

Catherine Walshe

List of Publications by Year in descending order

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Version: 2024-02-01

84
papers

1,518
citations

361413

20
h-index

395702

33
g-index

92
all docs

92
docs citations

92
times ranked

1648
citing authors

#	ARTICLE	IF	CITATIONS
1	Patterns of Access to Community Palliative Care Services: A Literature Review. <i>Journal of Pain and Symptom Management</i> , 2009, 37, 884-912.	1.2	99
2	Coping Well with Advanced Cancer: A Serial Qualitative Interview Study with Patients and Family Carers. <i>PLoS ONE</i> , 2017, 12, e0169071.	2.5	77
3	District nurses' role in palliative care provision: A realist review. <i>International Journal of Nursing Studies</i> , 2010, 47, 1167-1183.	5.6	76
4	'Necessity is the mother of invention': Specialist palliative care service innovation and practice change in response to COVID-19. Results from a multinational survey (CovPall). <i>Palliative Medicine</i> , 2021, 35, 814-829.	3.1	67
5	The views of patients with severe chronic obstructive pulmonary disease on advance care planning: A qualitative study. <i>Palliative Medicine</i> , 2013, 27, 265-272.	3.1	59
6	The Challenges of Caring for People Dying From COVID-19: A Multinational, Observational Study (CovPall). <i>Journal of Pain and Symptom Management</i> , 2021, 62, 460-470.	1.2	57
7	What are the views of hospital-based generalist palliative care professionals on what facilitates or hinders collaboration with in-patient specialist palliative care teams? A systematically constructed narrative synthesis. <i>Palliative Medicine</i> , 2016, 30, 240-256.	3.1	56
8	Using observation as a data collection method to help understand patient and professional roles and actions in palliative care settings. <i>Palliative Medicine</i> , 2012, 26, 1048-1054.	3.1	55
9	What influences referrals within community palliative care services? A qualitative case study. <i>Social Science and Medicine</i> , 2008, 67, 137-146.	3.8	54
10	Improving training in spiritual care: a qualitative study exploring patient perceptions of professional educational requirements. <i>Palliative Medicine</i> , 2009, 23, 601-607.	3.1	52
11	Implementation and impact of the Gold Standards Framework in community palliative care: a qualitative study of three primary care trusts. <i>Palliative Medicine</i> , 2008, 22, 736-743.	3.1	48
12	Judgements about fellow professionals and the management of patients receiving palliative care in primary care: a qualitative study. <i>British Journal of General Practice</i> , 2008, 58, 264-272.	1.4	41
13	Peer support for people with advanced cancer: a systematically constructed scoping review of quantitative and qualitative evidence. <i>Current Opinion in Supportive and Palliative Care</i> , 2018, 12, 308-322.	1.3	35
14	How effective are volunteers at supporting people in their last year of life? A pragmatic randomised wait-list trial in palliative care (ELSA). <i>BMC Medicine</i> , 2016, 14, 203.	5.5	34
15	Understanding and addressing challenges for advance care planning in the COVID-19 pandemic: An analysis of the UK CovPall survey data from specialist palliative care services. <i>Palliative Medicine</i> , 2021, 35, 1225-1237.	3.1	34
16	The evaluation of complex interventions in palliative care: An exploration of the potential of case study research strategies. <i>Palliative Medicine</i> , 2011, 25, 774-781.	3.1	31
17	'Being with' or 'doing for'? How the role of an end-of-life volunteer befriender can impact patient wellbeing: interviews from a multiple qualitative case study (ELSA). <i>Supportive Care in Cancer</i> , 2018, 26, 3163-3172.	2.2	29
18	Using the 'Social Marketing Mix Framework' to explore recruitment barriers and facilitators in palliative care randomised controlled trials? A narrative synthesis review. <i>Palliative Medicine</i> , 2018, 32, 990-1009.	3.1	27

