## Barbara A Daveson

List of Publications by Year in descending order

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84 papers

3,403 citations

33 h-index 55 g-index

88 all docs 88 docs citations

88 times ranked 3783 citing authors

#	Article	IF	CITATIONS
1	Preferences of Older People With a Life-Limiting Illness: A Discrete Choice Experiment. Journal of Pain and Symptom Management, 2022, 64, 137-145.	1.2	2
2	The PCOC Symptom Assessment Scale (SAS): A valid measure for daily use at point of care and in palliative care programs. PLoS ONE, 2021, 16, e0247250.	2.5	24
3	Sleeping-related distress in a palliative care population: A national, prospective, consecutive cohort. Palliative Medicine, 2021, 35, 1663-1670.	3.1	5
4	Patient-reported outcome measurement of symptom distress is feasible in most clinical scenarios in palliative care: an observational study involving routinely collected data. International Journal for Quality in Health Care, 2021, 33, .	1.8	15
5	Appetite-Related Distress Is Burdensome in the Last Sixty Days of Life of People Receiving Palliative Care: A National Longitudinal Consecutive Cohort Study. Journal of Palliative Medicine, 2021, , .	1.1	1
6	Associations between informal care costs, care quality, carer rewards, burden and subsequent grief: the international, access, rights and empowerment mortality follow-back study of the last 3Âmonths of life (IARE I study). BMC Medicine, 2020, 18, 344.	5 <b>.</b> 5	17
7	Drivers of care costs and quality in the last 3 months of life among older people receiving palliative care: A multinational mortality follow-back survey across England, Ireland and the United States. Palliative Medicine, 2020, 34, 513-523.	3.1	11
8	Optimal Cancer Care for Aboriginal and Torres Strait Islander People: A Shared Approach to System Level Change. JCO Global Oncology, 2020, 6, 108-114.	1.8	12
9	Patterns of emergency department attendance among older people in the last three months of life and factors associated with frequent attendance: a mortality follow-back survey. Age and Ageing, 2019, 48, 680-687.	1.6	10
10	A framework for complexity in palliative care: A qualitative study with patients, family carers and professionals. Palliative Medicine, 2018, 32, 1078-1090.	3.1	85
11	Strategies to address the shortcomings of commonly used advanced chronic heart failure descriptors to improve recruitment in palliative care research: A parallel mixed-methods feasibility study. Palliative Medicine, 2018, 32, 517-524.	3.1	7
12	Recommendations to reduce inequalities for LGBT people facing advanced illness: ACCESSCare national qualitative interview study. Palliative Medicine, 2018, 32, 23-35.	3.1	88
13	Understanding how a palliative-specific patient-reported outcome intervention works to facilitate patient-centred care in advanced heart failure: A qualitative study. Palliative Medicine, 2018, 32, 143-155.	3.1	39
14	Perspectives of patients, family caregivers and health professionals on the use of outcome measures in palliative care and lessons for implementation: a multi-method qualitative study. Annals of Palliative Medicine, 2018, 7, S137-S150.	1.2	28
15	How can a measure improve assessment and management of symptoms and concerns for people with dementia in care homes? A mixed-methods feasibility and process evaluation of IPOS-Dem. PLoS ONE, 2018, 13, e0200240.	2.5	59
16	How empowering is hospital care for older people with advanced disease? Barriers and facilitators from a cross-national ethnography in England, Ireland and the USA. Age and Ageing, 2017, 46, 300-309.	1.6	12
17	Professional Supervision as Storied Experience: Narrative Analysis Findings for Australian-Based Registered Music Therapists. Journal of Music Therapy, 2017, 54, 80-107.	0.9	1
18	Feasibility and acceptability of a patient-reported outcome intervention in chronic heart failure. BMJ Supportive and Palliative Care, 2017, 7, 470-479.	1.6	42

