

Kelli I Stajduhar

List of Publications by Year in descending order

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Version: 2024-02-01

124
papers

3,954
citations

117571

34
h-index

143943

57
g-index

125
all docs

125
docs citations

125
times ranked

3459
citing authors

#	ARTICLE	IF	CITATIONS
1	Prolonged Grief and Bereavement Supports Within a Caregiver Population Who Transition Through a Palliative Care Program in British Columbia, Canada. <i>American Journal of Hospice and Palliative Medicine</i> , 2022, 39, 361-369.	0.8	3
2	Feasibility of Routine Quality of Life Measurement for People Living With Dementia in Long-Term Care. <i>Journal of the American Medical Directors Association</i> , 2022, 23, 1221-1226.	1.2	6
3	Knowledge translation resources to support the use of quality of life assessment tools for the care of older adults living at home and their family caregivers. <i>Quality of Life Research</i> , 2022, 31, 1727-1747.	1.5	2
4	A realist review of the home care literature and its blind spots. <i>Journal of Evaluation in Clinical Practice</i> , 2022, 28, 680-689.	0.9	9
5	Equity-Oriented Healthcare: What It Is and Why We Need It in Oncology. <i>Current Oncology</i> , 2022, 29, 186-192.	0.9	10
6	A codevelopment process to advance methods for the use of patient-reported outcome measures and patient-reported experience measures with people who are homeless and experience chronic illness. <i>Health Expectations</i> , 2022, 25, 2264-2274.	1.1	3
7	Self-Perceived Competence of Nurses and Care Aides Providing a Palliative Approach in Home, Hospital, and Residential Care Settings: A Cross-Sectional Survey. <i>Canadian Journal of Nursing Research</i> , 2021, 53, 64-77.	0.6	9
8	"We are to be like machines" fill the bed before it gets cold: Exploring the emotional geographies of healthcare providers caring for dying residents in long-term care facilities. <i>Social Science and Medicine</i> , 2021, 272, 113749.	1.8	6
9	"Bare-bones" to "silver linings": lessons on integrating a palliative approach to care in long-term care in Western Canada. <i>BMC Health Services Research</i> , 2021, 21, 610.	0.9	7
10	Applying the concept of structural empowerment to interactions between families and home care nurses. <i>Nursing Inquiry</i> , 2020, 27, e12313.	1.1	3
11	Provocations on privilege in palliative care: Are we meeting our core mandate?. <i>Progress in Palliative Care</i> , 2020, 28, 89-93.	0.7	23
12	"Everybody in this community is at risk of dying": An ethnographic exploration on the potential of integrating a palliative approach to care among workers in inner-city settings. <i>Palliative and Supportive Care</i> , 2020, 18, 670-675.	0.6	10
13	Divergent Perspectives on the Use of the Edmonton Symptom Assessment System (Revised) in Palliative Care. <i>Journal of Hospice and Palliative Nursing</i> , 2020, 22, 75-81.	0.5	7
14	Caregiving at the margins: An ethnographic exploration of family caregivers experiences providing care for structurally vulnerable populations at the end-of-life. <i>Palliative Medicine</i> , 2020, 34, 946-953.	1.3	20
15	Team-Based Integrated Knowledge Translation for Enhancing Quality of Life in Long-term Care Settings: A Multi-method, Multi-sectoral Research Design. <i>International Journal of Health Policy and Management</i> , 2020, 9, 138-142.	0.5	8
16	"Just too busy living in the moment and surviving": barriers to accessing health care for structurally vulnerable populations at end-of-life. <i>BMC Palliative Care</i> , 2019, 18, 11.	0.8	65
17	A scoping review of palliative care for persons with severe persistent mental illness. <i>Palliative and Supportive Care</i> , 2019, 17, 479-487.	0.6	37
18	Micro-Meso-Macro Practice Tensions in Using Patient-Reported Outcome and Experience Measures in Hospital Palliative Care. <i>Qualitative Health Research</i> , 2019, 29, 510-521.	1.0	38

