

Barbara A Daveson

List of Publications by Year in descending order

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Version: 2024-02-01

84
papers

3,403
citations

126907

33
h-index

155660

55
g-index

88
all docs

88
docs citations

88
times ranked

3783
citing authors

#	ARTICLE	IF	CITATIONS
1	Preferences for place of death if faced with advanced cancer: a population survey in England, Flanders, Germany, Italy, the Netherlands, Portugal and Spain. <i>Annals of Oncology</i> , 2012, 23, 2006-2015.	1.2	430
2	Capture, Transfer, and Feedback of Patient-Centered Outcomes Data in Palliative Care Populations: Does It Make a Difference? A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2015, 49, 611-624.	1.2	158
3	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. <i>Palliative Medicine</i> , 2016, 30, 6-22.	3.1	149
4	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. <i>BMC Geriatrics</i> , 2017, 17, 271.	2.7	110
5	Implementing patient reported outcome measures (PROMs) in palliative care - users' cry for help. <i>Health and Quality of Life Outcomes</i> , 2011, 9, 27.	2.4	104
6	Emergency Department Attendance by Patients With Cancer in Their Last Month of Life: A Systematic Review and Meta-Analysis. <i>Journal of Clinical Oncology</i> , 2015, 33, 370-376.	1.6	99
7	Place and Cause of Death in Centenarians: A Population-Based Observational Study in England, 2001 to 2010. <i>PLoS Medicine</i> , 2014, 11, e1001653.	8.4	96
8	The use of two common palliative outcome measures in clinical care and research: A systematic review of POS and STAS. <i>Palliative Medicine</i> , 2011, 25, 304-313.	3.1	89
9	Recommendations to reduce inequalities for LGBT people facing advanced illness: ACCESSCare national qualitative interview study. <i>Palliative Medicine</i> , 2018, 32, 23-35.	3.1	88
10	Priorities for treatment, care and information if faced with serious illness: A comparative population-based survey in seven European countries. <i>Palliative Medicine</i> , 2014, 28, 101-110.	3.1	86
11	A framework for complexity in palliative care: A qualitative study with patients, family carers and professionals. <i>Palliative Medicine</i> , 2018, 32, 1078-1090.	3.1	85
12	A Systematic Review of the Use of the Palliative Care Outcome Scale and the Support Team Assessment Schedule in Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2015, 50, 842-853.e19.	1.2	75
13	“My body's falling apart.” Understanding the experiences of patients with advanced multimorbidity to improve care: serial interviews with patients and carers. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 60-65.	1.6	66
14	A critical review of advance directives in Germany: Attitudes, use and healthcare professionals' compliance. <i>Patient Education and Counseling</i> , 2012, 87, 277-288.	2.2	64
15	Factors associated with aggressive end of life cancer care. <i>Supportive Care in Cancer</i> , 2016, 24, 1079-1089.	2.2	64
16	The Selection and Use of Outcome Measures in Palliative and End-of-Life Care Research: The MORECare International Consensus Workshop. <i>Journal of Pain and Symptom Management</i> , 2013, 46, 925-937.	1.2	61
17	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. <i>PLoS ONE</i> , 2018, 13, e0200240.	2.5	59
18	The gap between policy and practice: a systematic review of patient-centred care interventions in chronic heart failure. <i>Heart Failure Reviews</i> , 2015, 20, 673-687.	3.9	58

