

Karla T Washington

List of Publications by Year in descending order

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Version: 2024-02-01

127
papers

1,992
citations

257450

24
h-index

361022

35
g-index

127
all docs

127
docs citations

127
times ranked

1915
citing authors

#	ARTICLE	IF	CITATIONS
1	Family Caregiversâ€™ Challenges in Cancer Pain Management for Patients Receiving Palliative Care. American Journal of Hospice and Palliative Medicine, 2023, 40, 43-51.	1.4	1
2	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
3	A Goal-Directed Model of Collaborative Decision Making in Hospice and Palliative Care. Journal of Palliative Care, 2022, 37, 120-124.	1.0	1
4	Examining spoken words and acoustic features of therapy sessions to understand family caregiversâ€™ anxiety and quality of life. International Journal of Medical Informatics, 2022, 160, 104716.	3.3	2
5	Opportunities and Challenges for Visual Qualitative Intervention Research on Facebook. International Journal of Qualitative Methods, The, 2022, 21, 160940692210744.	2.8	2
6	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
7	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
8	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
9	Perspectives of State Association Leaders During the COVID-19 Pandemic. American Journal of Hospice and Palliative Medicine, 2022, , 104990912210902.	1.4	1
10	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
11	Digital Storytelling as an Intervention for Bereaved Family Members. Omega: Journal of Death and Dying, 2021, 82, 570-586.	1.0	14
12	Toward a Practice-Informed Agenda for Hospice Intervention Research: What Are Staff Membersâ€™ Biggest Challenges?. American Journal of Hospice and Palliative Medicine, 2021, 38, 467-471.	1.4	2
13	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
14	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
15	Patient and Provider Satisfaction With Pediatric Urology Telemedicine Clinic. Journal of Patient Experience, 2021, 8, 237437352097573.	0.9	6
16	Comfort Needs of Cancer Family Caregivers in Outpatient Palliative Care. Journal of Hospice and Palliative Nursing, 2021, 23, 221-228.	0.9	7
17	Digital storytelling: Narrating meaning in bereavement. Death Studies, 2021, , 1-9.	2.7	5
18	Family Caregiver Problems in Outpatient Palliative Oncology. Journal of Palliative Medicine, 2021, 24, 1056-1060.	1.1	5

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19	Mixed methods analysis of hospice staff perceptions and shared decision making practices in hospice. <i>Supportive Care in Cancer</i> , 2021, , 1.	2.2	1
20	Anxiety, Depression, Quality of Life, Caregiver Burden, and Perceptions of Caregiver-Centered Communication among Black and White Hospice Family Caregivers. <i>Journal of Palliative Medicine</i> , 2021, , .	1.1	4
21	Hospice and Caregiving. , 2021, , 2451-2453.		0
22	A mixed-method evaluation of the adoption and implementation of the College Alcohol Intervention Matrix among prevention experts: a study protocol. <i>Implementation Science Communications</i> , 2021, 2, 142.	2.2	0
23	A systematic review of interventions for older adults living in jails and prisons. <i>Aging and Mental Health</i> , 2020, 24, 1019-1027.	2.8	21
24	Challenges in Implementing Hospice Clinical Trials: Preserving Scientific Integrity While Facing Change. <i>Journal of Pain and Symptom Management</i> , 2020, 59, 365-371.	1.2	15
25	Factors influencing engagement in an online support group for family caregivers of individuals with advanced cancer. <i>Journal of Psychosocial Oncology</i> , 2020, 38, 235-250.	1.2	10
26	Design and Preliminary Testing of the Caregiver-Centered Communication Questionnaire (CCCQ). <i>Journal of Palliative Care</i> , 2020, 35, 154-160.	1.0	9
27	Online social support groups for informal caregivers of hospice patients with cancer. <i>European Journal of Oncology Nursing</i> , 2020, 44, 101698.	2.1	40
28	Engaging Specialist Palliative Care in the Management of Amyotrophic Lateral Sclerosis: A Patient-, Family-, and Provider-Based Approach. <i>Journal of Palliative Care</i> , 2020, , 082585971989582.	1.0	2
29	Preliminary Results of Caregiver Speaks: A Storytelling Intervention for Bereaved Family Caregivers. <i>Journal of Loss and Trauma</i> , 2020, 25, 438-453.	1.5	3
30	The evidence supporting educational videos for patients and caregivers receiving hospice and palliative care: A systematic review. <i>Patient Education and Counseling</i> , 2020, 103, 1677-1691.	2.2	27
31	Spoken words as biomarkers: using machine learning to gain insight into communication as a predictor of anxiety. <i>Journal of the American Medical Informatics Association: JAMIA</i> , 2020, 27, 929-933.	4.4	7
32	Not Just Sitting By: Family Caregiversâ€™ Perspectives on Triadic Communication. <i>Innovation in Aging</i> , 2020, 4, 571-571.	0.1	2
33	ENVISION: A Tool to Improve Communication in Hospice Interdisciplinary Team Meetings. <i>Journal of Gerontological Nursing</i> , 2020, 46, 9-14.	0.6	7
34	Assessing the Reliability and Validity of a Brief Measure of Caregiver Quality of Life. <i>Journal of Pain and Symptom Management</i> , 2019, 58, 871-877.	1.2	5
35	Designing an Intervention. , 2019, , 41-59.		0
36	A Comparison of Rural and Urban Hospice Family Caregivers' Cancer Pain Knowledge and Experience. <i>Journal of Pain and Symptom Management</i> , 2019, 58, 685-689.	1.2	8