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19	Healthcare providers'™ perspectives on perceived barriers and facilitators of compassion: Results from a grounded theory study. <i>Journal of Clinical Nursing</i> , 2018, 27, 2083-2097.	1.4	37
20	What are healthcare providers'™ understandings and experiences of compassion? The healthcare compassion model: a grounded theory study of healthcare providers in Canada. <i>BMJ Open</i> , 2018, 8, e019701.	0.8	71
21	Place-based Experiences of Formal Healthcare Settings by People Experiencing Vulnerability at the End of Life. <i>Journal of Pain and Symptom Management</i> , 2018, 56, e56.	0.6	0
22	Design and introduction of a quality of life assessment and practice support system: perspectives from palliative care settings. <i>Journal of Patient-Reported Outcomes</i> , 2018, 2, 36.	0.9	20
23	Hospitals, clinics, and palliative care units: Place-based experiences of formal healthcare settings by people experiencing structural vulnerability at the end-of-life. <i>Health and Place</i> , 2018, 53, 43-51.	1.5	35
24	Discourses Reproducing Gender Inequities in Hospice Palliative Home Care. <i>Canadian Journal of Nursing Research</i> , 2018, 50, 189-201.	0.6	13
25	Embedding a Palliative Approach in Nursing Care Delivery. <i>Advances in Nursing Science</i> , 2017, 40, 263-279.	0.6	26
26	Structural impact on gendered expectations and exemptions for family caregivers in hospice palliative home care. <i>Nursing Inquiry</i> , 2017, 24, e12157.	1.1	12
27	Bereaved family members'™ perceptions of the quality of end-of-life care across four types of inpatient care settings. <i>BMC Palliative Care</i> , 2017, 16, 59.	0.8	19
28	When Cancer Hits the Streets. <i>Current Oncology</i> , 2017, 24, 149-150.	0.9	3
29	Rebuilding the Roots of Patient-Centred Care. <i>Canadian Journal of Nursing Leadership</i> , 2017, 30, 23-29.	0.6	1
30	“Close to”™ a palliative approach: nurses' and care aides' descriptions of caring for people with advancing chronic life-limiting conditions. <i>Journal of Clinical Nursing</i> , 2016, 25, 2189-2199.	1.4	20
31	RF1-B Experiences of Integrating Electronic Assessments Within Tertiary Palliative Care: Use of the Quality of Life Assessment and Practice Support System (QPSS). <i>Journal of Pain and Symptom Management</i> , 2016, 52, e5-e6.	0.6	0
32	RF2-C Barriers to Accessing Palliative Care for Structurally Vulnerable Populations. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e30.	0.6	1
33	G13-B A Scoping Review of Palliative Care for Those with Severe Persistent Mental Illness. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e63.	0.6	2
34	P102 Caregiver Coach Service: A Volunteer Service Aimed at Maintaining the Wellbeing of Family Caregivers. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e91.	0.6	0
35	P128 Desirable Features of a Quality of Life Assessment and Practice Support System (QPSS) in Palliative Acute and Home Care. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e99.	0.6	0
36	P335 Integrating Health Promotion and Palliative Care for a Palliative Approach: A Scoping Review. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e152-e153.	0.6	0

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37	Death Is a Social Justice Issue. <i>Advances in Nursing Science</i> , 2016, 39, 293-307.	0.6	59
38	Gendered Processes in Hospice Palliative Home Care for Seniors With Cancer and Their Family Caregivers. <i>Qualitative Health Research</i> , 2016, 26, 907-920.	1.0	10
39	Conceptual foundations of a palliative approach: a knowledge synthesis. <i>BMC Palliative Care</i> , 2016, 15, 5.	0.8	112
40	Identifying socio-environmental factors that facilitate resilience among Canadian palliative family caregivers: A qualitative case study. <i>Palliative and Supportive Care</i> , 2015, 13, 555-565.	0.6	26
41	What family caregivers learn when providing care at the end of life: A qualitative secondary analysis of multiple datasets. <i>Palliative and Supportive Care</i> , 2015, 13, 425-433.	0.6	15
42	Registered nurses' involvement in advance care planning: an integrative review. <i>International Journal of Palliative Nursing</i> , 2015, 21, 495-503.	0.2	30
43	OA–Caring for the family caregiver: working with volunteers to implement and improve a service to enable family caregivers to maintain their own wellbeing. <i>BMJ Supportive and Palliative Care</i> , 2015, 5, A3.1-A3.	0.8	2
44	Translational Scholarship and a Palliative Approach. <i>Advances in Nursing Science</i> , 2015, 38, 187-202.	0.6	6
45	Client, caregiver, and provider perspectives of safety in palliative home care: a mixed method design. <i>Safety in Health</i> , 2015, 1, .	0.7	17
46	(Re)theorizing Integrated Knowledge Translation. <i>Advances in Nursing Science</i> , 2015, 38, 175-186.	0.6	11
47	The Impact of the Carer Support Needs Assessment Tool (CSNAT) in Community Palliative Care Using a Stepped Wedge Cluster Trial. <i>PLoS ONE</i> , 2015, 10, e0123012.	1.1	86
48	Findings on Advance Care Plans among Cognitively Impaired Older Adults. <i>Canadian Journal on Aging</i> , 2015, 34, 165-175.	0.6	8
49	Palliative approach education for rural nurses and health-care workers: a mixed-method study. <i>International Journal of Palliative Nursing</i> , 2015, 21, 142-151.	0.2	11
50	Enabling a family caregiver-led assessment of support needs in home-based palliative care: Potential translation into practice. <i>Palliative Medicine</i> , 2015, 29, 929-938.	1.3	35
51	Supporting family caregivers to identify their own needs in end-of-life care: Qualitative findings from a stepped wedge cluster trial. <i>Palliative Medicine</i> , 2015, 29, 508-517.	1.3	76
52	Cultural influences on palliative family caregiving: service recommendations specific to the Vietnamese in Canada. <i>BMC Research Notes</i> , 2015, 8, 280.	0.6	5
53	Contrasting stories of life-threatening illness: A narrative inquiry. <i>International Journal of Nursing Studies</i> , 2015, 52, 207-215.	2.5	23
54	Examining the language“place“healthcare intersection in the context of Canadian homecare nursing. <i>Nursing Inquiry</i> , 2014, 21, 79-90.	1.1	9

