

# Irene J Higginson

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4161175/publications.pdf>

Version: 2024-02-01

669  
papers

39,680  
citations

3449

93  
h-index

7427

157  
g-index

706  
all docs

706  
docs citations

706  
times ranked

24011  
citing authors

#	ARTICLE	IF	CITATIONS
1	Effectiveness and cost-effectiveness of home palliative care services for adults with advanced illness and their caregivers. The Cochrane Library, 2022, 2022, CD007760.	1.5	376
2	Trajectories of disability in activities of daily living in advanced cancer or respiratory disease: a systematic review. Disability and Rehabilitation, 2022, 44, 1790-1801.	0.9	9
3	Intelligent Palliative Care Based on Patient-Reported Outcome Measures. Journal of Pain and Symptom Management, 2022, , .	0.6	5
4	Do guidelines influence breathlessness management in advanced lung diseases? A multinational survey of respiratory medicine and palliative care physicians. BMC Pulmonary Medicine, 2022, 22, 41.	0.8	3
5	Implementation of the Mental Capacity Act: a national observational study comparing resultant trends in place of death for older heart failure decedents with or without comorbid dementia. BMC Medicine, 2022, 20, 30.	2.3	1
6	“You can do it yourself and you can do it at your convenience” internet accessibility and willingness of people with chronic breathlessness to use an internet-based breathlessness self-management intervention during the COVID-19 pandemic. ERJ Open Research, 2022, 8, 00557-2021.	1.1	5
7	Experiences of staff providing specialist palliative care during COVID-19: a multiple qualitative case study. Journal of the Royal Society of Medicine, 2022, 115, 220-230.	1.1	22
8	Change in Activity of Palliative Care Services during the Covid-19 Pandemic: A Multinational Survey (CovPall). Journal of Palliative Medicine, 2022, 25, 465-471.	0.6	16
9	Association of primary and community care services with emergency visits and hospital admissions at the end of life in people with cancer: a retrospective cohort study. BMJ Open, 2022, 12, e054281.	0.8	2
10	COVID-19: Impact on Pediatric Palliative Care. Journal of Pain and Symptom Management, 2022, 64, e1-e5.	0.6	5
11	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. Palliative Medicine, 2022, 36, 319-331.	1.3	14
12	Primary care contacts, continuity, identification of palliative care needs, and hospital use: a population-based cohort study in people dying with dementia. British Journal of General Practice, 2022, 72, e684-e692.	0.7	9
13	Preferences of Older People With a Life-Limiting Illness: A Discrete Choice Experiment. Journal of Pain and Symptom Management, 2022, 64, 137-145.	0.6	2
14	Integrating palliative care into the ICU: a lasting and developing legacy. Intensive Care Medicine, 2022, 48, 939-942.	3.9	10
15	“The whole of humanity has lungs, doesn't it? We are not all the same sort of people” patient preferences and choices for an online, self-guided chronic breathlessness supportive intervention: SELF-BREATHE. ERJ Open Research, 2022, 8, 00093-2022.	1.1	2
16	Face and content validity, acceptability, feasibility, and implementability of a novel outcome measure for children with life-limiting or life-threatening illness in three sub-Saharan African countries. Palliative Medicine, 2022, 36, 1140-1153.	1.3	5
17	Symptom Control and Survival for People Severely ill With COVID: A Multicentre Cohort Study (CovPall-Symptom). Journal of Pain and Symptom Management, 2022, 64, 377-390.	0.6	3
18	Formal and Informal Costs of Care for People With Dementia Who Experience a Transition to Hospital at the End of Life: A Secondary Data Analysis. Journal of the American Medical Directors Association, 2022, , .	1.2	2

#	ARTICLE	IF	CITATIONS
19	Early specialised palliative care: interventions, symptoms, problems. <i>BMJ Supportive and Palliative Care</i> , 2021, 11, 444-453.	0.8	1
20	Disability in Basic Activities of Daily Living Is Associated With Symptom Burden in Older People With Advanced Cancer or Chronic Obstructive Pulmonary Disease: A Secondary Data Analysis. <i>Journal of Pain and Symptom Management</i> , 2021, 61, 1205-1214.	0.6	11
21	Exploring costs, cost components, and associated factors among people with dementia approaching the end of life: A systematic review. <i>Alzheimer's and Dementia: Translational Research and Clinical Interventions</i> , 2021, 7, e12198.	1.8	9
22	Preparedness and capacity of Indian palliative care services to respond to the COVID-19 pandemic: An online rapid assessment survey. <i>Indian Journal of Palliative Care</i> , 2021, 27, 152.	1.0	5
23	Characteristics and mortality rates among patients requiring intermediate care: a national cohort study using linked databases. <i>BMC Medicine</i> , 2021, 19, 48.	2.3	4
24	How many people will need palliative care in Scotland by 2040? A mixed-method study of projected palliative care need and recommendations for service delivery. <i>BMJ Open</i> , 2021, 11, e041317.	0.8	21
25	"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.	1.3	67
26	The impact of and response to the COVID-19 pandemic on a hospital palliative care team. <i>Future Healthcare Journal</i> , 2021, 8, 62-64.	0.6	5
27	Preparedness to Face the COVID-19 Pandemic in Hospice and Palliative Care Services in the Asia-Pacific Region: A Rapid Online Survey. <i>American Journal of Hospice and Palliative Medicine</i> , 2021, 38, 861-868.	0.8	7
28	Identification of palliative care needs among people with dementia and its association with acute hospital care and community service use at the end-of-life: A retrospective cohort study using linked primary, community and secondary care data. <i>Palliative Medicine</i> , 2021, 35, 1691-1700.	1.3	9
29	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.	1.3	34
30	Hospital-based specialist palliative care compared with usual care for adults with advanced illness and their caregivers: a systematic review. <i>Health Services and Delivery Research</i> , 2021, 9, 1-218.	1.4	2
31	Dying at home during the pandemic. <i>BMJ</i> , 2021, 373, n1437.	3.0	12
32	Community-based short-term integrated palliative and supportive care reduces symptom distress for older people with chronic noncancer conditions compared with usual care: A randomised controlled single-blind mixed method trial. <i>International Journal of Nursing Studies</i> , 2021, 120, 103978.	2.5	18
33	Changes in mortality patterns and place of death during the COVID-19 pandemic: A descriptive analysis of mortality data across four nations. <i>Palliative Medicine</i> , 2021, 35, 1975-1984.	1.3	38
34	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.	0.8	21
35	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, , .	0.5	12
36	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.	0.6	57

#	ARTICLE	IF	CITATIONS
37	Relationships between prolonged physical and social isolation during the COVID-19 pandemic, reduced physical activity and disability in activities of daily living among people with advanced respiratory disease. <i>Chronic Respiratory Disease</i> , 2021, 18, 147997312110358.	1.0	18
38	Breathlessness. <i>Respiratory Medicine</i> , 2021, , 89-113.	0.1	0
39	Pharmacological and nonpharmacological interventions to improve symptom control, functional exercise capacity and quality of life in interstitial lung disease: an evidence synthesis. <i>ERJ Open Research</i> , 2021, 7, 00107-2020.	1.1	4
40	Managing clinical uncertainty in older people towards the end of life: a systematic review of person-centred tools. <i>BMC Palliative Care</i> , 2021, 20, 168.	0.8	7
41	Preferences and priorities to manage clinical uncertainty for older people with frailty and multimorbidity: a discrete choice experiment and stakeholder consultations. <i>BMC Geriatrics</i> , 2021, 21, 553.	1.1	4
42	O-2â€œ...â€œGoing against the grain of all we doâ€œ™: hospice staff experiences of moral distress during COVID-19. , 2021, , .		0
43	Response to Zhou (2021) â€œComment on Evans et al (2021) â€œCommunity-based short-term integrated palliative and supportive care reduces symptom distress for older people with chronic noncancer conditions compared with usual careâ€œ™â€œ. <i>International Journal of Nursing Studies</i> , 2021, 125, 104119.	2.5	0
44	Common elements of service delivery models that optimise quality of life and health service use among older people with advanced progressive conditions: a tertiary systematic review. <i>BMJ Open</i> , 2021, 11, e048417.	0.8	5
45	Electronic palliative care coordination systems (EPaCCS): a systematic review. <i>BMJ Supportive and Palliative Care</i> , 2020, 10, 68-78.	0.8	31
46	End-of-life care and place of death in adults with serious mental illness: A systematic review and narrative synthesis. <i>Palliative Medicine</i> , 2020, 34, 49-68.	1.3	24
47	Exploratory analyses of the Danish Palliative Care Trial (DanPaCT): a randomized trial of early specialized palliative care plus standard care versus standard care in advanced cancer patients. <i>Supportive Care in Cancer</i> , 2020, 28, 2145-2155.	1.0	13
48	Intention-to-Treat Analyses for Randomized Controlled Trials in Hospice/Palliative Care: The Case for Analyses to be of People Exposed to the Intervention. <i>Journal of Pain and Symptom Management</i> , 2020, 59, 637-645.	0.6	9
49	Short-term integrated rehabilitation for people with newly diagnosed thoracic cancer: a multi-centre randomized controlled feasibility trial. <i>Clinical Rehabilitation</i> , 2020, 34, 205-219.	1.0	5
50	The effectiveness and cost-effectiveness of hospital-based specialist palliative care for adults with advanced illness and their caregivers. <i>The Cochrane Library</i> , 2020, 2020, CD012780.	1.5	61
51	Palliative care for patients with severe covid-19. <i>BMJ, The</i> , 2020, 370, m2710.	3.0	44
52	Intensive care utilization in patients with end-stage liver disease: A population-based comparative study of cohorts with and without comorbid hepatocellular carcinoma in Taiwan. <i>EClinicalMedicine</i> , 2020, 22, 100357.	3.2	0
53	Preparedness of African Palliative Care Services to Respond to the COVID-19 Pandemic: A Rapid Assessment. <i>Journal of Pain and Symptom Management</i> , 2020, 60, e10-e26.	0.6	12
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 3Âˆmonths of life (IARE I study). <i>BMC Medicine</i> , 2020, 18, 344.	2.3	17

