supportive Care, 2018, 16, 127-136.	1.0	5
89	Ways of Grieving Among Ojibwe Elders. Omega: Journal of Death and Dying, 2018, 78, 107-119.	1.0	5
90	Assessing the Reliability and Validity of a Brief Measure of Caregiver Quality of Life. Journal of Pain and Symptom Management, 2019, 58, 871-877.	1.2	5

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91	U.S. Providers' Perceptions of the Psychosocial Needs of Seriously Ill Patients of South Asian Origin: Implications for Health Social Work. Health and Social Work, 2019, 44, 177-184.	1.0	5
92	Digital storytelling: Narrating meaning in bereavement. Death Studies, 2021, , 1-9.	2.7	5
93	Family Caregiver Problems in Outpatient Palliative Oncology. Journal of Palliative Medicine, 2021, 24, 1056-1060.	1.1	5
94	Insomnia Symptoms Among Hospice Family Caregivers: Prevalence and Association with Caregiver Mental and Physical Health, Quality of Life, and Caregiver Burden. American Journal of Hospice and Palliative Medicine, 2023, 40, 517-528.	1.4	5
95	Delivering a psycho-educational intervention to Hispanic hospice informal caregivers. Palliative Medicine, 2011, 25, 185-186.	3.1	4
96	Commitment to care: A grounded theory of informal hospice caregivers' development as symptom managers. Qualitative Social Work, 2013, 12, 358-371.	1.4	4
97	A Pilot Study of An Intervention to Increase Family Member Involvement in Nursing Home Care Plan Meetings. Journal of Applied Gerontology, 2021, 40, 073346482094692.	2.0	4
98	Anxiety, Depression, Quality of Life, Caregiver Burden, and Perceptions of Caregiver-Centered Communication among Black and White Hospice Family Caregivers. Journal of Palliative Medicine, 2021,	1.1	4
99	Facebook Online Support Groups for Hospice Family Caregivers of Advanced Cancer Patients: Protocol, Facilitation Skills and Promising Outcomes. Journal of Social Work in End-of-Life and Palliative Care, 2022, 18, 146-159.	0.6	4
100	Rethinking Family Caregiving: Tailoring Cognitive-Behavioral Therapies to the Hospice Experience. Health and Social Work, 2014, 39, 244-250.	1.0	3
101	Testing the Factorial Validity of Scores From the Caregiver Pain Medicine Questionnaire. Journal of Pain and Symptom Management, 2014, 48, 99-109.	1.2	3
102	A Qualitative Analysis of Information Sharing in Hospice Interdisciplinary Group Meetings. American Journal of Hospice and Palliative Medicine, 2017, 34, 901-906.	1.4	3
103	Behind the doors of home hospice patients: A secondary qualitative analysis of hospice nurse communication with patients and families. Palliative and Supportive Care, 2019, 17, 579-583.	1.0	3
104	Preliminary Results of Caregiver Speaks: A Storytelling Intervention for Bereaved Family Caregivers. Journal of Loss and Trauma, 2020, 25, 438-453.	1.5	3
105	The Effect of Digital Literacy on Participation in Social Media Clinical Trials in Cancer: Tailoring an Informed Consent Process. Telemedicine Journal and E-Health, 2022, 28, 1682-1689.	2.8	3
106	Learning to Facilitate Advance Care Planning: The Novice Social Worker's Experience. Journal of Teaching in Social Work, 2014, 34, 514-530.	0.6	2
107	Coping Responses Among Hospice Family Caregivers: A Confirmatory Factor Analysis. Journal of Pain and Symptom Management, 2015, 50, 867-873.	1.2	2
108	Engaging Specialist Palliative Care in the Management of Amyotrophic Lateral Sclerosis: A Patient-, Family-, and Provider-Based Approach. Journal of Palliative Care, 2020, , 082585971989582.	1.0	2

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109	Toward a Practice-Informed Agenda for Hospice Intervention Research: What Are Staff Members' Biggest Challenges?. American Journal of Hospice and Palliative Medicine, 2021, 38, 467-471.	1.4	2
110	Not Just Sitting By: Family Caregivers' Perspectives on Triadic Communication. Innovation in Aging, 2020, 4, 571-571.	0.1	2
111	Examining spoken words and acoustic features of therapy sessions to understand family caregivers' anxiety and quality of life. International Journal of Medical Informatics, 2022, 160, 104716.	3.3	2
112	Opportunities and Challenges for Visual Qualitative Intervention Research on Facebook. International Journal of Qualitative Methods, The, 2022, 21, 160940692210744.	2.8	2
113	The Lived Experience of Physical Separation for Hospice Patients and Families amid COVID-19. Journal of Pain and Symptom Management, 2022, 63, 971-979.	1.2	2
114	Physical Health Problems: Shaping Transitions of Care. Annual Review of Gerontology and Geriatrics, 2011, 31, 65-91.	0.5	1
115	"l Have to Age in This Bodyâ€: Lesbian and Bisexual Older Women's Perspectives on a Health Behavior Intervention. Women's Health Issues, 2016, 26, S63-S70.	2.0	1
116	The Paradox of Hospice for Caregivers of Cancer Patients. Journal of Pain and Symptom Management, 2018, 56, e8-e11.	1.2	1
117	Development and Preliminary Evaluation of a Pain Management Manual for Hospice Providers to Support and Educate Family Caregivers. American Journal of Hospice and Palliative Medicine, 2019, 36, 207-215.	1.4	1
118	Lessons learned from a clinical-research partnership in outpatient palliative care Journal of Clinical Oncology, 2016, 34, 238-238.	1.6	1
119	Hospice and Caregiving. , 2019, , 1-3.		1
120	Mixed methods analysis of hospice staff perceptions and shared decision making practices in hospice. Supportive Care in Cancer, 2021, , 1.	2.2	1
121	A Goal-Directed Model of Collaborative Decision Making in Hospice and Palliative Care. Journal of Palliative Care, 2022, 37, 120-124.	1.0	1
122	Perspectives of State Association Leaders During the COVID-19 Pandemic. American Journal of Hospice and Palliative Medicine, 2022, , 104990912210902.	1.4	1
123	Family Caregivers' Challenges in Cancer Pain Management for Patients Receiving Palliative Care. American Journal of Hospice and Palliative Medicine, 2023, 40, 43-51.	1.4	1
124	Designing an Intervention., 2019,, 41-59.		0
125	EXPLORING SEXUAL HEALTH AND HIV COMMUNICATION BETWEEN HEALTHCARE PROVIDERS AND OLDER PATIENTS. Innovation in Aging, 2019, 3, S548-S549.	0.1	0
126	Hospice and Caregiving. , 2021, , 2451-2453.		О

#	Article	lF	CITATIONS
127	A mixed-method evaluation of the adoption and implementation of the College Alcohol Intervention Matrix among prevention experts: a study protocol. Implementation Science Communications, 2021, 2, 142.	2.2	O