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55	Educating Nurses for Palliative Care. <i>Journal of Hospice and Palliative Nursing</i> , 2014, 16, 47-54.	0.5	20
56	Applying Research Into Practice. <i>Home Healthcare Nurse</i> , 2014, 32, 88-95.	0.4	5
57	Changing communication needs and preferences across the cancer care trajectory: insights from the patient perspective. <i>Supportive Care in Cancer</i> , 2014, 22, 1009-1015.	1.0	92
58	Helene Hudson Lecture: Taking an "upstream" approach in the care of dying cancer patients: The case for a palliative approach. <i>Canadian Oncology Nursing Journal = Revue Canadienne De Nursing Oncologique</i> , 2014, 24, 144-148.	0.1	23
59	Taking an "upstream" approach in the care of dying cancer patients: the case for a palliative approach. <i>Canadian Oncology Nursing Journal = Revue Canadienne De Nursing Oncologique</i> , 2014, 24, 144-53.	0.1	13
60	Family caregiver learning "how family caregivers learn to provide care at the end of life: A qualitative secondary analysis of four datasets. <i>Palliative Medicine</i> , 2013, 27, 657-664.	1.3	57
61	Communicating shared decision-making: Cancer patient perspectives. <i>Patient Education and Counseling</i> , 2013, 90, 291-296.	1.0	75
62	A 10-year literature review of family caregiving for motor neurone disease: Moving from caregiver burden studies to palliative care interventions. <i>Palliative Medicine</i> , 2013, 27, 437-446.	1.3	116
63	Poor Communication in Cancer Care. <i>Cancer Nursing</i> , 2013, 36, 445-453.	0.7	42
64	Analysis and Proposed Model of Family Caregivers' Relationships With Home Health Providers and Perceptions of the Quality of Formal Services. <i>Journal of Applied Gerontology</i> , 2013, 32, 188-206.	1.0	18
65	Burdens of Family Caregiving at the End of Life. <i>Clinical and Investigative Medicine</i> , 2013, 36, 121.	0.3	50
66	Defining and measuring a palliative approach in primary care. <i>Canadian Family Physician</i> , 2013, 59, 1149-50.	0.1	23
67	Exposing barriers to end-of-life communication in heart failure: an integrative review. <i>Canadian Journal of Cardiovascular Nursing = Journal Canadien En Soins Infirmiers Cardio-vasculaires</i> , 2013, 23, 12-8.	0.1	6
68	End of Life in Residential Care from the Perspective of Care Aides. <i>Canadian Journal on Aging</i> , 2012, 31, 411-421.	0.6	18
69	Patient perceptions of communications on the threshold of cancer survivorship: implications for provider responses. <i>Journal of Cancer Survivorship</i> , 2012, 6, 229-237.	1.5	49
70	Family caregivers' ideal expectations of Canada's Compassionate Care Benefit. <i>Health and Social Care in the Community</i> , 2012, 20, 172-180.	0.7	5
71	People living with serious illness: stories of spirituality. <i>Journal of Clinical Nursing</i> , 2012, 21, 2347-2356.	1.4	35
72	Legitimising and rationalising in talk about satisfaction with formal healthcare among bereaved family members. <i>Sociology of Health and Illness</i> , 2012, 34, 1010-1024.	1.1	12