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19	Development of a caregiver-reported measure to support systematic assessment of people with dementia in long-term care: The Integrated Palliative care Outcome Scale for Dementia. Palliative Medicine, 2017, 31, 651-660.	3.1	57
20	Social and clinical determinants of preferences and their achievement at the end of life: prospective cohort study of older adults receiving palliative care in three countries. BMC Geriatrics, 2017, 17, 271.	2.7	110
21	My body's falling apart.' Understanding the experiences of patients with advanced multimorbidity to improve care: serial interviews with patients and carers. BMJ Supportive and Palliative Care, 2016, 6, 60-65.	1.6	66
22	"What makes it worth the investment to say it?―– Care experiences of lesbian, gay, bisexual and/or trans* people facing life-limiting illness: a qualitative interview study. BMJ Supportive and Palliative Care, 2016, 6, 388.1-388.	1.6	2
23	"l'll be in a safe place†a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. BMJ Supportive and Palliative Care, 2016, 6, 394.2-394.	1.6	3
24	â€ï'lll be in a safe place': a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. BMJ Open, 2016, 6, e012134.	1.9	45
25	RF1-A How Empowering Is Hospital Care for Older Adults, and What Difference Does Palliative Care Make? A Cross-National Ethnography in England, Ireland and the USA. Journal of Pain and Symptom Management, 2016, 52, e5.	1.2	0
26	Of apples and oranges: Lessons learned from the preparation of research protocols for systematic reviews exploring the effectiveness of Specialist Palliative Care. BMC Palliative Care, 2016, 15, 43.	1.8	11
27	Measures to assess commonly experienced symptoms for people with dementia in long-term care settings: a systematic review. BMC Medicine, 2016, 14, 38.	5.5	32
28	Music Therapy Assessment Tool for Awareness in Disorders of Consciousness (MATADOC): Reliability and Validity of a Measure to Assess Awareness in Patients with Disorders of Consciousness. Journal of Music Therapy, 2016, 53, 1-26.	0.9	27
29	EAPC White Paper on outcome measurement in palliative care: Improving practice, attaining outcomes and delivering quality services – Recommendations from the European Association for Palliative Care (EAPC) Task Force on Outcome Measurement. Palliative Medicine, 2016, 30, 6-22.	3.1	149
30	Effects of professional music therapy supervision on clinical outcomes and therapist competency: a systematic review involving narrative synthesis. Nordic Journal of Music Therapy, 2016, 25, 185-208.	1.1	9
31	Factors associated with aggressive end of life cancer care. Supportive Care in Cancer, 2016, 24, 1079-1089.	2.2	64
32	Charting the Terrain of Grounded Theory Research in Music Therapy. , 2015, , .		2
33	Capture, Transfer, and Feedback of Patient-Centered Outcomes Data in Palliative Care Populations: Does It Make a Difference? A Systematic Review. Journal of Pain and Symptom Management, 2015, 49, 611-624.	1.2	158
34	Emergency Department Attendance by Patients With Cancer in Their Last Month of Life: A Systematic Review and Meta-Analysis. Journal of Clinical Oncology, 2015, 33, 370-376.	1.6	99
35	Results of a transparent expert consultation on patient and public involvement in palliative care research. Palliative Medicine, 2015, 29, 939-949.	3.1	35
36	The Need for Palliative Care in Ireland: A Population-Based Estimate of Palliative Care Using Routine Mortality Data, Inclusive of Nonmalignant Conditions. Journal of Pain and Symptom Management, 2015, 49, 726-733.e1.	1.2	54