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37	Adapting the Resilience Framework for Family Caregivers of Hospice Patients With Dementia. <i>American Journal of Alzheimer's Disease and Other Dementias</i> , 2019, 34, 399-411.	1.9	13
38	U.S. Providers'™ Perceptions of the Psychosocial Needs of Seriously Ill Patients of South Asian Origin: Implications for Health Social Work. <i>Health and Social Work</i> , 2019, 44, 177-184.	1.0	5
39	Family caregivers'™ perspectives on communication with cancer care providers. <i>Journal of Psychosocial Oncology</i> , 2019, 37, 777-790.	1.2	20
40	Innovative Tools to Support Family Caregivers of Persons with Cancer: The Role of Information Technology. <i>Seminars in Oncology Nursing</i> , 2019, 35, 384-388.	1.5	21
41	A Problem-Solving Intervention for Hospice Family Caregivers: A Randomized Clinical Trial. <i>Journal of the American Geriatrics Society</i> , 2019, 67, 1345-1352.	2.6	28
42	Stakeholder Perspectives on the Biopsychosocial and Spiritual Realities of Living With ALS: Implications for Palliative Care Teams. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 851-857.	1.4	18
43	Health-Care Providers'™ Perspectives on Decision-Making Among Seriously Ill Patients of South Asian Origin in the United States. <i>Journal of Palliative Care</i> , 2019, 34, 181-188.	1.0	9
44	Behind the doors of home hospice patients: A secondary qualitative analysis of hospice nurse communication with patients and families. <i>Palliative and Supportive Care</i> , 2019, 17, 579-583.	1.0	3
45	Accounts of Family Conflict in Home Hospice Care: The Central Role of Autonomy for Informal Caregiver Resilience. <i>Journal of Family Nursing</i> , 2019, 25, 190-218.	1.9	15
46	EXPLORING SEXUAL HEALTH AND HIV COMMUNICATION BETWEEN HEALTHCARE PROVIDERS AND OLDER PATIENTS. <i>Innovation in Aging</i> , 2019, 3, S548-S549.	0.1	0
47	Development and Preliminary Evaluation of a Pain Management Manual for Hospice Providers to Support and Educate Family Caregivers. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 207-215.	1.4	1
48	Hospice and Caregiving. , 2019, , 1-3.		1
49	“Just Let Me Go” End-of-Life Planning Among Ojibwe Elders. <i>Gerontologist</i> , The, 2018, 58, gnw151.	3.9	9
50	Communicating Caregivers' Challenges With Cancer Pain Management: An Analysis of Home Hospice Visits. <i>Journal of Pain and Symptom Management</i> , 2018, 55, 1296-1303.	1.2	13
51	Sleep Problems, Anxiety, and Global Self-Rated Health Among Hospice Family Caregivers. <i>American Journal of Hospice and Palliative Medicine</i> , 2018, 35, 244-249.	1.4	29
52	Differential effects of patient symptom subtypes on informal hospice caregiver depression. <i>Palliative and Supportive Care</i> , 2018, 16, 127-136.	1.0	5
53	Ways of Grieving Among Ojibwe Elders. <i>Omega: Journal of Death and Dying</i> , 2018, 78, 107-119.	1.0	5
54	Quality Hospice Care in Adult Family Homes: Barriers and Facilitators. <i>Journal of the American Medical Directors Association</i> , 2018, 19, 136-140.	2.5	8