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73	End-of-life Care and Interprofessional Communication: Not Simply a Matter of "More". Health and Interprofessional Practice, 2012, 1, .	0.3	9
74	Perceptions regarding death and dying of individuals with chronic kidney disease. Nephrology Nursing Journal, 2012, 39, 197-204.	0.1	5
75	Bereaved Family Members' assessments of the quality of End-Of-Life Care: What is Important?. Journal of Palliative Care, 2011, 27, 261-269.	0.4	33
76	An exploration of empowerment discourse within home-care nurses' accounts of practice. Nursing Inquiry, 2011, 18, 66-76.	1.1	21
77	Home care nurses' decisions about the need for and amount of service at the end of life. Journal of Advanced Nursing, 2011, 67, 276-286.	1.5	40
78	The influence of culture on home-based family caregiving at end-of-life: A case study of Dutch reformed family care givers in Ontario, Canada. Social Science and Medicine, 2011, 72, 338-346.	1.8	35
79	Innovations in Research and Training in Palliative and End-of-Life Care: The Canadian Perspective (307). Journal of Pain and Symptom Management, 2011, 41, 183-184.	0.6	0
80	Canada's Compassionate Care Benefit: Is it an adequate public health response to addressing the issue of caregiver burden in end-of-life care?. BMC Public Health, 2011, 11, 335.	1.2	34
81	Exploring Family Caregivers' Rationales for Nonuse of Formal Home Health Services When Caring for a Dying Family Member. Home Health Care Management and Practice, 2011, 23, 318-328.	0.4	6
82	Articulating the Role of Relationships in Access to Home Care Nursing at the End of Life. Qualitative Health Research, 2011, 21, 117-131.	1.0	19
83	Chronic illness, palliative care, and the problematic nature of dying. Canadian Journal of Nursing Research, 2011, 43, 7-15.	0.6	27
84	Core aspects of "empowering" caregivers as articulated by leaders in home health care: palliative and chronic illness contexts. Canadian Journal of Nursing Research, 2011, 43, 78-94.	0.6	12
85	Case study: a model patient or impression management?. Nephrology Nursing Journal, 2011, 38, 511-3.	0.1	0
86	Bereaved family members' assessments of the quality of end-of-life care: what is important?. Journal of Palliative Care, 2011, 27, 261-9.	0.4	13
87	Patient perceptions of helpful communication in the context of advanced cancer. Journal of Clinical Nursing, 2010, 19, 2039-2047.	1.4	43
88	Helpful communications during the diagnostic period: an interpretive description of patient preferences. European Journal of Cancer Care, 2010, 19, 746-754.	0.7	38
89	A critical analysis of health promotion and "empowerment" in the context of palliative family care-giving. Nursing Inquiry, 2010, 17, 221-230.	1.1	37
90	A systematic review of instruments related to family caregivers of palliative care patients. Palliative Medicine, 2010, 24, 656-668.	1.3	98