#	ARTICLE	IF	CITATIONS
55	Changing patterns of mortality during the COVID-19 pandemic: Population-based modelling to understand palliative care implications. <i>Palliative Medicine</i> , 2020, 34, 1193-1201.	1.3	49
56	Palliative Care in Acute Heart Failure. <i>Current Heart Failure Reports</i> , 2020, 17, 424-437.	1.3	6
57	Effect of Short-term Integrated Palliative Care on Patient-Reported Outcomes Among Patients Severely Affected With Long-term Neurological Conditions. <i>JAMA Network Open</i> , 2020, 3, e2015061.	2.8	26
58	Determining the prevalence of palliative needs and exploring screening accuracy of depression and anxiety items of the integrated palliative care outcome scale – a multi-centre study. <i>BMC Palliative Care</i> , 2020, 19, 69.	0.8	7
59	Phase II mixed methods™ feasibility cluster randomised controlled trial of a novel community-based enhanced care intervention to improve person-centred outcomes for people living with HIV in Ghana. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 2020, 32, 107-118.	0.6	5
60	Towards person-centred quality care for children with life-limiting and life-threatening illness: Self-reported symptoms, concerns and priority outcomes from a multi-country qualitative study. <i>Palliative Medicine</i> , 2020, 34, 319-335.	1.3	18
61	Standards, Guidelines, and Quality Measures for Successful Specialty Palliative Care Integration Into Oncology: Current Approaches and Future Directions. <i>Journal of Clinical Oncology</i> , 2020, 38, 987-994.	0.8	30
62	Palliative Care and the Management of Common Distressing Symptoms in Advanced Cancer: Pain, Breathlessness, Nausea and Vomiting, and Fatigue. <i>Journal of Clinical Oncology</i> , 2020, 38, 905-914.	0.8	154
63	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.	1.3	11
64	Equal or not? Women hold less prestigious roles at respiratory medicine conferences than men. <i>European Respiratory Journal</i> , 2020, 55, 1900701.	3.1	15
65	Response and role of palliative care during the COVID-19 pandemic: A national telephone survey of hospices in Italy. <i>Palliative Medicine</i> , 2020, 34, 889-895.	1.3	108
66	Primary care service use by end-of-life cancer patients: a nationwide population-based cohort study in the United Kingdom. <i>BMC Family Practice</i> , 2020, 21, 76.	2.9	11
67	The Role and Response of Palliative Care and Hospice Services in Epidemics and Pandemics: A Rapid Review to Inform Practice During the COVID-19 Pandemic. <i>Journal of Pain and Symptom Management</i> , 2020, 60, e31-e40.	0.6	204
68	Characteristics, Symptom Management, and Outcomes of 101 Patients With COVID-19 Referred for Hospital Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2020, 60, e77-e81.	0.6	160
69	Regional variations in geographic access to inpatient hospices and Place of death: A Population-based study in England, UK. <i>PLoS ONE</i> , 2020, 15, e0231666.	1.1	11
70	Managing the supportive care needs of those affected by COVID-19. <i>European Respiratory Journal</i> , 2020, 55, 2000815.	3.1	95
71	What is the evidence for mirtazapine in treating cancer-related symptomatology? A systematic review. <i>Supportive Care in Cancer</i> , 2020, 28, 1597-1606.	1.0	19
72	How can we achieve person-centred care for people living with HIV/AIDS? A qualitative interview study with healthcare professionals and patients in Ghana. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 2020, 32, 1479-1488.	0.6	12

#	ARTICLE	IF	CITATIONS
73	Towards patient-centred cancer care: cross-cultural validity and responsiveness of the Turkish Integrated Palliative care Outcome Scale. <i>Health and Quality of Life Outcomes</i> , 2020, 18, 312.	1.0	11
74	Immediate versus delayed short-term integrated palliative care for advanced long-term neurological conditions: the OPTCARE Neuro RCT. <i>Health Services and Delivery Research</i> , 2020, 8, 1-80.	1.4	4
75	Randomised, double-blind, multicentre, mixed-methods, dose-escalation feasibility trial of mirtazapine for better treatment of severe breathlessness in advanced lung disease (BETTER-B feasibility). <i>Thorax</i> , 2020, 75, 176-179.	2.7	4
76	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.	0.7	10
77	Establishing key criteria to define and compare models of specialist palliative care: A mixed-methods study using qualitative interviews and Delphi survey. <i>Palliative Medicine</i> , 2019, 33, 1114-1124.	1.3	27
78	A Rasch Analysis of the Integrated Palliative Care Outcome Scale. <i>Journal of Pain and Symptom Management</i> , 2019, 57, 290-296.	0.6	27
79	Development and Evaluation of Complex Interventions in Palliative Care. , 2019, , 1669-1679.		1
80	Understanding which people with dementia are at risk of inappropriate care and avoidable transitions to hospital near the end-of-life: a retrospective cohort study. <i>Age and Ageing</i> , 2019, 48, 672-679.	0.7	25
81	Which outcome domains are important in palliative care and when? An international expert consensus workshop, using the nominal group technique. <i>Palliative Medicine</i> , 2019, 33, 1058-1068.	1.3	27
82	The escalating global burden of serious health-related suffering: projections to 2060 by world regions, age groups, and health conditions. <i>The Lancet Global Health</i> , 2019, 7, e883-e892.	2.9	395
83	A brief, patient- and proxy-reported outcome measure in advanced illness: Validity, reliability and responsiveness of the Integrated Palliative care Outcome Scale (IPOS). <i>Palliative Medicine</i> , 2019, 33, 1045-1057.	1.3	197
84	Place of death and other factors associated with unnatural mortality in patients with serious mental disorders: population-based retrospective cohort study. <i>BJPsych Open</i> , 2019, 5, e23.	0.3	8
85	Symptoms and anxiety predict declining health-related quality of life in multiple myeloma: A prospective, multi-centre longitudinal study. <i>Palliative Medicine</i> , 2019, 33, 541-551.	1.3	30
86	Urban and rural differences in geographical accessibility to inpatient palliative and end-of-life (PEoLC) facilities and place of death: a national population-based study in England, UK. <i>International Journal of Health Geographics</i> , 2019, 18, 8.	1.2	36
87	Socioeconomic position and use of healthcare in the last year of life: A systematic review and meta-analysis. <i>PLoS Medicine</i> , 2019, 16, e1002782.	3.9	102
88	Service Delivery Models to Maximize Quality of Life for Older People at the End of Life: A Rapid Review. <i>Milbank Quarterly</i> , 2019, 97, 113-175.	2.1	55
89	Change in multidimensional problems and quality of life over three months after HIV diagnosis: a multicentre longitudinal study in Kenya and Uganda. <i>BMC Infectious Diseases</i> , 2019, 19, 248.	1.3	1
90	Symptom dimensions in people affected by long-term neurological conditions: a factor analysis of a patient-centred palliative care outcome symptom scale. <i>Scientific Reports</i> , 2019, 9, 4972.	1.6	8

#	ARTICLE	IF	CITATIONS
91	The views and experiences of older people with conservatively managed renal failure: a qualitative study of communication, information and decision-making. <i>BMC Nephrology</i> , 2019, 20, 38.	0.8	19
92	Recommendations for services for people living with chronic breathlessness in advanced disease: Results of a transparent expert consultation. <i>Chronic Respiratory Disease</i> , 2019, 16, 147997311881644.	1.0	13
93	In response to Ballantyne and Schaefer's "Consent and the ethical duty to participate in health data research". <i>Journal of Medical Ethics</i> , 2019, 45, 351-352.	1.0	5
94	The King's Brief Interstitial Lung Disease (KBILD) questionnaire: an updated minimal clinically important difference. <i>BMJ Open Respiratory Research</i> , 2019, 6, e000363.	1.2	30
95	Invisible and intangible illness: a qualitative interview study of patients' experiences and understandings of conservatively managed end-stage kidney disease. <i>Annals of Palliative Medicine</i> , 2019, 8, 121-129.	0.5	16
96	Predicting outcomes following holistic breathlessness services: A pooled analysis of individual patient data. <i>Palliative Medicine</i> , 2019, 33, 462-466.	1.3	8
97	The impact of population ageing on end-of-life care in Scotland: projections of place of death and recommendations for future service provision. <i>BMC Palliative Care</i> , 2019, 18, 112.	0.8	35
98	Place of death and factors associated with hospital death in patients who have died from liver disease in England: a national population-based study. <i>The Lancet Gastroenterology and Hepatology</i> , 2019, 4, 52-62.	3.7	14
99	Finding a "new normal" following acute illness: A qualitative study of influences on frail older people's care preferences. <i>Palliative Medicine</i> , 2019, 33, 301-311.	1.3	21
100	Translation and cognitive testing of the Italian Integrated Palliative Outcome Scale (IPOS) among patients and healthcare professionals. <i>PLoS ONE</i> , 2019, 14, e0208536.	1.1	19
101	Symptom prevalence and quality of life of patients with end-stage liver disease: A systematic review and meta-analysis. <i>Palliative Medicine</i> , 2019, 33, 24-36.	1.3	157
102	Symptoms and Concerns Among Children and Young People with Life-Limiting and Life-Threatening Conditions: A Systematic Review Highlighting Meaningful Health Outcomes. <i>Patient</i> , 2019, 12, 15-55.	1.1	50
103	Holistic services for people with advanced disease and chronic breathlessness: a systematic review and meta-analysis. <i>Thorax</i> , 2019, 74, 270-281.	2.7	96
104	Factors associated with older people's emergency department attendance towards the end of life: a systematic review. <i>European Journal of Public Health</i> , 2019, 29, 67-74.	0.1	35
105	Control and Context Are Central for People With Advanced Illness Experiencing Breathlessness: A Systematic Review and Thematic Synthesis. <i>Journal of Pain and Symptom Management</i> , 2019, 57, 140-155.e2.	0.6	26
106	"Keep All Thee 'Til the End": Reclaiming the Lifeworld for Patients in the Hospice Setting. <i>Omega: Journal of Death and Dying</i> , 2019, 78, 390-403.	0.7	7
107	Leaky pipeline, gender bias, self-selection or all three? A quantitative analysis of gender balance at an international palliative care research conference. <i>BMJ Supportive and Palliative Care</i> , 2019, 9, 146-148.	0.8	27
108	The role of service factors on variations in place of death: an observational study. <i>Health Services and Delivery Research</i> , 2019, 7, 1-58.	1.4	2