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55	Pain Management Concerns From the Hospice Family Caregiversâ€™ Perspective. <i>American Journal of Hospice and Palliative Medicine</i> , 2018, 35, 601-611.	1.4	39
56	Shared Decision Making in Home Hospice Nursing Visits: A Qualitative Study. <i>Journal of Pain and Symptom Management</i> , 2018, 55, 922-929.	1.2	9
57	Relationships among Symptom Management Burden, Coping Responses, and Caregiver Psychological Distress at End of Life. <i>Journal of Palliative Medicine</i> , 2018, 21, 1234-1241.	1.1	14
58	The Paradox of Hospice for Caregivers of Cancer Patients. <i>Journal of Pain and Symptom Management</i> , 2018, 56, e8-e11.	1.2	1
59	Delivering problemâ€solving therapy to family caregivers of people with cancer: A feasibility study in outpatient palliative care. <i>Psycho-Oncology</i> , 2018, 27, 2494-2499.	2.3	31
60	Exploring the Challenges that Family Caregivers Faced When Caring for Hospice Patients with Heart Failure. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2018, 14, 162-176.	0.6	7
61	Perceptions of Parenting Residents Among Family Medicine Residency Directors. <i>Family Medicine</i> , 2018, 50, 756-762.	0.5	6
62	Challenges and Strategies for Hospice Caregivers: A Qualitative Analysis. <i>Gerontologist</i> , The, 2017, 57, gnw054.	3.9	22
63	A Qualitative Analysis of Information Sharing in Hospice Interdisciplinary Group Meetings. <i>American Journal of Hospice and Palliative Medicine</i> , 2017, 34, 901-906.	1.4	3
64	Communication Challenges and Strategies of U.S. Health Professionals Caring for Seriously Ill South Asian Patients and Their Families. <i>Journal of Palliative Medicine</i> , 2017, 20, 611-617.	1.1	19
65	Hospice Family Caregiver Involvement in Care Plan Meetings: A Mixed-Methods Randomized Controlled Trial. <i>American Journal of Hospice and Palliative Medicine</i> , 2017, 34, 849-859.	1.4	33
66	The Prevalence and Risks for Depression and Anxiety in Hospice Caregivers. <i>Journal of Palliative Medicine</i> , 2017, 20, 366-371.	1.1	36
67	Team functioning in hospice interprofessional meetings: An exploratory study of providersâ€™ perspectives. <i>Journal of Interprofessional Care</i> , 2017, 31, 455-462.	1.7	11
68	The Effect of Internet Group Support for Caregivers on Social Support, Self-Efficacy, and Caregiver Burden: A Meta-Analysis. <i>Telemedicine Journal and E-Health</i> , 2017, 23, 621-629.	2.8	44
69	Digital Storytelling: Familiesâ€™ Search for Meaning after Child Death. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2017, 13, 239-250.	0.6	15
70	The Social Convoy for Family Caregivers Over the Course of Hospice. <i>Journal of Pain and Symptom Management</i> , 2016, 51, 213-219.	1.2	21
71	â€œ Have to Age in This Bodyâ€: Lesbian and Bisexual Older Women's Perspectives on a Health Behavior Intervention. <i>Women's Health Issues</i> , 2016, 26, S63-S70.	2.0	1
72	Pain in Hospice Patients With Dementia. <i>American Journal of Alzheimer's Disease and Other Dementias</i> , 2016, 31, 524-529.	1.9	22