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19	Systematic Review of the Primary Research on Minority Ethnic Groups and End-of-Life Care From the United Kingdom. <i>Journal of Pain and Symptom Management</i> , 2012, 43, 261-286.	1.2	57
20	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. <i>Palliative Medicine</i> , 2017, 31, 651-660.	3.1	57
21	The Need for Palliative Care in Ireland: A Population-Based Estimate of Palliative Care Using Routine Mortality Data, Inclusive of Nonmalignant Conditions. <i>Journal of Pain and Symptom Management</i> , 2015, 49, 726-733.e1.	1.2	54
22	Coordination of care for individuals with advanced progressive conditions: a multi-site ethnographic and serial interview study. <i>British Journal of General Practice</i> , 2013, 63, e580-e588.	1.4	51
23	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. <i>Neuropsychological Rehabilitation</i> , 2014, 24, 101-124.	1.6	45
24	‘I’ll be in a safe place’: a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. <i>BMJ Open</i> , 2016, 6, e012134.	1.9	45
25	The PRISMA Symposium 1: Outcome Tool Use. Disharmony in European Outcomes Research for Palliative and Advanced Disease Care: Too Many Tools in Practice. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 493-500.	1.2	43
26	Feasibility and acceptability of a patient-reported outcome intervention in chronic heart failure. <i>BMJ Supportive and Palliative Care</i> , 2017, 7, 470-479.	1.6	42
27	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. <i>Palliative Medicine</i> , 2013, 27, 418-427.	3.1	41
28	Public preferences and priorities for end-of-life care in Kenya: a population-based street survey. <i>BMC Palliative Care</i> , 2014, 13, 4.	1.8	41
29	Understanding how a palliative-specific patient-reported outcome intervention works to facilitate patient-centred care in advanced heart failure: A qualitative study. <i>Palliative Medicine</i> , 2018, 32, 143-155.	3.1	39
30	Outcomes ‘out of africa’: the selection and implementation of outcome measures for palliative care in Africa. <i>BMC Palliative Care</i> , 2012, 11, 1.	1.8	38
31	Coordination of end-of-life care for patients with lung cancer and those with advanced COPD: are there transferable lessons? A longitudinal qualitative study. <i>Primary Care Respiratory Journal: Journal of the General Practice Airways Group</i> , 2014, 23, 46-51.	2.3	38
32	If you had less than a year to live, would you want to know? A seven-country European population survey of public preferences for disclosure of poor prognosis. <i>Psycho-Oncology</i> , 2013, 22, 2298-2305.	2.3	37
33	The Real-World Problem of Care Coordination: A Longitudinal Qualitative Study with Patients Living with Advanced Progressive Illness and Their Unpaid Caregivers. <i>PLoS ONE</i> , 2014, 9, e95523.	2.5	37
34	‘Burden to others’ as a public concern in advanced cancer: a comparative survey in seven European countries. <i>BMC Cancer</i> , 2013, 13, 105.	2.6	35
35	Results of a transparent expert consultation on patient and public involvement in palliative care research. <i>Palliative Medicine</i> , 2015, 29, 939-949.	3.1	35
36	Measures to assess commonly experienced symptoms for people with dementia in long-term care settings: a systematic review. <i>BMC Medicine</i> , 2016, 14, 38.	5.5	32

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37	Cultural Competence in End-of-Life Care: Terms, Definitions, and Conceptual Models from the British Literature. <i>Journal of Palliative Medicine</i> , 2012, 15, 812-820.	1.1	31
38	Diversity in Defining End of Life Care: An Obstacle or the Way Forward?. <i>PLoS ONE</i> , 2013, 8, e68002.	2.5	30
39	A descriptive study exploring the role of music therapy in prisons. <i>Arts in Psychotherapy</i> , 2001, 28, 137-141.	1.2	29
40	Enhancing patient-reported outcome measurement in research and practice of palliative and end-of-life care. <i>Supportive Care in Cancer</i> , 2012, 20, 1573-1578.	2.2	29
41	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. <i>Annals of Palliative Medicine</i> , 2018, 7, S137-S150.	1.2	28
42	Are we heading in the same direction? European and African doctors' and nurses' views and experiences regarding outcome measurement in palliative care. <i>Palliative Medicine</i> , 2012, 26, 242-249.	3.1	27
43	Culture Is a Priority for Research in End-of-Life Care in Europe: A Research Agenda. <i>Journal of Pain and Symptom Management</i> , 2012, 44, 285-294.	1.2	27
44	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. <i>Journal of Music Therapy</i> , 2016, 53, 1-26.	0.9	27
45	Choosing care homes as the least preferred place to die: a cross-national survey of public preferences in seven European countries. <i>BMC Palliative Care</i> , 2014, 13, 48.	1.8	26
46	The PCOC Symptom Assessment Scale (SAS): A valid measure for daily use at point of care and in palliative care programs. <i>PLoS ONE</i> , 2021, 16, e0247250.	2.5	24
47	"A quiet still voice that just touches" music's relevance for adults living with life-threatening cancer diagnoses. <i>Supportive Care in Cancer</i> , 2014, 22, 1037-1047.	2.2	23
48	A pan-European survey of research in end-of-life cancer care. <i>Supportive Care in Cancer</i> , 2012, 20, 39-48.	2.2	21
49	Emergency department attendance by patients with cancer in the last month of life: a systematic review and meta-analysis. <i>Lancet, The</i> , 2015, 385, S41.	13.7	19
50	A Project Investigating Music Therapy Referral Trends within Palliative Care: An Australian Perspective. <i>Journal of Music Therapy</i> , 2007, 44, 139-155.	0.9	18
51	Constructing Understandings of End-of-Life Care in Europe: A Qualitative Study Involving Cognitive Interviewing with Implications for Cross-National Surveys. <i>Journal of Palliative Medicine</i> , 2011, 14, 343-349.	1.1	18
52	Learning from the public: citizens describe the need to improve end-of-life care access, provision and recognition across Europe. <i>European Journal of Public Health</i> , 2014, 24, 521-527.	0.3	18
53	Which questions of two commonly used multidimensional palliative care patient reported outcome measures are most useful? Results from the European and African PRISMA survey. <i>BMJ Supportive and Palliative Care</i> , 2012, 2, 36-42.	1.6	17
54	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 3 months of life (IARE I study). <i>BMC Medicine</i> , 2020, 18, 344.	5.5	17