#	ARTICLE	IF	CITATIONS
109	Holistic services for people with advanced disease and chronic or refractory breathlessness: a mixed-methods evidence synthesis. <i>Health Services and Delivery Research</i> , 2019, 7, 1-104.	1.4	8
110	The AMBER care bundle for hospital inpatients with uncertain recovery nearing the end of life: the ImproveCare feasibility cluster RCT. <i>Health Technology Assessment</i> , 2019, 23, 1-150.	1.3	11
111	Population-based palliative care planning in Ireland: how many people will live and die with serious illness to 2046?. <i>HRB Open Research</i> , 2019, 2, 35.	0.3	14
112	A framework for complexity in palliative care: A qualitative study with patients, family carers and professionals. <i>Palliative Medicine</i> , 2018, 32, 1078-1090.	1.3	85
113	Influences on Care Preferences of Older People with Advanced Illness: A Systematic Review and Thematic Synthesis. <i>Journal of the American Geriatrics Society</i> , 2018, 66, 1031-1039.	1.3	49
114	Changing health behaviour with rehabilitation in thoracic cancer: A systematic review and synthesis. <i>Psycho-Oncology</i> , 2018, 27, 1675-1694.	1.0	13
115	Effect of participation in a randomised controlled trial of an integrated palliative care intervention on HIV-associated stigma. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 2018, 30, 1180-1188.	0.6	12
116	Development and validation of a casemix classification to predict costs of specialist palliative care provision across inpatient hospice, hospital and community settings in the UK: a study protocol. <i>BMJ Open</i> , 2018, 8, e020071.	0.8	11
117	Palliative care needs of heart failure patients in China: putting people first. <i>Current Opinion in Supportive and Palliative Care</i> , 2018, 12, 10-15.	0.5	6
118	Integrating palliative care into neurology services: what do the professionals say?. <i>BMJ Supportive and Palliative Care</i> , 2018, 8, 41-44.	0.8	24
119	Time to go beyond observing the problem. Response to: Dying in hospital: socioeconomic inequality trends in England, DOI: 10.1177/1355819616686807. <i>Journal of Health Services Research and Policy</i> , 2018, , 135581961775018.	0.8	0
120	Implementing Outcome Measures in Palliative Care. <i>Journal of Palliative Medicine</i> , 2018, 21, 414-414.	0.6	1
121	What factors influence emergency department visits by patients with cancer at the end of life? Analysis of a 124,030 patient cohort. <i>Palliative Medicine</i> , 2018, 32, 426-438.	1.3	41
122	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.	1.3	7
123	Seeing is believing – healthcare professionals’ perceptions of a complex intervention to improve care towards the end of life: A qualitative interview study. <i>Palliative Medicine</i> , 2018, 32, 525-532.	1.3	14
124	Strategies used in improving and assessing the level of reporting of implementation fidelity in randomised controlled trials of palliative care complex interventions: A systematic review. <i>Palliative Medicine</i> , 2018, 32, 500-516.	1.3	19
125	Alleviating the access abyss in palliative care and pain relief – an imperative of universal health coverage: the Lancet Commission report. <i>Lancet</i> , The, 2018, 391, 1391-1454.	6.3	732
126	What is the impact of population ageing on the future provision of end-of-life care? Population-based projections of place of death. <i>Palliative Medicine</i> , 2018, 32, 329-336.	1.3	218



#	ARTICLE	IF	CITATIONS
127	Predictors of emergency department attendance by people with dementia in their last year of life: Retrospective cohort study using linked clinical and administrative data. <i>Alzheimer's and Dementia</i> , 2018, 14, 20-27.	0.4	46
128	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.	1.3	39
129	Development and Evaluation of Complex Interventions in Palliative Care. , 2018, , 1-11.		0
130	Outcome measurement—a scoping review of the literature and future developments in palliative care clinical practice. <i>Annals of Palliative Medicine</i> , 2018, 7, S196-S206.	0.5	19
131	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.	0.5	28
132	Developing an integrated rehabilitation model for thoracic cancer services: views of patients, informal carers and clinicians. <i>Pilot and Feasibility Studies</i> , 2018, 4, 160.	0.5	10
133	Integration of oncology and palliative care: a Lancet Oncology Commission. <i>Lancet Oncology</i> , The, 2018, 19, e588-e653.	5.1	452
134	The future of end-of-life care. <i>Lancet</i> , The, 2018, 392, 915-916.	6.3	10
135	A Population-Based Conceptual Framework for Evaluating the Role of Healthcare Services in Place of Death. <i>Healthcare (Switzerland)</i> , 2018, 6, 107.	1.0	13
136	Patient empowerment, what does it mean for adults in the advanced stages of a life-limiting illness: A systematic review using critical interpretive synthesis. <i>Palliative Medicine</i> , 2018, 32, 1288-1304.	1.3	47
137	Use of mirtazapine in patients with chronic breathlessness: A case series. <i>Palliative Medicine</i> , 2018, 32, 1518-1521.	1.3	20
138	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.	1.1	59
139	Is end-of-life care a priority for policymakers? Qualitative documentary analysis of health care strategies. <i>Palliative Medicine</i> , 2018, 32, 1474-1486.	1.3	20
140	Active ingredients of a person-centred intervention for people on HIV treatment: analysis of mixed methods trial data. <i>BMC Infectious Diseases</i> , 2018, 18, 27.	1.3	12
141	Exercise Deficiency Diseases of Ageing: The Primacy of Exercise and Muscle Strengthening as First-Line Therapeutic Agents to Combat Frailty. <i>Journal of the American Medical Directors Association</i> , 2018, 19, 741-743.	1.2	39
142	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.	0.7	12
143	The intensity of caregiving is a more important predictor of adverse bereavement outcomes for adult child than spousal caregivers of patients who die of cancer. <i>Psycho-Oncology</i> , 2017, 26, 316-322.	1.0	10
144	Measuring geographical accessibility to palliative and end of life (PEoLC) related facilities: a comparative study in an area with well-developed specialist palliative care (SPC) provision. <i>BMC Palliative Care</i> , 2017, 16, 14.	0.8	11

#	ARTICLE	IF	CITATIONS
145	Variations in the cost of formal and informal health care for patients with advanced chronic disease and refractory breathlessness: A cross-sectional secondary analysis. <i>Palliative Medicine</i> , 2017, 31, 369-377.	1.3	38
146	Which patients with advanced respiratory disease die in hospital? A 14-year population-based study of trends and associated factors. <i>BMC Medicine</i> , 2017, 15, 19.	2.3	47
147	The Myeloma Patient Outcome Scale is the first quality of life tool developed for clinical use and validated in patients with follicular lymphoma. <i>European Journal of Haematology</i> , 2017, 98, 508-516.	1.1	7
148	Home palliative care works: but how? A meta-ethnography of the experiences of patients and family caregivers. <i>BMJ Supportive and Palliative Care</i> , 2017, 7, 00.2-00.	0.8	96
149	Randomised clinical trial of early specialist palliative care plus standard care versus standard care alone in patients with advanced cancer: The Danish Palliative Care Trial. <i>Palliative Medicine</i> , 2017, 31, 814-824.	1.3	67
150	Does the EQ-5D capture the concerns measured by the Palliative care Outcome Scale? Mapping the Palliative care Outcome Scale onto the EQ-5D using statistical methods. <i>Palliative Medicine</i> , 2017, 31, 716-725.	1.3	23
151	Breathlessness during daily activity: The psychometric properties of the London Chest Activity of Daily Living Scale in patients with advanced disease and refractory breathlessness. <i>Palliative Medicine</i> , 2017, 31, 868-875.	1.3	11
152	About the "œsurprise question" Cmaj, 2017, 189, E807-E807.	0.9	5
153	The Effectiveness of Singing or Playing a Wind Instrument in Improving Respiratory Function in Patients with Long-Term Neurological Conditions: A Systematic Review. <i>Journal of Music Therapy</i> , 2017, 54, 108-131.	0.6	5
154	Development of a Patient-Reported Palliative Care-Specific Health Classification System: The POS-E. <i>Patient</i> , 2017, 10, 353-365.	1.1	14
155	Do we have adequate tools and skills to manage uncertainty among patients and families in ICU?. <i>Intensive Care Medicine</i> , 2017, 43, 463-464.	3.9	0
156	Universal coverage for palliative care in respiratory disease and critical care. <i>Lancet Respiratory Medicine</i> , 2017, 5, 915-916.	5.2	4
157	Valuing Attributes of Home Palliative Care With Service Users: A Pilot Discrete Choice Experiment. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 973-985.	0.6	11
158	Palliative care and management of troublesome symptoms for people with chronic obstructive pulmonary disease. <i>Lancet, The</i> , 2017, 390, 988-1002.	6.3	147
159	Feasibility and acceptability of a patient-reported outcome intervention in chronic heart failure. <i>BMJ Supportive and Palliative Care</i> , 2017, 7, 470-479.	0.8	42
160	Longitudinal validity and reliability of the Myeloma Patient Outcome Scale (MyPOS) was established using traditional, generalizability and Rasch psychometric methods. <i>Quality of Life Research</i> , 2017, 26, 2931-2947.	1.5	5
161	The Effect of Communication Skills Training for Generalist Palliative Care Providers on Patient-Reported Outcomes and Clinician Behaviors: A Systematic Review and Meta-analysis. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 404-416.e5.	0.6	70
162	A Systematic Review of End-of-Life Care Communication Skills Training for Generalist Palliative Care Providers: Research Quality and Reporting Guidance. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 417-425.	0.6	58