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91	Part 1: Home-based family caregiving at the end of life: a comprehensive review of published quantitative research (1998-2008). <i>Palliative Medicine</i> , 2010, 24, 573-593.	1.3	275
92	Part 2: Home-based family caregiving at the end of life: a comprehensive review of published qualitative research (1998-2008). <i>Palliative Medicine</i> , 2010, 24, 594-607.	1.3	243
93	What makes grief difficult? Perspectives from bereaved family caregivers and healthcare providers of advanced cancer patients. <i>Palliative and Supportive Care</i> , 2010, 8, 277-289.	0.6	52
94	Palliative family caregivers' accounts of health care experiences: The importance of "security". <i>Palliative and Supportive Care</i> , 2009, 7, 435-447.	0.6	51
95	Supporting lay carers in end of life care: current gaps and future priorities. <i>Palliative Medicine</i> , 2009, 23, 339-344.	1.3	179
96	Interviewing Family Caregivers: Implications of the Caregiving Context for the Research Interview. <i>Qualitative Health Research</i> , 2009, 19, 859-867.	1.0	34
97	Patient Real-Time and 12-Month Retrospective Perceptions of Difficult Communications in the Cancer Diagnostic Period. <i>Qualitative Health Research</i> , 2009, 19, 1383-1394.	1.0	44
98	Resilience from the perspective of the illicit injection drug user: An exploratory descriptive study. <i>International Journal of Drug Policy</i> , 2009, 20, 309-316.	1.6	26
99	Time-related communication skills from the cancer patient perspective. <i>Psycho-Oncology</i> , 2009, 18, 500-507.	1.0	33
100	Use of palliative care services in a semirural program in British Columbia. <i>Canadian Journal of Rural Medicine: the Official Journal of the Society of Rural Physicians of Canada = Journal Canadien De La Médecine Rurale: Le Journal Officiel De La Société Des Médecins Ruraux Du Canada</i> , 2009, 14, 10-5.	0.2	1
101	Situated/being situated: Client and co-worker roles of family caregivers in hospice palliative care. <i>Social Science and Medicine</i> , 2008, 67, 1789-1797.	1.8	45
102	Nursing Graduate Supervision of Theses and Projects at a Distance: Issues and Challenges. <i>International Journal of Nursing Education Scholarship</i> , 2008, 5, 1-12.	0.4	7
103	Short Report: Preferences for location of death of seriously ill hospitalized patients: perspectives from Canadian patients and their family caregivers. <i>Palliative Medicine</i> , 2008, 22, 85-88.	1.3	69
104	Planning for End-of-Life Care: Findings from the Canadian Study of Health and Aging. <i>Canadian Journal on Aging</i> , 2008, 27, 11-21.	0.6	15
105	Factors Influencing Family Caregivers' Ability to Cope With Providing End-of-Life Cancer Care at Home. <i>Cancer Nursing</i> , 2008, 31, 77-85.	0.7	86
106	Planning for End-of-Life Care: Findings from the Canadian Study of Health and Aging. <i>Canadian Journal on Aging</i> , 2008, 27, 11-21.	0.6	11
107	Balancing dual roles in end-of-life research. <i>Canadian Oncology Nursing Journal = Revue Canadienne De Nursing Oncologique</i> , 2007, 17, 141-143.	0.1	1
108	The information transfer and knowledge acquisition geographies of family caregivers: an analysis of Canada's Compassionate Care Benefit. <i>Canadian Journal of Nursing Research</i> , 2007, 39, 36-54.	0.6	23

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109	Using Quality Improvement to Enhance Research Readiness in Palliative Care. <i>Journal for Healthcare Quality: Official Publication of the National Association for Healthcare Quality</i> , 2006, 28, 22-28.	0.3	4
110	Canada's Compassionate Care Benefit: Views of family caregivers in chronic illness. <i>International Journal of Palliative Nursing</i> , 2006, 12, 438-445.	0.2	16
111	The uses of provincial administrative health databases for research on palliative care: Insights from British Columbia, Canada. <i>BMC Palliative Care</i> , 2005, 4, 2.	0.8	16
112	Variations in and factors influencing family members' decisions for palliative home care. <i>Palliative Medicine</i> , 2005, 19, 21-32.	1.3	107
113	Missed opportunities: injection drug use and HIV/AIDS in Victoria, Canada. <i>International Journal of Drug Policy</i> , 2004, 15, 171-181.	1.6	9
114	Examining the Perspectives of Family Members Involved in the Delivery of Palliative Care at Home. <i>Journal of Palliative Care</i> , 2003, 19, 27-35.	0.4	114
115	Examining the perspectives of family members involved in the delivery of palliative care at home. <i>Journal of Palliative Care</i> , 2003, 19, 27-35.	0.4	36
116	A Qualitative Evaluation of an HIV/AIDS Respite Care Service in Victoria, Canada. <i>Evaluation and the Health Professions</i> , 2002, 25, 321-344.	0.9	3
117	Effective partnerships. Bringing nursing research alive in the practice setting. <i>The Canadian Nurse</i> , 2002, 98, 14-8.	0.0	0
118	A case for the "middle ground": exploring the tensions of postmodern thought in nursing. <i>Nursing Philosophy</i> , 2001, 2, 72-82.	0.9	14
119	Examining the process of community development. <i>Journal of Advanced Nursing</i> , 2001, 33, 828-835.	1.5	17
120	Thyroid cancer: patients' experiences of receiving iodine-131 therapy. <i>Oncology Nursing Forum</i> , 2000, 27, 1213-8.	0.5	32
121	Home Away from Home: Essential Elements in Developing Housing Options for People Living with HIV/AIDS. <i>AIDS Patient Care and STDs</i> , 1999, 13, 481-491.	1.1	2
122	Creating effective nursing partnerships: relating community development to participatory action research. <i>Journal of Advanced Nursing</i> , 1999, 29, 1238-1245.	1.5	19
123	Death at Home: Challenges for Families and Directions for the Future. <i>Journal of Palliative Care</i> , 1998, 14, 8-14.	0.4	81
124	Palliative Care at Home: Reflections on HIV/AIDS Family Caregiving Experiences. <i>Journal of Palliative Care</i> , 1998, 14, 14-22.	0.4	41