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19	Improving living and dying for people with advanced dementia living in care homes: a realist review of Namaste Care and other multisensory interventions. <i>BMC Geriatrics</i> , 2018, 18, 303.	2.7	27
20	A group intervention to improve quality of life for people with advanced dementia living in care homes: the Namaste feasibility cluster RCT. <i>Health Technology Assessment</i> , 2020, 24, 1-140.	2.8	26
21	“It’s not what they were expecting”: A systematic review and narrative synthesis of the role and experience of the hospital palliative care volunteer. <i>Palliative Medicine</i> , 2020, 34, 589-604.	3.1	24
22	Experiences of staff providing specialist palliative care during COVID-19: a multiple qualitative case study. <i>Journal of the Royal Society of Medicine</i> , 2022, 115, 220-230.	2.0	22
23	Specialist palliative care services response to ethnic minority groups with COVID-19: equal but inequitable? an observational study. <i>BMJ Supportive and Palliative Care</i> , 2021, , bmjspcare-2021-003083.	1.6	21
24	Protocol for the End-of-Life Social Action Study (ELSA): a randomised wait-list controlled trial and embedded qualitative case study evaluation assessing the causal impact of social action befriending services on end of life experience. <i>BMC Palliative Care</i> , 2016, 15, 60.	1.8	20
25	Translation and psychometric assessment of the Persian version of the Rhodes Index of Nausea, Vomiting and Retching (INVR) scale for the assessment of chemotherapy-induced nausea and vomiting. <i>European Journal of Cancer Care</i> , 2014, 23, 811-818.	1.5	19
26	A revised model for coping with advanced cancer. Mapping concepts from a longitudinal qualitative study of patients and carers coping with advanced cancer onto Folkman and Greer's theoretical model of appraisal and coping. <i>Psycho-Oncology</i> , 2018, 27, 229-235.	2.3	18
27	Evaluating partnership working: lessons for palliative care. <i>European Journal of Cancer Care</i> , 2007, 16, 48-54.	1.5	17
28	Namaste Care in nursing care homes for people with advanced dementia: protocol for a feasibility randomised controlled trial. <i>BMJ Open</i> , 2018, 8, e026531.	1.9	17
29	Perspectives of elders and their adult children of Black and minority ethnic heritage on end-of-life conversations: A meta-ethnography. <i>Palliative Medicine</i> , 2020, 34, 195-208.	3.1	16
30	Change in Activity of Palliative Care Services during the Covid-19 Pandemic: A Multinational Survey (CovPall). <i>Journal of Palliative Medicine</i> , 2022, 25, 465-471.	1.1	16
31	“Busyness” and the preclusion of quality palliative district nursing care. <i>Nursing Ethics</i> , 2013, 20, 893-903.	3.4	14
32	Are nurse-led chemotherapy clinics really nurse-led? An ethnographic study. <i>International Journal of Nursing Studies</i> , 2017, 69, 1-8.	5.6	14
33	Perceptions of a Good Death in Children with Life-Shortening Conditions: An Integrative Review. <i>Journal of Palliative Medicine</i> , 2019, 22, 714-723.	1.1	14
34	Peer support to maintain psychological wellbeing in people with advanced cancer: findings from a feasibility study for a randomised controlled trial. <i>BMC Palliative Care</i> , 2020, 19, 129.	1.8	14
35	Understanding the impact of the Covid-19 pandemic on delivery of rehabilitation in specialist palliative care services: An analysis of the CovPall-Rehab survey data. <i>Palliative Medicine</i> , 2022, 36, 319-331.	3.1	14
36	Ward social workers’ views of what facilitates or hinders collaboration with specialist palliative care team social workers: A grounded theory. <i>BMC Palliative Care</i> , 2018, 17, 7.	1.8	12

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37	Prohibit, Protect, or Adapt? The Changing Role of Volunteers in Palliative and Hospice Care Services During the COVID-19 Pandemic. A Multinational Survey (Covpall). <i>International Journal of Health Policy and Management</i> , 2021, , .	0.9	12
38	Protocol for a longitudinal qualitative interview study: maintaining psychological well-being in advanced cancerâ€”what can we learn from patientsâ€™ and carersâ€™ own coping strategies?: TableÂ1. <i>BMJ Open</i> , 2013, 3, e003046.	1.9	11
39	Communicative constructions of person-centred and non-person-centred caring in nurse-led consultations. <i>European Journal of Oncology Nursing</i> , 2019, 40, 10-21.	2.1	11
40	How Grandparents Experience the Death of a Grandchild With a Life-Limiting Condition. <i>Journal of Family Nursing</i> , 2019, 25, 109-127.	1.9	11
41	Whom to help? An exploration of the assessment of grief. <i>International Journal of Palliative Nursing</i> , 1997, 3, 132-137.	0.5	10
42	Understanding the bereavement experience of grandparents following the death of a grandchild from a lifeâ€limiting condition: A metaâ€ethnography. <i>Journal of Advanced Nursing</i> , 2019, 75, 1406-1417.	3.3	10
43	Peer Mentors for People with Advanced Cancer: Lessons Learnt from Recruiting and Training Peer Mentors for a Feasibility Randomized Controlled Trial. <i>Journal of Cancer Education</i> , 2021, 36, 710-718.	1.3	10
44	Smiles behind the masks: A systematic review and narrative synthesis exploring how family members of seriously ill or dying patients are supported during infectious disease outbreaks. <i>Palliative Medicine</i> , 2021, 35, 1452-1467.	3.1	10
45	What are the Emotional Experiences of Being a Volunteer in Palliative and End-of-Life Care Settings? A Systematic Review and Thematic Synthesis. <i>Journal of Pain and Symptom Management</i> , 2021, 62, e232-e247.	1.2	10
46	A poststructural rethinking of the ethics of technology in relation to the provision of palliative home care by district nurses. <i>Nursing Philosophy</i> , 2016, 17, 59-70.	1.9	9
47	Nurses' feelings of 'ownership' of palliative care patients: findings from a qualitative case study. <i>Progress in Palliative Care</i> , 2010, 18, 346-351.	1.2	8
48	Quality care as ethical care: a poststructural analysis of palliative and supportive district nursing care. <i>Nursing Inquiry</i> , 2016, 23, 12-23.	2.1	8
49	Exploring the experiences of living with Lewy body dementia: An integrative review. <i>Journal of Advanced Nursing</i> , 2021, 77, 4632-4645.	3.3	8
50	The impact on emotional well-being of being a palliative care volunteer: An interpretative phenomenological analysis. <i>Palliative Medicine</i> , 2021, , 026921632110647.	3.1	8
51	Palliative care research: State of play and journal direction. <i>Palliative Medicine</i> , 2017, 31, 3-4.	3.1	7
52	Quality of Life Trends in People With and Without Cancer Referred to Volunteer-Provided Palliative Care Services (ELSA): A Longitudinal Study. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 689-698.	1.2	7
53	A four-stage process for intervention description and guide development of a practice-based intervention: refining the Namaste Care intervention implementation specification for people with advanced dementia prior to a feasibility cluster randomised trial. <i>BMC Geriatrics</i> , 2019, 19, 275.	2.7	7
54	Perspectives on COVID-19 and palliative care research. <i>Palliative Medicine</i> , 2021, 35, 4-5.	3.1	7

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55	Aims, actions and advance care planning by district nurses providing palliative care: an ethnographic observational study. <i>British Journal of Community Nursing</i> , 2020, 25, 276-286.	0.4	6
56	COVID-19: A personal perspective. <i>Palliative Medicine</i> , 2020, 34, 687-688.	3.1	5
57	The experience of informal caregivers in providing patient care in hospitals in low- and middle-income countries: A qualitative meta-synthesis. <i>Journal of Health Services Research and Policy</i> , 2022, 27, 321-329.	1.7	5
58	Communication patterns in nurse-led chemotherapy clinics: A mixed-method study. <i>Patient Education and Counseling</i> , 2020, 103, 1538-1545.	2.2	4
59	Nonparametric time series summary statistics for high-frequency accelerometry data from individuals with advanced dementia. <i>PLoS ONE</i> , 2020, 15, e0239368.	2.5	4
60	Palliative care research: Has it come of age?. <i>Palliative Medicine</i> , 2013, 27, 883-884.	3.1	3
61	Mixed Method Research in Palliative Care. , 2019, , 1681-1699.		3
62	Hospital-based social workersâ€™ perceptions of generalist- and specialist-level palliative social work activities. <i>Journal of Social Work</i> , 2021, 21, 416-434.	1.4	3
63	What should we report? Lessons learnt from the development and implementation of serious adverse event reporting procedures in non-pharmacological trials in palliative care. <i>BMC Palliative Care</i> , 2021, 20, 19.	1.8	3
64	A good death in the child with life shortening illness: A qualitative multiple-case study. <i>Palliative Medicine</i> , 2021, 35, 1878-1888.	3.1	3
65	The experience of hospitalization in people with advanced chronic obstructive pulmonary disease: A qualitative, phenomenological study. <i>Chronic Illness</i> , 2023, 19, 339-353.	1.5	3
66	Symptom Control and Survival for People Severely ill With COVID: A Multicentre Cohort Study (CovPall-Symptom). <i>Journal of Pain and Symptom Management</i> , 2022, 64, 377-390.	1.2	3
67	Protocol for a systematic review on the experience of informal caregivers for people with a moderate to advanced dementia within a domestic home setting. <i>Systematic Reviews</i> , 2020, 9, 270.	5.3	2
68	Exploring the psychological impact of life-limiting illness using the Attitude to Health Change scales: A qualitative focus group study in a hospice palliative care setting. <i>European Journal of Cancer Care</i> , 2020, 29, e13302.	1.5	2
69	Constructing a new role for family carers and volunteers providing care towards the end of life: an action research approach exploring a new model of hospice care. <i>Health and Social Care in the Community</i> , 2021, 29, 837-845.	1.6	2
70	A Systematic Review with Thematic Synthesis of the Experience of Hospitalization in People with Advanced Chronic Obstructive Pulmonary Disease. <i>COPD: Journal of Chronic Obstructive Pulmonary Disease</i> , 2021, 18, 1-9.	1.6	2
71	Do journals contribute to the international publication of research in their field? A bibliometric analysis of palliative care journal data. <i>Palliative Medicine</i> , 2020, 34, 541-546.	3.1	2
72	Exploring a New Model of End-of-Life Care for Older People That Operates in the Space Between the Life World and the Healthcare System: A Qualitative Case Study. <i>International Journal of Health Policy and Management</i> , 2020, 9, 344-351.	0.9	2

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73	Mixed Method Research in Palliative Care. , 2018, , 1-19.		1
74	Exploring the experience of recurrence with advanced cancer for people who perceived themselves to be cancer free: a grounded theory study. Supportive Care in Cancer, 2021, 29, 3885-3894.	2.2	1
75	What Do We Mean by Palliative Care. , 2011, , 17-29.		1
76	New authorsâ€™ guidelines for Palliative Medicine: More work for authors, reviewers and editors or an essential tool?. Palliative Medicine, 2010, 24, 559-560.	3.1	0
77	Supporting and promoting palliative care for older people. Journal of Health Visiting, 2013, 1, 27-30.	0.1	0
78	ADVANCE CARE PLANNING TRAINING IN NURSING HOMES: A SUCCESS STORY?. BMJ Supportive and Palliative Care, 2014, 4, A32.3-A33.	1.6	0
79	THE VALUE OF LONGITUDINAL INTERVIEWS IN EXPLORING COPING STRATEGIES OF PATIENTS WITH ADVANCED CANCER AND THEIR CARERS. BMJ Supportive and Palliative Care, 2014, 4, A40.1-A40.	1.6	0
80	HEALTH PROFESSIONALSâ€™ PERSPECTIVES ABOUT ADVANCE CARE PLANNING IN PARKINSON'S DISEASE: A QUALITATIVE STUDY. BMJ Supportive and Palliative Care, 2014, 4, A45.1-A45.	1.6	0
81	<i>Palliative Medicine</i> journal: What have we done, where should we be going and why?. Palliative Medicine, 2015, 29, 673-674.	3.1	0
82	Applying an Analytical Process to Longitudinal Narrative Interviews With Couples Living and Dying With Lewy Body Dementia. International Journal of Qualitative Methods, The, 2021, 20, 160940692110606.	2.8	0
83	Exploring Specialist Palliative Care Practitioner Perspectives on the Face Validity of the Attitude to Health Change Scales in Assessing the Impact of Life-limiting Illness on Patients and Carers. Journal of Palliative Care, 2021, , 082585972110640.	1.0	0
84	A tribute to Derek Doyle and Cynthia Goh. Palliative Medicine, 0, , 026921632211093.	3.1	0