#	ARTICLE	IF	CITATIONS
163	Disability in activities of daily living among adults with cancer: A systematic review and meta-analysis. <i>Cancer Treatment Reviews</i> , 2017, 61, 94-106.	3.4	166
164	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.	1.3	57
165	The effectiveness and cost-effectiveness of inpatient specialist palliative care in acute hospitals for adults with advanced illness and their caregivers. <i>The Cochrane Library</i> , 2017, , .	1.5	4
166	Hospice and Palliative Care. , 2017, , 69-76.		1
167	Cognitive-emotional interventions for breathlessness in adults with advanced diseases. <i>The Cochrane Library</i> , 2017, , .	1.5	0
168	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.	1.1	110
169	Editorial. <i>Palliative Medicine</i> , 2017, 31, 681-683.	1.3	0
170	The TEAM Approach to Improving Oncology Outcomes by Incorporating Palliative Care in Practice. <i>Journal of Oncology Practice</i> , 2017, 13, 557-566.	2.5	40
171	Effectiveness and cost-effectiveness of home palliative care services for adults with advanced illness and their caregivers. <i>Sao Paulo Medical Journal</i> , 2016, 134, 93-94.	0.4	15
172	Psychometric Properties of a Generic, Patient-Centred Palliative Care Outcome Measure of Symptom Burden for People with Progressive Long Term Neurological Conditions. <i>PLoS ONE</i> , 2016, 11, e0165379.	1.1	19
173	The impact of disease-related symptoms and palliative care concerns on health-related quality of life in multiple myeloma: a multi-centre study. <i>BMC Cancer</i> , 2016, 16, 427.	1.1	111
174	“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.	0.8	66
175	“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 Supportive and Palliative Care</i> , 2016, 6, 394.2-394.	0.8	3
176	Evidence-based policy in palliative care: time to learn from our mistakes. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 417-417.	0.8	2
177	“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.	0.8	45
178	Place of death in children and young people with cancer and implications for end of life care: a population-based study in England, 1993-2014. <i>BMC Cancer</i> , 2016, 16, 727.	1.1	31
179	“Can Breathe Again!” Patients' Self-Management Strategies for Episodic Breathlessness in Advanced Disease, Derived From Qualitative Interviews. <i>Journal of Pain and Symptom Management</i> , 2016, 52, 228-234.	0.6	35
180	Past trends and projections of hospital deaths to inform the integration of palliative care in one of the most ageing countries in the world. <i>Palliative Medicine</i> , 2016, 30, 363-373.	1.3	40

#	ARTICLE	IF	CITATIONS
181	How integrated are neurology and palliative care services? Results of a multicentre mapping exercise. <i>BMC Neurology</i> , 2016, 16, 63.	0.8	41
182	Dignity Through Integrated Symptom Management: Lessons From the Breathlessness Support Service. <i>Journal of Pain and Symptom Management</i> , 2016, 52, 515-524.	0.6	18
183	Factors Associated with Transition from Community Settings to Hospital as Place of Death for Adults Aged 75 and Older: A Population-Based Mortality Follow-Back Survey. <i>Journal of the American Geriatrics Society</i> , 2016, 64, 2210-2217.	1.3	34
184	Prevalence of symptoms in patients with multiple myeloma: a systematic review and meta-analysis. <i>European Journal of Haematology</i> , 2016, 97, 416-429.	1.1	104
185	“Outcomes” Is Not an Oxymoron in Hospice/Palliative Care. <i>Journal of Palliative Medicine</i> , 2016, 19, 1128-1129.	0.6	3
186	Neuromuscular electrical stimulation for muscle weakness in adults with advanced disease. <i>The Cochrane Library</i> , 2016, 2016, CD009419.	1.5	131
187	Benzodiazepines for the relief of breathlessness in advanced malignant and non-malignant diseases in adults. <i>The Cochrane Library</i> , 2016, 2016, CD007354.	1.5	133
188	Developing a model of short-term integrated palliative and supportive care for frail older people in community settings: perspectives of older people, carers and other key stakeholders. <i>Age and Ageing</i> , 2016, 45, 863-873.	0.7	70
189	Using routine data to improve palliative and end of life care. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 257-262.	0.8	54
190	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.	0.8	11
191	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.	2.3	32
192	Discovering the hidden benefits of cognitive interviewing in two languages: The first phase of a validation study of the Integrated Palliative care Outcome Scale. <i>Palliative Medicine</i> , 2016, 30, 599-610.	1.3	109
193	Neuromuscular electrical stimulation to improve exercise capacity in patients with severe COPD – Authors’ reply. <i>Lancet Respiratory Medicine</i> , 2016, 4, e16.	5.2	3
194	Validity, reliability and responsiveness to change of the Italian palliative care outcome scale: a multicenter study of advanced cancer patients. <i>BMC Palliative Care</i> , 2016, 15, 23.	0.8	22
195	The clinical and cost effectiveness of a Breathlessness Intervention Service for patients with advanced non-malignant disease and their informal carers: mixed findings of a mixed method randomised controlled trial. <i>Trials</i> , 2016, 17, 185.	0.7	94
196	Reporting of clinically diagnosed dementia on death certificates: retrospective cohort study. <i>Age and Ageing</i> , 2016, 45, 667-672.	0.7	55
197	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.	1.3	149
198	Patients’ experiences of a new integrated breathlessness support service for patients with refractory breathlessness: Results of a postal survey. <i>Palliative Medicine</i> , 2016, 30, 313-322.	1.3	26

#	ARTICLE	IF	CITATIONS
199	Research challenges in palliative and end of life care. <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 2-4.	0.8	59
200	The need to research refractory breathlessness. <i>European Respiratory Journal</i> , 2016, 47, 342-343.	3.1	32
201	Neuromuscular electrical stimulation to improve exercise capacity in patients with severe COPD: a randomised double-blind, placebo-controlled trial. <i>Lancet Respiratory Medicine</i> , 2016, 4, 27-36.	5.2	110
202	“I’ll continue as long as I can, and die when I can’t help it”: a qualitative exploration of the views of end-of-life care by those affected by head and neck cancer (HNC). <i>BMJ Supportive and Palliative Care</i> , 2016, 6, 43-51.	0.8	18
203	Home care by general practitioners for cancer patients in the last 3 months of life: An epidemiological study of quality and associated factors. <i>Palliative Medicine</i> , 2016, 30, 64-74.	1.3	35
204	The changing demographics of inpatient hospice death: Population-based cross-sectional study in England, 1993–2012. <i>Palliative Medicine</i> , 2016, 30, 45-53.	1.3	54
205	Decision Making About Gastrostomy and Noninvasive Ventilation in Amyotrophic Lateral Sclerosis. <i>Qualitative Health Research</i> , 2016, 26, 1366-1381.	1.0	36
206	Factors associated with aggressive end of life cancer care. <i>Supportive Care in Cancer</i> , 2016, 24, 1079-1089.	1.0	64
207	Factors Associated with Participation, Active Refusals and Reasons for Not Taking Part in a Mortality Followback Survey Evaluating End-of-Life Care. <i>PLoS ONE</i> , 2016, 11, e0146134.	1.1	13
208	How should we manage information needs, family anxiety, depression, and breathlessness for those affected by advanced disease: development of a Clinical Decision Support Tool using a Delphi design. <i>BMC Medicine</i> , 2015, 13, 263.	2.3	34
209	Is dying in hospital better than home in incurable cancer and what factors influence this? A population-based study. <i>BMC Medicine</i> , 2015, 13, 235.	2.3	111
210	How does a new breathlessness support service affect patients?. <i>European Respiratory Journal</i> , 2015, 46, 1515-1518.	3.1	17
211	Cochrane Reviews: Four Proposals for Improvement. <i>Journal of Palliative Medicine</i> , 2015, 18, 906-907.	0.6	3
212	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.	0.6	158
213	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.	0.8	99
214	Results of a transparent expert consultation on patient and public involvement in palliative care research. <i>Palliative Medicine</i> , 2015, 29, 939-949.	1.3	35
215	Invisible suffering: breathlessness in and beyond the clinic—a reply. <i>Lancet Respiratory Medicine</i> , 2015, 3, e29.	5.2	1
216	Rasch analysis and impact factor methods both yield valid and comparable measures of health status in interstitial lung disease. <i>Journal of Clinical Epidemiology</i> , 2015, 68, 1019-1027.	2.4	10

#	ARTICLE	IF	CITATIONS
217	Screening for Depression in Advanced Disease: Psychometric Properties, Sensitivity, and Specificity of Two Items of the Palliative Care Outcome Scale (POS). <i>Journal of Pain and Symptom Management</i> , 2015, 49, 277-288.	0.6	14
218	Bioelectrical impedance phase angle relates to function, disease severity and prognosis in stable chronic obstructive pulmonary disease. <i>Clinical Nutrition</i> , 2015, 34, 1245-1250.	2.3	75
219	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.	0.6	54
220	Nurse-led palliative care for HIV-positive patients taking antiretroviral therapy in Kenya: a randomised controlled trial. <i>Lancet HIV</i> , 2015, 2, e328-e334.	2.1	45
221	Comparative Analysis of Informal Caregiver Burden in Advanced Cancer, Dementia, and Acquired Brain Injury. <i>Journal of Pain and Symptom Management</i> , 2015, 50, 445-452.	0.6	79
222	Palliative care for patients with advanced fibrotic lung disease: a randomised controlled phase II and feasibility trial of a community case conference intervention. <i>Thorax</i> , 2015, 70, 830-839.	2.7	97
223	“Less ticking the boxes, more providing support”: A qualitative study on health professionals’ concerns towards the Liverpool Care of the Dying Pathway. <i>Palliative Medicine</i> , 2015, 29, 529-537.	1.3	10
224	Characteristics of episodic breathlessness as reported by patients with advanced chronic obstructive pulmonary disease and lung cancer: Results of a descriptive cohort study. <i>Palliative Medicine</i> , 2015, 29, 420-428.	1.3	33
225	Public Health and Palliative Care in 2015. <i>Clinics in Geriatric Medicine</i> , 2015, 31, 253-263.	1.0	18
226	Twitter discussions from a respirology journal club “ Authors’ reply. <i>Lancet Respiratory Medicine</i> , 2015, 3, e10-e11.	5.2	0
227	Improving the assessment of quality of life in the clinical care of myeloma patients: the development and validation of the Myeloma Patient Outcome Scale (MyPOS). <i>BMC Cancer</i> , 2015, 15, 280.	1.1	37
228	Patient and carer experiences of clinical uncertainty and deterioration, in the face of limited reversibility: A comparative observational study of the AMBER care bundle. <i>Palliative Medicine</i> , 2015, 29, 797-807.	1.3	36
229	Breathlessness and crises in the context of advanced illness: A comparison between COPD and lung cancer patients. <i>Palliative and Supportive Care</i> , 2015, 13, 229-237.	0.6	18
230	“It doesn't do the care for you’: a qualitative study of health care professionals’ perceptions of the benefits and harms of integrated care pathways for end of life care: Figure 1. <i>BMJ Open</i> , 2015, 5, e008242.	0.8	28
231	Exploring meanings of illness causation among those severely affected by multiple sclerosis: a comparative qualitative study of Black Caribbean and White British people. <i>BMC Palliative Care</i> , 2015, 14, 13.	0.8	12
232	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.	0.6	75
233	Managing Cancer Pain at the End of Life with Multiple Strong Opioids: A Population-Based Retrospective Cohort Study in Primary Care. <i>PLoS ONE</i> , 2014, 9, e79266.	1.1	18
234	Does Ethnicity Affect Where People with Cancer Die? A Population-Based 10 Year Study. <i>PLoS ONE</i> , 2014, 9, e95052.	1.1	32

#	ARTICLE	IF	CITATIONS
235	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.	1.1	37
236	What Next in Refractory Breathlessness? Breathlessness? Research Questions for Palliative Care. <i>Journal of Palliative Care</i> , 2014, 30, 271-278.	0.4	3
237	The HIV basic care package: where is it available and who receives it? Findings from a mixed methods evaluation in Kenya and Uganda. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 2014, 26, 1155-1163.	0.6	2
238	A randomised controlled trial to assess the effectiveness of a nurse-led palliative care intervention for HIV positive patients on antiretroviral therapy: recruitment, refusal, randomisation and missing data. <i>BMC Research Notes</i> , 2014, 7, 600.	0.6	9
239	Detailed statistical analysis plan for the Danish Palliative Care Trial (DanPaCT). <i>Trials</i> , 2014, 15, 376.	0.7	18
240	Benefits and Costs of Home Palliative Care Compared With Usual Care for Patients With Advanced Illness and Their Family Caregivers. <i>JAMA - Journal of the American Medical Association</i> , 2014, 311, 1060.	3.8	53
241	Place and Cause of Death in Centenarians: A Population-Based Observational Study in England, 2001 to 2010. <i>PLoS Medicine</i> , 2014, 11, e1001653.	3.9	96
242	Psychological as well as illness factors influence acceptance of non-invasive ventilation (NIV) and gastrostomy in amyotrophic lateral sclerosis (ALS): A prospective population study. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2014, 15, 376-387.	1.1	42
243	End-of-life care in general practice: A cross-sectional, retrospective survey of "cancer", "organ failure" and "old-age/dementia" patients. <i>Palliative Medicine</i> , 2014, 28, 965-975.	1.3	53
244	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.	0.1	18
245	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.	1.3	86
246	International variation in classification of dialysis withdrawal: a systematic review. <i>Nephrology Dialysis Transplantation</i> , 2014, 29, 625-635.	0.4	37
247	Implementing patient-reported outcome measures in palliative care clinical practice: A systematic review of facilitators and barriers. <i>Palliative Medicine</i> , 2014, 28, 158-175.	1.3	266
248	An integrated palliative and respiratory care service for patients with advanced disease and refractory breathlessness: a randomised controlled trial. <i>Lancet Respiratory Medicine</i> , 2014, 2, 979-987.	5.2	464
249	The presence of CD4 counts for the management of HIV patients in East Africa: A multicentred study. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 2014, 26, 613-618.	0.6	8
250	Palliative care-related self-report problems among cancer patients in East Africa: a two-country study. <i>Supportive Care in Cancer</i> , 2014, 22, 3185-3192.	1.0	15
251	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.	0.8	26
252	Wellbeing among sub-Saharan African patients with advanced HIV and/or cancer: an international multicentred comparison study of two outcome measures. <i>Health and Quality of Life Outcomes</i> , 2014, 12, 80.	1.0	8

#	ARTICLE	IF	CITATIONS
253	Quality of life and wellbeing among HIV outpatients in East Africa: a multicentre observational study. <i>BMC Infectious Diseases</i> , 2014, 14, 613.	1.3	16
254	Is a specialist breathlessness service more effective and cost-effective for patients with advanced cancer and their carers than standard care? Findings of a mixed-method randomised controlled trial. <i>BMC Medicine</i> , 2014, 12, 194.	2.3	155
255	The Liverpool Care Pathway for cancer patients dying in hospital medical wards: A before-and-after cluster phase II trial of outcomes reported by family members. <i>Palliative Medicine</i> , 2014, 28, 10-17.	1.3	28
256	How many people need palliative care? A study developing and comparing methods for population-based estimates. <i>Palliative Medicine</i> , 2014, 28, 49-58.	1.3	339
257	Episodic Breathlessness: A Clinically Relevant and Rising Issue. <i>Annals of the American Thoracic Society</i> , 2014, 11, 274-274.	1.5	2
258	Patterns of dignity-related distress at the end of life: A cross-sectional study of patients with advanced cancer and care home residents. <i>Palliative Medicine</i> , 2014, 28, 1118-1127.	1.3	17
259	Comparisons of Costs between Black Caribbean and White British Patients with Advanced Multiple Sclerosis in the UK. <i>Multiple Sclerosis International</i> , 2014, 2014, 1-12.	0.4	5
260	Stressors and Resources of Caregivers of Patients With Incurable Progressive Illness in Sub-Saharan Africa. <i>Qualitative Health Research</i> , 2014, 24, 317-328.	1.0	34
261	I've had a good life, what's left is a bonus: Factor analysis of the Mental Adjustment to Cancer Scale in a palliative care population. <i>Palliative Medicine</i> , 2014, 28, 243-255.	1.3	5
262	How Are Patient Populations Characterized in Studies Investigating Depression in Advanced Cancer? Results From a Systematic Literature Review. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 678-698.	0.6	15
263	Maintaining dignity for residents of care homes: A qualitative study of the views of care home staff, community nurses, residents and their families. <i>Geriatric Nursing</i> , 2014, 35, 55-60.	0.9	56
264	Breathlessness, Functional Status, Distress, and Palliative Care Needs Over Time in Patients With Advanced Chronic Obstructive Pulmonary Disease or Lung Cancer: A Cohort Study. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 569-581.e1.	0.6	90
265	Public preferences and priorities for end-of-life care in Kenya: a population-based street survey. <i>BMC Palliative Care</i> , 2014, 13, 4.	0.8	41
266	Liverpool Care Pathway for patients with cancer in hospital: a cluster randomised trial. <i>Lancet</i> , The, 2014, 383, 226-237.	6.3	76
267	Making Palliative Care Matter. <i>Journal of Palliative Medicine</i> , 2014, 17, 1292-1293.	0.6	1
268	End-of-life care—what do cancer patients want?. <i>Nature Reviews Clinical Oncology</i> , 2014, 11, 100-108.	12.5	74
269	Inclusion of end-of-life care in the global health agenda. <i>The Lancet Global Health</i> , 2014, 2, e375-e376.	2.9	27
270	Understanding what matters most to people with multiple myeloma: a qualitative study of views on quality of life. <i>BMC Cancer</i> , 2014, 14, 496.	1.1	65