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55	Music therapy student supervision: considering aspects of resistance and parallel processes in the supervisory relationship with students in final clinical placement. <i>Arts in Psychotherapy</i> , 2004, 31, 67-76.	1.2	16
56	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. <i>Postgraduate Medical Journal</i> , 2012, 88, 451-457.	1.8	15
57	Patient-reported outcome measurement of symptom distress is feasible in most clinical scenarios in palliative care: an observational study involving routinely collected data. <i>International Journal for Quality in Health Care</i> , 2021, 33, .	1.8	15
58	Social-cultural factors in end-of-life care in Belgium: A scoping of the research literature. <i>Palliative Medicine</i> , 2013, 27, 131-143.	3.1	14
59	Indigenous music therapy theory building through grounded theory research: The developing indigenous theory framework. <i>Arts in Psychotherapy</i> , 2008, 35, 280-286.	1.2	13
60	The PRISMA Symposium 4: How Should Europe Progress End-of-Life and Palliative Clinical Care Research? Recommendations From the Proceedings. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 511-516.	1.2	13
61	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. <i>Age and Ageing</i> , 2017, 46, 300-309.	1.6	12
62	Optimal Cancer Care for Aboriginal and Torres Strait Islander People: A Shared Approach to System Level Change. <i>JCO Global Oncology</i> , 2020, 6, 108-114.	1.8	12
63	A Role for Music Therapy in Special Education. <i>International Journal of Disability Development and Education</i> , 1998, 45, 449-457.	1.1	11
64	An Audit about Music Therapy Assessments and Recommendations for Adult Patients Suspected to be in a Low, Awareness State. <i>Journal of Music Therapy</i> , 2010, 47, 408-422.	0.9	11
65	Of apples and oranges: Lessons learned from the preparation of research protocols for systematic reviews exploring the effectiveness of Specialist Palliative Care. <i>BMC Palliative Care</i> , 2016, 15, 43.	1.8	11
66	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. <i>Palliative Medicine</i> , 2020, 34, 513-523.	3.1	11
67	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. <i>Age and Ageing</i> , 2019, 48, 680-687.	1.6	10
68	The music therapy assessment tool for low awareness states. <i>International Journal of Therapy and Rehabilitation</i> , 2007, 14, 544-549.	0.3	9
69	Avoid "prognostic paralysis" just get ahead and plan and co-ordinate care. <i>Npj Primary Care Respiratory Medicine</i> , 2014, 24, 14085.	2.6	9
70	Effects of professional music therapy supervision on clinical outcomes and therapist competency: a systematic review involving narrative synthesis. <i>Nordic Journal of Music Therapy</i> , 2016, 25, 185-208.	1.1	9
71	Music Therapy Methods with Children, Adolescents, and Adults with Severe Neurobehavioral Disorders Due to Brain Injury. <i>Music Therapy Perspectives</i> , 2011, 29, 5-13.	0.5	7
72	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. <i>Palliative Medicine</i> , 2018, 32, 517-524.	3.1	7

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73	Investigating the Dimension of Time: Findings from a Modified Grounded Theory Study about Clients' Experiences and Descriptions of Temporality or Time within Music Therapy. <i>Journal of Music Therapy</i> , 2011, 48, 28-54.	0.9	6
74	A philosophical inquiry into the role of rhythm in music therapy. <i>Arts in Psychotherapy</i> , 2002, 29, 265-270.	1.2	5
75	Sleeping-related distress in a palliative care population: A national, prospective, consecutive cohort. <i>Palliative Medicine</i> , 2021, 35, 1663-1670.	3.1	5
76	Public opinion on preferences and priorities for end-of-life care in sub-Saharan Africa: piloting a novel method of street surveying: Table 1. <i>BMJ Supportive and Palliative Care</i> , 2012, 2, 72-74.	1.6	3
77	“I’d like to be in a safe place” a qualitative study of the decisions taken by people with advanced cancer to seek emergency department care. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 394.2-394.	1.6	3
78	Charting the Terrain of Grounded Theory Research in Music Therapy. , 2015, , .		2
79	“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. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 388.1-388.	1.6	2
80	Preferences of Older People With a Life-Limiting Illness: A Discrete Choice Experiment. <i>Journal of Pain and Symptom Management</i> , 2022, 64, 137-145.	1.2	2
81	Reflecting on the Long-term (Neurological) End-of-Life Conference. <i>British Journal of Neuroscience Nursing</i> , 2010, 6, 188-190.	0.2	1
82	Professional Supervision as Storied Experience: Narrative Analysis Findings for Australian-Based Registered Music Therapists. <i>Journal of Music Therapy</i> , 2017, 54, 80-107.	0.9	1
83	Appetite-Related Distress Is Burdensome in the Last Sixty Days of Life of People Receiving Palliative Care: A National Longitudinal Consecutive Cohort Study. <i>Journal of Palliative Medicine</i> , 2021, , .	1.1	1
84	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. <i>Journal of Pain and Symptom Management</i> , 2016, 52, e5.	1.2	0