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73	Social Workers as Behavioral Health Consultants in the Primary Care Clinic. <i>Health and Social Work</i> , 2016, 41, 196-200.	1.0	16
74	A multimethod analysis of shared decision-making in hospice interdisciplinary team meetings including family caregivers. <i>Palliative Medicine</i> , 2016, 30, 270-278.	3.1	33
75	Family Members' Experience With Hospice in Nursing Homes. <i>American Journal of Hospice and Palliative Medicine</i> , 2016, 33, 354-362.	1.4	14
76	Perspectives of Health Care Providers on US South Asians' Attitudes Toward Pain Management at End of Life. <i>American Journal of Hospice and Palliative Medicine</i> , 2016, 33, 849-857.	1.4	21
77	Trends in Engagement in Advance Care Planning Behaviors and the Role of Socioeconomic Status. <i>American Journal of Hospice and Palliative Medicine</i> , 2016, 33, 651-657.	1.4	22
78	Hospice and palliative social workers' experiences with clients at risk of suicide. <i>Palliative and Supportive Care</i> , 2016, 14, 664-671.	1.0	9
79	Lessons learned from a clinical-research partnership in outpatient palliative care.. <i>Journal of Clinical Oncology</i> , 2016, 34, 238-238.	1.6	1
80	Anxiety among informal hospice caregivers: An exploratory study. <i>Palliative and Supportive Care</i> , 2015, 13, 567-573.	1.0	20
81	Gender Differences in Caregiving at End of Life: Implications for Hospice Teams. <i>Journal of Palliative Medicine</i> , 2015, 18, 1048-1053.	1.1	32
82	Randomized clinical trials in US hospices: challenges and the current state of the art. <i>Clinical Investigation</i> , 2015, 5, 839-846.	0.0	6
83	"It is the "starting over" part that is so hard" Using an online group to support hospice bereavement. <i>Palliative and Supportive Care</i> , 2015, 13, 351-357.	1.0	17
84	The Stress of Sadness. <i>American Journal of Hospice and Palliative Medicine</i> , 2015, 32, 745-749.	1.4	13
85	Coping Responses Among Hospice Family Caregivers: A Confirmatory Factor Analysis. <i>Journal of Pain and Symptom Management</i> , 2015, 50, 867-873.	1.2	2
86	Unique characteristics of informal hospice cancer caregiving. <i>Supportive Care in Cancer</i> , 2015, 23, 2121-2128.	2.2	23
87	Experiences of Sexual and Gender Minorities Caring for Adults with Non-AIDS-Related Chronic Illnesses. <i>Social Work Research</i> , 2015, 39, 71-81.	0.6	18
88	Lessons Learned from a Secret Facebook Support Group. <i>Health and Social Work</i> , 2015, 40, 125-133.	1.0	24
89	Learning to Facilitate Advance Care Planning: The Novice Social Worker's Experience. <i>Journal of Teaching in Social Work</i> , 2014, 34, 514-530.	0.6	2
90	Caregiver Evaluation of the ACTIVE Intervention. <i>American Journal of Hospice and Palliative Medicine</i> , 2014, 31, 444-453.	1.4	23

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91	Ethical Dilemmas Faced by Hospice Social Workers. <i>Social Work in Health Care</i> , 2014, 53, 950-968.	1.6	13
92	Rethinking Family Caregiving: Tailoring Cognitive-Behavioral Therapies to the Hospice Experience. <i>Health and Social Work</i> , 2014, 39, 244-250.	1.0	3
93	The Promise of Secret Facebook Groups for Active Family Caregivers of Hospice Patients. <i>Journal of Palliative Medicine</i> , 2014, 17, 1199-1200.	1.1	12
94	Testing the Factorial Validity of Scores From the Caregiver Pain Medicine Questionnaire. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 99-109.	1.2	3
95	Understanding Social Support Burden Among Family Caregivers. <i>Health Communication</i> , 2014, 29, 901-910.	3.1	41
96	Hospice Family Members' Perceptions of and Experiences With End-of-Life Care in the Nursing Home. <i>Journal of the American Medical Directors Association</i> , 2014, 15, 744-750.	2.5	18
97	Hospice Caregivers' Experiences With Pain Management: "I'm Not a Doctor, and I Don't Know if I Helped Her Go Faster or Slower" <i>Journal of Pain and Symptom Management</i> , 2013, 46, 846-858.	1.2	40
98	Hospice Caregiver Depression: The Evidence Surrounding the Greatest Pain of All. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2013, 9, 256-271.	0.6	36
99	Job Satisfaction, Intent To Stay, and Recommended Job Improvements: The Palliative Nursing Assistant Speaks. <i>Journal of Palliative Medicine</i> , 2013, 16, 1356-1361.	1.1	12
100	Application of the VALUE Communication Principles in ACTIVE Hospice Team Meetings. <i>Journal of Palliative Medicine</i> , 2013, 16, 60-66.	1.1	5
101	Commitment to care: A grounded theory of informal hospice caregivers' development as symptom managers. <i>Qualitative Social Work</i> , 2013, 12, 358-371.	1.4	4
102	Stress Variances Among Informal Hospice Caregivers. <i>Qualitative Health Research</i> , 2012, 22, 1114-1125.	2.1	38
103	Qualitative evaluation of a problem-solving intervention for informal hospice caregivers. <i>Palliative Medicine</i> , 2012, 26, 1018-1024.	3.1	14
104	A Noninferiority Trial of a Problem-Solving Intervention for Hospice Caregivers: In Person versus Videophone. <i>Journal of Palliative Medicine</i> , 2012, 15, 653-660.	1.1	52
105	A Systematic Review of the Evidence Base for Telehospice. <i>Telemedicine Journal and E-Health</i> , 2012, 18, 38-47.	2.8	41
106	Problem Solving Interventions: An Opportunity for Hospice Social Workers to Better Meet Caregiver Needs. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2012, 8, 3-9.	0.6	8
107	Family Perspectives on the Hospice Experience in Adult Family Homes. <i>Journal of Gerontological Social Work</i> , 2011, 54, 159-174.	1.0	5
108	Information needs of informal caregivers of older adults with chronic health conditions. <i>Patient Education and Counseling</i> , 2011, 83, 37-44.	2.2	126