#	ARTICLE	IF	CITATIONS
271	Feasibility of assessing quality of care at the end of life in two cluster trials using an after-death approach with multiple assessments. <i>BMC Palliative Care</i> , 2014, 13, 36.	0.8	2
272	Self-report measurement of pain & symptoms in palliative care patients: a comparison of verbal, visual and hand scoring methods in Sub-Saharan Africa. <i>Health and Quality of Life Outcomes</i> , 2014, 12, 118.	1.0	20
273	Reversal of English trend towards hospital death in dementia: a population-based study of place of death and associated individual and regional factors, 2001–2010. <i>BMC Neurology</i> , 2014, 14, 59.	0.8	88
274	Definition, Categorization, and Terminology of Episodic Breathlessness: Consensus by an International Delphi Survey. <i>Journal of Pain and Symptom Management</i> , 2014, 47, 828-838.	0.6	67
275	Are There Differences in the Prevalence of Palliative Care-Related Problems in People Living With Advanced Cancer and Eight Non-Cancer Conditions? A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 660-677.	0.6	322
276	A Psychometric Evaluation of the Functional Assessment of Chronic Illness Therapy-Palliative Care (FACIT-Pal) Scale With Palliative Care Samples in Three African Countries. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 983-991.	0.6	14
277	Experience of persistent psychological symptoms and perceived stigma among people with HIV on antiretroviral therapy (ART): A systematic review. <i>International Journal of Nursing Studies</i> , 2014, 51, 1171-1189.	2.5	146
278	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.	2.5	38
279	Avoid “prognostic paralysis” just get ahead and plan and co-ordinate care. <i>Npj Primary Care Respiratory Medicine</i> , 2014, 24, 14085.	1.1	9
280	Availability of essential drugs for managing HIV-related pain and symptoms within 120 PEPFAR-funded health facilities in East Africa: A cross-sectional survey with onsite verification. <i>Palliative Medicine</i> , 2014, 28, 293-301.	1.3	43
281	Holistic models for end of life care: Establishing the place of culture. <i>Progress in Palliative Care</i> , 2014, 22, 80-87.	0.7	12
282	Geographical and temporal Understanding In place of Death in England (1984–2010): analysis of trends and associated factors to improve end-of-life Care (GUIDE_Care) – primary research. <i>Health Services and Delivery Research</i> , 2014, 2, 1-104.	1.4	13
283	Meanings of happiness among two ethnic groups living with advanced cancer in south London: a qualitative study. <i>Psycho-Oncology</i> , 2013, 22, 1096-1103.	1.0	16
284	“Peace” and “life worthwhile” as measures of spiritual well-being in African palliative care: a mixed-methods study. <i>Health and Quality of Life Outcomes</i> , 2013, 11, 94.	1.0	37
285	“Burden to others” as a public concern in advanced cancer: a comparative survey in seven European countries. <i>BMC Cancer</i> , 2013, 13, 105.	1.1	35
286	Heterogeneity and changes in preferences for dying at home: a systematic review. <i>BMC Palliative Care</i> , 2013, 12, 7.	0.8	710
287	Evaluating complex interventions in End of Life Care: the MORECare Statement on good practice generated by a synthesis of transparent expert consultations and systematic reviews. <i>BMC Medicine</i> , 2013, 11, 111.	2.3	256
288	Cognitive interviewing of bereaved relatives to improve the measurement of health outcomes and care utilisation at the end of life in a mortality followback survey. <i>Supportive Care in Cancer</i> , 2013, 21, 2835-2844.	1.0	11

#	ARTICLE	IF	CITATIONS
289	How to Analyze Palliative Care Outcome Data for Patients in Sub-Saharan Africa: An International, Multicenter, Factor Analytic Examination of the APCA African POS. <i>Journal of Pain and Symptom Management</i> , 2013, 45, 746-752.	0.6	25
290	Episodic Breathlessness in Patients With Advanced Disease: A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2013, 45, 561-578.	0.6	55
291	'My dreams are shattered down and it hurts lots'—a qualitative study of palliative care needs and their management by HIV outpatient services in Kenya and Uganda. <i>BMC Palliative Care</i> , 2013, 12, 35.	0.8	33
292	Development and evaluation of the feasibility and effects on staff, patients, and families of a new tool, the Psychosocial Assessment and Communication Evaluation (PACE), to improve communication and palliative care in intensive care and during clinical uncertainty. <i>BMC Medicine</i> , 2013, 11, 213.	2.3	41
293	A randomised, multicentre clinical trial of specialised palliative care plus standard treatment versus standard treatment alone for cancer patients with palliative care needs: the Danish palliative care trial (DanPaCT) protocol. <i>BMC Palliative Care</i> , 2013, 12, 37.	0.8	17
294	Neuromuscular electrical stimulation for muscle weakness in adults with advanced disease. , 2013, , CDO09419.		60
295	Symptom prevalence, severity and palliative care needs assessment using the Palliative Outcome Scale: A cross-sectional study of patients with Parkinson's disease and related neurological conditions. <i>Palliative Medicine</i> , 2013, 27, 722-731.	1.3	77
296	Mixed Methods Research in the Development and Evaluation of Complex Interventions in Palliative and End-of-Life Care: Report on the MORECare Consensus Exercise. <i>Journal of Palliative Medicine</i> , 2013, 16, 1550-1560.	0.6	34
297	Living with breathlessness: A survey of caregivers of breathless patients with lung cancer or heart failure. <i>Palliative Medicine</i> , 2013, 27, 647-656.	1.3	82
298	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.	0.6	61
299	Episodic and Continuous Breathlessness: A New Categorization of Breathlessness. <i>Journal of Pain and Symptom Management</i> , 2013, 45, 1019-1029.	0.6	56
300	'Best practice' in developing and evaluating palliative and end-of-life care services: A meta-synthesis of research methods for the MORECare project. <i>Palliative Medicine</i> , 2013, 27, 885-898.	1.3	73
301	Research into palliative care in sub-Saharan Africa. <i>Lancet Oncology</i> , The, 2013, 14, e183-e188.	5.1	113
302	'It Makes You Feel That Somebody Is Out There Caring': A Qualitative Study of Intervention and Control Participants' Perceptions of the Benefits of Taking Part in an Evaluation of Dignity Therapy for People With Advanced Cancer. <i>Journal of Pain and Symptom Management</i> , 2013, 45, 712-725.	0.6	33
303	The minimal important difference of the King's Brief Interstitial Lung Disease Questionnaire (K-BILD) and forced vital capacity in interstitial lung disease. <i>Respiratory Medicine</i> , 2013, 107, 1438-1443.	1.3	39
304	A Psychometric Validation of Two Brief Measures to Assess Palliative Need in Patients Severely Affected by Multiple Sclerosis. <i>Journal of Pain and Symptom Management</i> , 2013, 46, 406-412.	0.6	28
305	Interventions to improve symptoms and quality of life of patients with fibrotic interstitial lung disease: a systematic review of the literature. <i>Thorax</i> , 2013, 68, 867-879.	2.7	86
306	'I wish I knew more ...' the end-of-life planning and information needs for end-stage fibrotic interstitial lung disease: views of patients, carers and health professionals: Table 1. <i>BMJ Supportive and Palliative Care</i> , 2013, 3, 84-90.	0.8	71

#	ARTICLE	IF	CITATIONS
307	Capturing activity, costs, and outcomes: The challenges to be overcome for successful economic evaluation in palliative care. <i>Progress in Palliative Care</i> , 2013, 21, 232-235.	0.7	10
308	MORECare research methods guidance development: Recommendations for ethical issues in palliative and end-of-life care research. <i>Palliative Medicine</i> , 2013, 27, 908-917.	1.3	68
309	Recommendations for managing missing data, attrition and response shift in palliative and end-of-life care research: Part of the MORECare research method guidance on statistical issues. <i>Palliative Medicine</i> , 2013, 27, 899-907.	1.3	80
310	Place of death, and its relation with underlying cause of death, in Parkinson's disease, motor neurone disease, and multiple sclerosis: A population-based study. <i>Palliative Medicine</i> , 2013, 27, 840-846.	1.3	60
311	The development and validation of the King's Sarcoidosis Questionnaire for the assessment of health status. <i>Thorax</i> , 2013, 68, 57-65.	2.7	92
312	Changing Patterns in Place of Cancer Death in England: A Population-Based Study. <i>PLoS Medicine</i> , 2013, 10, e1001410.	3.9	138
313	Time for a prospective study to evaluate the Amber Care Bundle. <i>BMJ Supportive and Palliative Care</i> , 2013, 3, 376-377.	0.8	13
314	Social-cultural factors in end-of-life care in Belgium: A scoping of the research literature. <i>Palliative Medicine</i> , 2013, 27, 131-143.	1.3	14
315	Episodes of breathlessness: Types and patterns – a qualitative study exploring experiences of patients with advanced diseases. <i>Palliative Medicine</i> , 2013, 27, 524-532.	1.3	54
316	Non-pharmacological interventions for breathlessness in advanced stages of malignant and non-malignant diseases. <i>The Cochrane Library</i> , 2013, , CD005623.	1.5	80
317	Epidemiology of Pain in Cancer. , 2013, , 5-24.		8
318	“It makes me feel that I’m still relevant”: A qualitative study of the views of nursing home residents on dignity therapy and taking part in a phase II randomised controlled trial of a palliative care psychotherapy. <i>Palliative Medicine</i> , 2013, 27, 358-366.	1.3	23
319	Breathlessness – current and emerging mechanisms, measurement and management: A discussion from an European Association of Palliative Care workshop. <i>Palliative Medicine</i> , 2013, 27, 932-938.	1.3	25
320	The palliative care needs for fibrotic interstitial lung disease: A qualitative study of patients, informal caregivers and health professionals. <i>Palliative Medicine</i> , 2013, 27, 869-876.	1.3	131
321	Advancing Palliative Care by Learning. <i>Journal of Palliative Medicine</i> , 2013, 16, 1164-1165.	0.6	0
322	Characterizing Episodic Breathlessness in Patients with Advanced Disease. <i>Journal of Palliative Medicine</i> , 2013, 16, 1275-1279.	0.6	28
323	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.	1.3	41
324	Standardize records of place of death. <i>Nature</i> , 2013, 495, 449-449.	13.7	3