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37	Emergency department attendance by patients with cancer in the last month of life: a systematic review and meta-analysis. Lancet, The, 2015, 385, S41.	13.7	19
38	The gap between policy and practice: a systematic review of patient-centred care interventions in chronic heart failure. Heart Failure Reviews, 2015, 20, 673-687.	3.9	58
39	A Systematic Review of the Use of the Palliative Care Outcome Scale and the Support Team Assessment Schedule in Palliative Care. Journal of Pain and Symptom Management, 2015, 50, 842-853.e19.	1.2	75
40	The Real-World Problem of Care Coordination: A Longitudinal Qualitative Study with Patients Living with Advanced Progressive Illness and Their Unpaid Caregivers. PLoS ONE, 2014, 9, e95523.	2.5	37
41	Place and Cause of Death in Centenarians: A Population-Based Observational Study in England, 2001 to 2010. PLoS Medicine, 2014, 11, e1001653.	8.4	96
42	Learning from the public: citizens describe the need to improve end-of-life care access, provision and recognition across Europe. European Journal of Public Health, 2014, 24, 521-527.	0.3	18
43	Priorities for treatment, care and information if faced with serious illness: A comparative population-based survey in seven European countries. Palliative Medicine, 2014, 28, 101-110.	3.1	86
44	Music Therapy Assessment Tool for Awareness in Disorders of Consciousness (MATADOC): Standardisation of the principal subscale to assess awareness in patients with disorders of consciousness. Neuropsychological Rehabilitation, 2014, 24, 101-124.	1.6	45
45	Choosing care homes as the least preferred place to die: a cross-national survey of public preferences in seven European countries. BMC Palliative Care, 2014, 13, 48.	1.8	26
46	Public preferences and priorities for end-of-life care in Kenya: a population-based street survey. BMC Palliative Care, 2014, 13, 4.	1.8	41
47	"A quiet still voice that just touches― music's relevance for adults living with life-threatening cancer diagnoses. Supportive Care in Cancer, 2014, 22, 1037-1047.	2.2	23
48	Coordination of end-of-life care for patients with lung cancer and those with advanced COPD: are there transferable lessons? A longitudinal qualitative study. Primary Care Respiratory Journal: Journal of the General Practice Airways Group, 2014, 23, 46-51.	2.3	38
49	Avoid †prognostic paralysis†m— just get ahead and plan and co-ordinate care. Npj Primary Care Respiratory Medicine, 2014, 24, 14085.	2.6	9
50	†Burden to others' as a public concern in advanced cancer: a comparative survey in seven European countries. BMC Cancer, 2013, 13, 105.	2.6	35
51	If you had less than a year to live, would you want to know? A sevenâ€country European population survey of public preferences for disclosure of poor prognosis. Psycho-Oncology, 2013, 22, 2298-2305.	2.3	37
52	The Selection and Use of Outcome Measures in Palliative and End-of-Life Care Research: The MORECare International Consensus Workshop. Journal of Pain and Symptom Management, 2013, 46, 925-937.	1.2	61
53	Social-cultural factors in end-of-life care in Belgium: A scoping of the research literature. Palliative Medicine, 2013, 27, 131-143.	3.1	14
54	To be involved or not to be involved: A survey of public preferences for self-involvement in decision-making involving mental capacity (competency) within Europe. Palliative Medicine, 2013, 27, 418-427.	3.1	41

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55	Coordination of care for individuals with advanced progressive conditions: a multi-site ethnographic and serial interview study. British Journal of General Practice, 2013, 63, e580-e588.	1.4	51
56	Diversity in Defining End of Life Care: An Obstacle or the Way Forward?. PLoS ONE, 2013, 8, e68002.	2.5	30
57	Public opinion on preferences and priorities for end-of-life care in sub-Saharan Africa: piloting a novel method of street surveying: Table 1. BMJ Supportive and Palliative Care, 2012, 2, 72-74.	1.6	3
58	Preferences for place of death if faced with advanced cancer: a population survey in England, Flanders, Germany, Italy, the Netherlands, Portugal and Spain. Annals of Oncology, 2012, 23, 2006-2015.	1.2	430
59	Republished: Which questions of two commonly used multidimensional palliative care patient reported outcome measures are most useful? Results from the European and African PRISMA survey. Postgraduate Medical Journal, 2012, 88, 451-457.	1.8	15
60	Are we heading in the same direction? European and African doctors' and nurses' views and experiences regarding outcome measurement in palliative care. Palliative Medicine, 2012, 26, 242-249.	3.1	27
61	Which questions of two commonly used multidimensional palliative care patient reported outcome measures are most useful? Results from the European and African PRISMA survey. BMJ Supportive and Palliative Care, 2012, 2, 36-42.	1.6	17
62	Cultural Competence in End-of-Life Care: Terms, Definitions, and Conceptual Models from the British Literature. Journal of Palliative Medicine, 2012, 15, 812-820.	1.1	31
63	Culture Is a Priority for Research in End-of-Life Care in Europe: A Research Agenda. Journal of Pain and Symptom Management, 2012, 44, 285-294.	1.2	27
64	Outcomes 'out of africa': the selection and implementation of outcome measures for palliative care in Africa. BMC Palliative Care, 2012, 11, 1.	1.8	38
65	Enhancing patient-reported outcome measurement in research and practice of palliative and end-of-life care. Supportive Care in Cancer, 2012, 20, 1573-1578.	2.2	29
66	Systematic Review of the Primary Research on Minority Ethnic Groups and End-of-Life Care From the United Kingdom. Journal of Pain and Symptom Management, 2012, 43, 261-286.	1.2	57
67	A critical review of advance directives in Germany: Attitudes, use and healthcare professionals' compliance. Patient Education and Counseling, 2012, 87, 277-288.	2.2	64
68	A pan-European survey of research in end-of-life cancer care. Supportive Care in Cancer, 2012, 20, 39-48.	2.2	21
69	The use of two common palliative outcome measures in clinical care and research: A systematic review of POS and STAS. Palliative Medicine, 2011, 25, 304-313.	3.1	89
70	The PRISMA Symposium 4: How Should Europe Progress End-of-Life and Palliative Clinical Care Research? Recommendations From the Proceedings. Journal of Pain and Symptom Management, 2011, 42, 511-516.	1.2	13
71	The PRISMA Symposium 1: Outcome Tool Use. Disharmony in European Outcomes Research for Palliative and Advanced Disease Care: Too Many Tools in Practice. Journal of Pain and Symptom Management, 2011, 42, 493-500.	1.2	43
72	Implementing patient reported outcome measures (PROMs) in palliative care - users' cry for help. Health and Quality of Life Outcomes, 2011, 9, 27.	2.4	104

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73	Constructing Understandings of End-of-Life Care in Europe: A Qualitative Study Involving Cognitive Interviewing with Implications for Cross-National Surveys. Journal of Palliative Medicine, 2011, 14, 343-349.	1.1	18
74	Music Therapy Methods with Children, Adolescents, and Adults with Severe Neurobehavioral Disorders Due to Brain Injury. Music Therapy Perspectives, 2011, 29, 5-13.	0.5	7
75	Investigating the Dimension of Time: Findings from a Modified Grounded Theory Study about Clients' Experiences and Descriptions of Temporality or Time within Music Therapy. Journal of Music Therapy, 2011, 48, 28-54.	0.9	6
76	Reflecting on the Long-term (Neurological) End-of-Life Conference. British Journal of Neuroscience Nursing, 2010, 6, 188-190.	0.2	1
77	An Audit about Music Therapy Assessments and Recommendations for Adult Patients Suspected to be in a Low, Awareness State. Journal of Music Therapy, 2010, 47, 408-422.	0.9	11
78	Indigenous music therapy theory building through grounded theory research: The developing indigenous theory framework. Arts in Psychotherapy, 2008, 35, 280-286.	1.2	13
79	A Project Investigating Music Therapy Referral Trends within Palliative Care: An Australian Perspective. Journal of Music Therapy, 2007, 44, 139-155.	0.9	18
80	The music therapy assessment tool for low awareness states. International Journal of Therapy and Rehabilitation, 2007, 14, 544-549.	0.3	9
81	Music therapy student supervision: considering aspects of resistance and parallel processes in the supervisory relationship with students in final clinical placement. Arts in Psychotherapy, 2004, 31, 67-76.	1.2	16
82	A philosophical inquiry into the role of rhythm in music therapy. Arts in Psychotherapy, 2002, 29, 265-270.	1.2	5
83	A descriptive study exploring the role of music therapy in prisons. Arts in Psychotherapy, 2001, 28, 137-141.	1.2	29
84	A Role for Music Therapy in Special Education. International Journal of Disability Development and Education, 1998, 45, 449-457.	1.1	11