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109	Delivering a psycho-educational intervention to Hispanic hospice informal caregivers. <i>Palliative Medicine</i> , 2011, 25, 185-186.	3.1	4
110	Physical Health Problems: Shaping Transitions of Care. <i>Annual Review of Gerontology and Geriatrics</i> , 2011, 31, 65-91.	0.5	1
111	Use of videophones to deliver a cognitive-behavioural therapy to hospice caregivers. <i>Journal of Telemedicine and Telecare</i> , 2011, 17, 142-145.	2.7	33
112	The Role of Human Factors in Telehealth. <i>Telemedicine Journal and E-Health</i> , 2010, 16, 446-453.	2.8	48
113	Recruitment Challenges and Strategies in a Home-Based Telehealth Study. <i>Telemedicine Journal and E-Health</i> , 2010, 16, 839-843.	2.8	13
114	A Problem Solving Intervention for Hospice Caregivers: A Pilot Study. <i>Journal of Palliative Medicine</i> , 2010, 13, 1005-1011.	1.1	47
115	Question Asking by Family Caregivers in Hospice Interdisciplinary Team Meetings. <i>Research in Gerontological Nursing</i> , 2010, 3, 82-88.	0.6	12
116	Job Satisfaction Among Child Welfare Workers in Public and Performance-Based Contracting Environments. <i>Journal of Public Child Welfare</i> , 2009, 3, 159-172.	0.9	7
117	Telehospice Acceptance Among Providers: A Multidisciplinary Comparison. <i>American Journal of Hospice and Palliative Medicine</i> , 2009, 25, 452-457.	1.4	22
118	Treating Perpetrators of Child Physical Abuse. <i>Trauma, Violence, and Abuse</i> , 2009, 10, 115-124.	6.2	6
119	“They’re part of the team”: participant evaluation of the ACTIVE intervention. <i>Palliative Medicine</i> , 2009, 23, 549-555.	3.1	29
120	Social Work Role in Hospice Pain Management: A National Survey. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2009, 5, 61-74.	0.6	26
121	Barriers to Pain Management: Caregiver Perceptions and Pain Talk by Hospice Interdisciplinary Teams. <i>Journal of Pain and Symptom Management</i> , 2008, 36, 374-382.	1.2	57
122	A study of information flow in hospice interdisciplinary team meetings. <i>Journal of Interprofessional Care</i> , 2008, 22, 621-629.	1.7	29
123	Informal Hospice Caregiving: The Toll on Quality of Life. <i>Journal of Social Work in End-of-Life and Palliative Care</i> , 2008, 4, 312-332.	0.6	22
124	Barriers to Hospice Use among African Americans: A Systematic Review. <i>Health and Social Work</i> , 2008, 33, 267-274.	1.0	62
125	Use of the Time, Interaction, and Performance Theory to Study Hospice Interdisciplinary Team Meetings. <i>Journal of Hospice and Palliative Nursing</i> , 2008, 10, 376-381.	0.9	5
126	Research Review: Sibling placement in foster care: a review of the evidence. <i>Child and Family Social Work</i> , 2007, 12, 426-433.	1.4	20

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127	Home Internet Use among Hospice Service Recipients: Recommendations for Web-Based Interventions. Journal of Medical Systems, 2007, 31, 385-389.	3.6	8