#	ARTICLE	IF	CITATIONS
325	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.	0.7	51
326	Dying at home “ is it better: A narrative appraisal of the state of the science. <i>Palliative Medicine</i> , 2013, 27, 918-924.	1.3	132
327	Evidence on home palliative care: Charting past, present, and future at the Cicely Saunders Institute “ WHO Collaborating Centre for Palliative Care, Policy and Rehabilitation. <i>Progress in Palliative Care</i> , 2013, 21, 204-213.	0.7	13
328	Progression, Symptoms and Psychosocial Concerns among Those Severely Affected by Multiple Sclerosis: A Mixed-Methods Cross-Sectional Study of Black Caribbean and White British People. <i>PLoS ONE</i> , 2013, 8, e75431.	1.1	28
329	Multidimensional Patient-Reported Problems within Two Weeks of HIV Diagnosis in East Africa: A Multicentre Observational Study. <i>PLoS ONE</i> , 2013, 8, e57203.	1.1	14
330	Diversity in Defining End of Life Care: An Obstacle or the Way Forward?. <i>PLoS ONE</i> , 2013, 8, e68002.	1.1	30
331	Bereaved relatives’s™ views about participating in cancer research. <i>Palliative Medicine</i> , 2012, 26, 379-383.	1.3	35
332	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.	0.8	3
333	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.	0.6	430
334	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.	0.9	15
335	Challenges in defining “palliative care”™ for the purposes of clinical trials. <i>Current Opinion in Supportive and Palliative Care</i> , 2012, 6, 471-482.	0.5	17
336	The development and validation of the King’s Brief Interstitial Lung Disease (K-BILD) health status questionnaire. <i>Thorax</i> , 2012, 67, 804-810.	2.7	180
337	Opioid Prescribing for Cancer Pain During the Last 3 Months of Life: Associated Factors and 9-Year Trends in a Nationwide United Kingdom Cohort Study. <i>Journal of Clinical Oncology</i> , 2012, 30, 4373-4379.	0.8	62
338	Are we heading in the same direction? European and African doctors’s™ and nurses’s™ views and experiences regarding outcome measurement in palliative care. <i>Palliative Medicine</i> , 2012, 26, 242-249.	1.3	27
339	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.	0.8	17
340	Feasibility, acceptability and potential effectiveness of Dignity Therapy for older people in care homes: A phase II randomized controlled trial of a brief palliative care psychotherapy. <i>Palliative Medicine</i> , 2012, 26, 703-712.	1.3	82
341	Adjusting to the caregiving role: the importance of coping and support. <i>International Journal of Palliative Nursing</i> , 2012, 18, 541-545.	0.2	24
342	Predictors of non-remission of depression in a palliative care population. <i>Palliative Medicine</i> , 2012, 26, 683-695.	1.3	18

#	ARTICLE	IF	CITATIONS
343	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.	0.6	31
344	Intensity and correlates of multidimensional problems in HIV patients receiving integrated palliative care in sub-Saharan Africa. <i>Sexually Transmitted Infections</i> , 2012, 88, 607-611.	0.8	14
345	Reversal of the British trends in place of death: Time series analysis 2004–2010. <i>Palliative Medicine</i> , 2012, 26, 102-107.	1.3	160
346	Research into end-of-life cancer care—investment is needed. <i>Lancet, The</i> , 2012, 379, 519.	6.3	20
347	Integration of palliative care throughout HIV disease. <i>Lancet Infectious Diseases, The</i> , 2012, 12, 571-575.	4.6	49
348	What are the perceived needs and challenges of informal caregivers in home cancer palliative care? Qualitative data to construct a feasible psycho-educational intervention. <i>Supportive Care in Cancer</i> , 2012, 20, 1975-1982.	1.0	73
349	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.	0.6	27
350	Population, mortality and place of death in Germany (1950–2050) – Implications for end-of-life care in the future. <i>Public Health</i> , 2012, 126, 937-946.	1.4	44
351	Acceptability and Preferences of Six Different Routes of Drug Application for Acute Breathlessness: A Comparison Study between the United Kingdom and Germany. <i>Journal of Palliative Medicine</i> , 2012, 15, 1374-1381.	0.6	20
352	The “Spirit 8” successfully captured spiritual well-being in African palliative care: factor and Rasch analysis. <i>Journal of Clinical Epidemiology</i> , 2012, 65, 434-443.	2.4	27
353	Treatment outcomes in palliative care: the TOPCare study. A mixed methods phase III randomised controlled trial to assess the effectiveness of a nurse-led palliative care intervention for HIV positive patients on antiretroviral therapy. <i>BMC Infectious Diseases</i> , 2012, 12, 288.	1.3	21
354	What issues matter most to people with multiple myeloma and how well are we measuring them? A systematic review of quality of life tools. <i>European Journal of Haematology</i> , 2012, 89, 437-457.	1.1	64
355	Patient, caregiver, health professional and researcher views and experiences of participating in research at the end of life: a critical interpretive synthesis of the literature. <i>BMC Medical Research Methodology</i> , 2012, 12, 123.	1.4	75
356	Development, effectiveness and cost-effectiveness of a new out-patient Breathlessness Support Service: study protocol of a phase III fast-track randomised controlled trial. <i>BMC Pulmonary Medicine</i> , 2012, 12, 58.	0.8	34
357	Outcomes 'out of africa': the selection and implementation of outcome measures for palliative care in Africa. <i>BMC Palliative Care</i> , 2012, 11, 1.	0.8	38
358	“The greatest thing in the world is the family”™: the meaning of social support among Black Caribbean and White British patients living with advanced cancer. <i>Psycho-Oncology</i> , 2012, 21, 400-408.	1.0	19
359	Using the 12-item General Health Questionnaire to screen psychological distress from survivorship to end-of-life care: dimensionality and item quality. <i>Psycho-Oncology</i> , 2012, 21, 954-961.	1.0	29
360	Specialist Palliative Care is More Than Drugs: A Retrospective Study of ILD Patients. <i>Lung</i> , 2012, 190, 215-220.	1.4	94

#	ARTICLE	IF	CITATIONS
361	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.	1.0	29
362	End-of-life care across Southern Europe: A critical review of cultural similarities and differences between Italy, Spain and Portugal. <i>Critical Reviews in Oncology/Hematology</i> , 2012, 82, 387-401.	2.0	80
363	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.	0.6	57
364	Illness Perceptions, Adjustment to Illness, and Depression in a Palliative Care Population. <i>Journal of Pain and Symptom Management</i> , 2012, 43, 819-832.	0.6	34
365	Prevalence, Burden, and Correlates of Physical and Psychological Symptoms Among HIV Palliative Care Patients in Sub-Saharan Africa: An International Multicenter Study. <i>Journal of Pain and Symptom Management</i> , 2012, 44, 1-9.	0.6	62
366	A critical review of advance directives in Germany: Attitudes, use and healthcare professionals' compliance. <i>Patient Education and Counseling</i> , 2012, 87, 277-288.	1.0	64
367	A pan-European survey of research in end-of-life cancer care. <i>Supportive Care in Cancer</i> , 2012, 20, 39-48.	1.0	21
368	Circumstances of death and dying. , 2012, , 35-49.		2
369	Culture and End of Life Care: A Scoping Exercise in Seven European Countries. <i>PLoS ONE</i> , 2012, 7, e34188.	1.1	97
370	Symptoms and Quality of Life in Late Stage Parkinson Syndromes: A Longitudinal Community Study of Predictive Factors. <i>PLoS ONE</i> , 2012, 7, e46327.	1.1	83
371	The clinical epidemiology of depression in palliative care and the predictive value of somatic symptoms: Cross-sectional survey with four-week follow-up. <i>Palliative Medicine</i> , 2011, 25, 229-241.	1.3	72
372	How common are palliative care needs among older people who die in the emergency department?. <i>BMJ Supportive and Palliative Care</i> , 2011, 1, 184-188.	0.8	11
373	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.	1.3	89
374	The prevalence and burden of symptoms amongst cancer patients attending palliative care in two African countries. <i>European Journal of Cancer</i> , 2011, 47, 51-56.	1.3	95
375	The development of evidence-based European guidelines on the management of depression in palliative cancer care. <i>European Journal of Cancer</i> , 2011, 47, 702-712.	1.3	122
376	Breathlessness in cancer patients – Implications, management and challenges. <i>European Journal of Oncology Nursing</i> , 2011, 15, 459-469.	0.9	41
377	Antidepressants for the treatment of depression in neurological disorders: a systematic review and meta-analysis of randomised controlled trials. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2011, 82, 914-923.	0.9	73
378	The MVQOLI successfully captured quality of life in African palliative care: a factor analysis. <i>Journal of Clinical Epidemiology</i> , 2011, 64, 913-924.	2.4	20

#	ARTICLE	IF	CITATIONS
379	The effect of Palliative Daycare on Hope: A comparison of Daycare Patients with Two Control Groups. <i>Journal of Palliative Care</i> , 2011, 27, 216-223.	0.4	12
380	Interventions for improving palliative care for older people living in nursing care homes. <i>The Cochrane Library</i> , 2011, , CD007132.	1.5	90
381	Evaluating psycho-educational interventions for informal carers of patients receiving cancer care or palliative care: Strengths and limitations of different study designs. <i>Palliative Medicine</i> , 2011, 25, 345-356.	1.3	39
382	End-Stage Renal Disease: A New Trajectory of Functional Decline in the Last Year of Life. <i>Journal of the American Geriatrics Society</i> , 2011, 59, 304-308.	1.3	96
383	Prescription patterns of analgesics in the last 3 months of life: a retrospective analysis of 10202 lung cancer patients. <i>British Journal of Cancer</i> , 2011, 104, 1704-1710.	2.9	14
384	The Measurement of Spirituality in Palliative Care and the Content of Tools Validated Cross-Culturally: A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2011, 41, 728-753.	0.6	111
385	Identifying Psychological Distress at Key Stages of the Cancer Illness Trajectory: A Systematic Review of Validated Self-Report Measures. <i>Journal of Pain and Symptom Management</i> , 2011, 41, 619-636.	0.6	75
386	A Psychometric Evaluation of Measures of Spirituality Validated in Culturally Diverse Palliative Care Populations. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 604-622.	0.6	36
387	What Palliative Care-Related Problems Do Patients Experience at HIV Diagnosis? A Systematic Review of the Evidence. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 734-753.	0.6	46
388	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.	0.6	13
389	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.	0.6	43
390	Study Protocol: Phase III single-blinded fast-track pragmatic randomised controlled trial of a complex intervention for breathlessness in advanced disease. <i>Trials</i> , 2011, 12, 130.	0.7	40
391	Implementing a quality improvement programme in palliative care in care homes: a qualitative study. <i>BMC Geriatrics</i> , 2011, 11, 31.	1.1	38
392	Expert opinion on detecting and treating depression in palliative care: A Delphi study. <i>BMC Palliative Care</i> , 2011, 10, 10.	0.8	34
393	The lived experience of breathlessness and its implications for care: a qualitative comparison in cancer, COPD, heart failure and MND. <i>BMC Palliative Care</i> , 2011, 10, 15.	0.8	108
394	Appraisal of literature reviews on end-of-life care for minority ethnic groups in the UK and a critical comparison with policy recommendations from the UK end-of-life care strategy. <i>BMC Health Services Research</i> , 2011, 11, 141.	0.9	39
395	Quality of life among patients receiving palliative care in South Africa and Uganda: a multi-centred study. <i>Health and Quality of Life Outcomes</i> , 2011, 9, 21.	1.0	64
396	Implementing patient reported outcome measures (PROMs) in palliative care - users' cry for help. <i>Health and Quality of Life Outcomes</i> , 2011, 9, 27.	1.0	104

#	ARTICLE	IF	CITATIONS
397	Recruiting patients with advanced malignant and non-malignant disease: lessons learned from a palliative care RCT. <i>Trials</i> , 2011, 12, .	0.7	1
398	The Global Palliative Care Lottery: Can We Overcome It?. <i>Journal of Palliative Medicine</i> , 2011, 14, 384-385.	0.6	4
399	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.	0.6	18
400	Measuring the Net Benefits of Hospice and Palliative Care: A Composite Measure for Multiple Audiencesâ€™ Palliative Net Benefit. <i>Journal of Palliative Medicine</i> , 2011, 14, 264-265.	0.6	9
401	A novel approach to enhancing hope in patients with advanced cancer: a randomised phase II trial of dignity therapy. <i>BMJ Supportive and Palliative Care</i> , 2011, 1, 315-321.	0.8	85
402	Antidepressants for the treatment of depression in palliative care: systematic review and meta-analysis. <i>Palliative Medicine</i> , 2011, 25, 36-51.	1.3	144
403	The randomized fast-track trial in palliative care: Role, utility and ethics in the evaluation of interventions in palliative care?. <i>Palliative Medicine</i> , 2011, 25, 741-747.	1.3	25
404	Prevalence, course and associations of desire for hastened death in a UK palliative population: a cross-sectional study. <i>BMJ Supportive and Palliative Care</i> , 2011, 1, 140-148.	0.8	22
405	Evaluation of a new model of short-term palliative care for people severely affected with multiple sclerosis: a randomised fast-track trial to test timing of referral and how long the effect is maintained. <i>Postgraduate Medical Journal</i> , 2011, 87, 769-775.	0.9	58
406	How common are palliative care needs among older people who die in the emergency department?. <i>Emergency Medicine Journal</i> , 2011, 28, 491-495.	0.4	62
407	Palliative Cancer Care: An Epidemiologic Study. <i>Journal of Clinical Oncology</i> , 2011, 29, 646-650.	0.8	46
408	Assessing quality-of-life in older people in care homes. <i>Age and Ageing</i> , 2011, 40, 507-512.	0.7	39
409	Trajectories of Illness in Stage 5 Chronic Kidney Disease. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2011, 6, 1580-1590.	2.2	101
410	Developing a breathlessness intervention service for patients with palliative and supportive care needs, irrespective of diagnosis. <i>Journal of Palliative Care</i> , 2011, 27, 28-36.	0.4	20
411	What Is the Evidence That Palliative Care Teams Improve Outcomes for Cancer Patients and Their Families?. <i>Cancer Journal (Sudbury, Mass)</i> , 2010, 16, 423-435.	1.0	313
412	Benzodiazepines for the relief of breathlessness in advanced malignant and non-malignant diseases in adults. , 2010, , CD007354.		105
413	Does a social model of hospice day care affect advanced cancer patientsâ€™ use of other health and social services? A prospective quasi-experimental trial. <i>Supportive Care in Cancer</i> , 2010, 18, 627-637.	1.0	10
414	The experience of using the SEIQoL-DW with patients with advanced chronic obstructive pulmonary disease (COPD): issues of process and outcome. <i>Quality of Life Research</i> , 2010, 19, 619-629.	1.5	17



#	ARTICLE	IF	CITATIONS
415	The Experience of Breathlessness: The Social Course of Chronic Obstructive Pulmonary Disease. <i>Journal of Pain and Symptom Management</i> , 2010, 39, 555-563.	0.6	70
416	Psychological Well-Being and Quality of Care: A Factor-Analytic Examination of the Palliative Care Outcome Scale. <i>Journal of Pain and Symptom Management</i> , 2010, 40, 67-74.	0.6	52
417	Symptoms in the Month Before Death for Stage 5 Chronic Kidney Disease Patients Managed Without Dialysis. <i>Journal of Pain and Symptom Management</i> , 2010, 40, 342-352.	0.6	109
418	Variations in the quality and costs of end-of-life care, preferences and palliative outcomes for cancer patients by place of death: the QUALYCARE study. <i>BMC Cancer</i> , 2010, 10, 400.	1.1	66
419	Multi-centred mixed-methods PEPFAR HIV care & support public health evaluation: study protocol. <i>BMC Public Health</i> , 2010, 10, 584.	1.2	16
420	Validation of a core outcome measure for palliative care in Africa: the APCA African Palliative Outcome Scale. <i>Health and Quality of Life Outcomes</i> , 2010, 8, 10.	1.0	143
421	Effectiveness of a hand-held fan for breathlessness: a randomised phase II trial. <i>BMC Palliative Care</i> , 2010, 9, 22.	0.8	71
422	Review: Delivering research in end-of-life care: problems, pitfalls and future priorities. <i>Palliative Medicine</i> , 2010, 24, 456-461.	1.3	50
423	Palliative care for people severely affected by multiple sclerosis: evaluation of a novel palliative care service. <i>Multiple Sclerosis Journal</i> , 2010, 16, 627-636.	1.4	119
424	Clinical Effectiveness and Safety of Acupuncture in the Treatment of Irradiation-Induced Xerostomia in Patients with Head and Neck Cancer: A Systematic Review. <i>Acupuncture in Medicine</i> , 2010, 28, 191-199.	0.4	58
425	Palliative Care in the Tennessees: What Can We Expect?. <i>Journal of Palliative Medicine</i> , 2010, 13, 355-356.	0.6	0
426	Time to get it right: are preferences for place of death more stable than we think?. <i>Palliative Medicine</i> , 2010, 24, 352-353.	1.3	23
427	Fear of dying in an ethnically diverse society: cross-sectional studies of people aged 65+ in Britain. <i>Postgraduate Medical Journal</i> , 2010, 86, 197-202.	0.9	15
428	â€™A softening of edgesâ€™: a comparison of yoga classes at palliative care services in New Delhi and London. <i>International Journal of Palliative Nursing</i> , 2010, 16, 548-554.	0.2	9
429	Costâ€™effectiveness of lenalidomide in multiple myeloma. <i>Expert Review of Pharmacoeconomics and Outcomes Research</i> , 2010, 10, 229-238.	0.7	7
430	Results of a pilot investigation into a complex intervention for breathlessness in advanced chronic obstructive pulmonary disease (COPD): Brief report. <i>Palliative and Supportive Care</i> , 2010, 8, 143-149.	0.6	36
431	Individual breathlessness trajectories do not match summary trajectories in advanced cancer and chronic obstructive pulmonary disease: results from a longitudinal study. <i>Palliative Medicine</i> , 2010, 24, 777-786.	1.3	70
432	Short-form Zarit Caregiver Burden Interviews were valid in advanced conditions. <i>Journal of Clinical Epidemiology</i> , 2010, 63, 535-542.	2.4	230

#	ARTICLE	IF	CITATIONS
433	PRISMA: A pan-European co-ordinating action to advance the science in end-of-life cancer care. European Journal of Cancer, 2010, 46, 1493-1501.	1.3	54
434	Clinical priorities, barriers and solutions in end-of-life cancer care research across Europe. Report from a workshop. European Journal of Cancer, 2010, 46, 1815-1822.	1.3	37
435	Palliative care in cancer: How to improve clinical research. European Journal of Cancer, 2010, 46, 1486-1487.	1.3	1
436	Psychological distress in cancer from survivorship to end of life care: Prevalence, associated factors and clinical implications. European Journal of Cancer, 2010, 46, 2036-2044.	1.3	112
437	Refractory breathlessness: oxygen or room air?. Lancet, The, 2010, 376, 746-748.	6.3	5
438	Understanding Breathlessness: Cross-Sectional Comparison of Symptom Burden and Palliative Care Needs in Chronic Obstructive Pulmonary Disease and Cancer. Journal of Palliative Medicine, 2010, 13, 1109-1118.	0.6	202
439	Antidepressants for depression in physically ill people. The Cochrane Library, 2010, , CD007503.	1.5	130
440	Caring for a person in advanced illness and suffering from breathlessness at home: Threats and resources. Palliative and Supportive Care, 2009, 7, 153-162.	0.6	99
441	The Place Where They Have to Take You In or the Place Where You Prefer to Be?. Journal of Palliative Medicine, 2009, 12, 763-764.	0.6	2
442	Fast-Track Trials in Palliative Care: An Alternative Randomized Controlled Trial Design. Journal of Palliative Medicine, 2009, 12, 213-213.	0.6	15
443	Self-management for breathlessness in COPD: the role of pulmonary rehabilitation. Chronic Respiratory Disease, 2009, 6, 133-140.	1.0	29
444	Living and dying with dignity: a qualitative study of the views of older people in nursing homes. