

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	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
2	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
3	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
4	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
5	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
6	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
7	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
8	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
9	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
10	Perspectives on COVID-19 and palliative care research. <i>Palliative Medicine</i> , 2021, 35, 4-5.	3.1	7
11	Exploring the experience of recurrence with advanced cancer for people who perceived themselves to be cancer free: a grounded theory study. <i>Supportive Care in Cancer</i> , 2021, 29, 3885-3894.	2.2	1
12	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
13	â€˜Necessity is the mother of inventionâ€™: 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
14	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
15	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
16	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
17	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
18	Specialist palliative care services response to ethnic minority groups with COVID-19: equal but inequitableâ€™ an observational study. <i>BMJ Supportive and Palliative Care</i> , 2021, , bmjpspcare-2021-003083.	1.6	21

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19	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
20	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
21	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
22	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
23	Applying an Analytical Process to Longitudinal Narrative Interviews With Couples Living and Dying With Lewy Body Dementia. <i>International Journal of Qualitative Methods</i> , The, 2021, 20, 160940692110606.	2.8	0
24	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. <i>Journal of Palliative Care</i> , 2021, , 082585972110640.	1.0	0
25	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
26	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
27	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
28	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
29	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
30	Communication patterns in nurse-led chemotherapy clinics: A mixed-method study. <i>Patient Education and Counseling</i> , 2020, 103, 1538-1545.	2.2	4
31	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
32	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
33	COVID-19: A personal perspective. <i>Palliative Medicine</i> , 2020, 34, 687-688.	3.1	5
34	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
35	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
36	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

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37	Nonparametric time series summary statistics for high-frequency accelerometry data from individuals with advanced dementia. PLoS ONE, 2020, 15, e0239368.	2.5	4
38	Mixed Method Research in Palliative Care. , 2019, , 1681-1699.		3
39	Communicative constructions of person-centred and non-person-centred caring in nurse-led consultations. European Journal of Oncology Nursing, 2019, 40, 10-21.	2.1	11
40	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. BMC Geriatrics, 2019, 19, 275.	2.7	7
41	Understanding the bereavement experience of grandparents following the death of a grandchild from a life-limiting condition: A meta-ethnography. Journal of Advanced Nursing, 2019, 75, 1406-1417.	3.3	10
42	How Grandparents Experience the Death of a Grandchild With a Life-Limiting Condition. Journal of Family Nursing, 2019, 25, 109-127.	1.9	11
43	Perceptions of a Good Death in Children with Life-Shortening Conditions: An Integrative Review. Journal of Palliative Medicine, 2019, 22, 714-723.	1.1	14
44	Using the "Social Marketing Mix Framework"™ to explore recruitment barriers and facilitators in palliative care randomised controlled trials? A narrative synthesis review. Palliative Medicine, 2018, 32, 990-1009.	3.1	27
45	"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). Supportive Care in Cancer, 2018, 26, 3163-3172.	2.2	29
46	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. Psycho-Oncology, 2018, 27, 229-235.	2.3	18
47	Mixed Method Research in Palliative Care. , 2018, , 1-19.		1
48	Improving living and dying for people with advanced dementia living in care homes: a realist review of Namaste Care and other multisensory interventions. BMC Geriatrics, 2018, 18, 303.	2.7	27
49	Namaste Care in nursing care homes for people with advanced dementia: protocol for a feasibility randomised controlled trial. BMJ Open, 2018, 8, e026531.	1.9	17
50	Peer support for people with advanced cancer: a systematically constructed scoping review of quantitative and qualitative evidence. Current Opinion in Supportive and Palliative Care, 2018, 12, 308-322.	1.3	35
51	Ward social workers'™ views of what facilitates or hinders collaboration with specialist palliative care team social workers: A grounded theory. BMC Palliative Care, 2018, 17, 7.	1.8	12
52	Quality of Life Trends in People With and Without Cancer Referred to Volunteer-Provided Palliative Care Services (ELSA): A Longitudinal Study. Journal of Pain and Symptom Management, 2018, 56, 689-698.	1.2	7
53	Are nurse-led chemotherapy clinics really nurse-led? An ethnographic study. International Journal of Nursing Studies, 2017, 69, 1-8.	5.6	14
54	Palliative care research: State of play and journal direction. Palliative Medicine, 2017, 31, 3-4.	3.1	7

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55	Coping Well with Advanced Cancer: A Serial Qualitative Interview Study with Patients and Family Carers. PLoS ONE, 2017, 12, e0169071.	2.5	77
56	A poststructural rethinking of the ethics of technology in relation to the provision of palliative home care by district nurses. Nursing Philosophy, 2016, 17, 59-70.	1.9	9
57	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. BMC Palliative Care, 2016, 15, 60.	1.8	20
58	How effective are volunteers at supporting people in their last year of life? A pragmatic randomised wait-list trial in palliative care (ELSA). BMC Medicine, 2016, 14, 203.	5.5	34
59	Quality care as ethical care: a poststructural analysis of palliative and supportive district nursing care. Nursing Inquiry, 2016, 23, 12-23.	2.1	8
60	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. Palliative Medicine, 2016, 30, 240-256.	3.1	56
61	<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
62	ADVANCE CARE PLANNING TRAINING IN NURSING HOMES: A SUCCESS STORY?. BMJ Supportive and Palliative Care, 2014, 4, A32.3-A33.	1.6	0
63	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
64	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. European Journal of Cancer Care, 2014, 23, 811-818.	1.5	19
65	HEALTH PROFESSIONALS'S 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
66	Protocol for a longitudinal qualitative interview study: maintaining psychological well-being in advanced cancer's what can we learn from patients' and carers' own coping strategies?: Table 1. BMJ Open, 2013, 3, e003046.	1.9	11
67	'Business' and the preclusion of quality palliative district nursing care. Nursing Ethics, 2013, 20, 893-903.	3.4	14
68	Palliative care research: Has it come of age?. Palliative Medicine, 2013, 27, 883-884.	3.1	3
69	The views of patients with severe chronic obstructive pulmonary disease on advance care planning: A qualitative study. Palliative Medicine, 2013, 27, 265-272.	3.1	59
70	Supporting and promoting palliative care for older people. Journal of Health Visiting, 2013, 1, 27-30.	0.1	0
71	Using observation as a data collection method to help understand patient and professional roles and actions in palliative care settings. Palliative Medicine, 2012, 26, 1048-1054.	3.1	55
72	The evaluation of complex interventions in palliative care: An exploration of the potential of case study research strategies. Palliative Medicine, 2011, 25, 774-781.	3.1	31

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73	What Do We Mean by Palliative Care. , 2011, , 17-29.		1
74	Nurses' feelings of 'ownership' of palliative care patients: findings from a qualitative case study. Progress in Palliative Care, 2010, 18, 346-351.	1.2	8
75	District nurses' role in palliative care provision: A realist review. International Journal of Nursing Studies, 2010, 47, 1167-1183.	5.6	76
76	New authors' 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	Improving training in spiritual care: a qualitative study exploring patient perceptions of professional educational requirements. Palliative Medicine, 2009, 23, 601-607.	3.1	52
78	Patterns of Access to Community Palliative Care Services: A Literature Review. Journal of Pain and Symptom Management, 2009, 37, 884-912.	1.2	99
79	What influences referrals within community palliative care services? A qualitative case study. Social Science and Medicine, 2008, 67, 137-146.	3.8	54
80	Implementation and impact of the Gold Standards Framework in community palliative care: a qualitative study of three primary care trusts. Palliative Medicine, 2008, 22, 736-743.	3.1	48
81	Judgements about fellow professionals and the management of patients receiving palliative care in primary care: a qualitative study. British Journal of General Practice, 2008, 58, 264-272.	1.4	41
82	Evaluating partnership working: lessons for palliative care. European Journal of Cancer Care, 2007, 16, 48-54.	1.5	17
83	Whom to help? An exploration of the assessment of grief. International Journal of Palliative Nursing, 1997, 3, 132-137.	0.5	10
84	A tribute to Derek Doyle and Cynthia Goh. Palliative Medicine, 0, , 026921632211093.	3.1	0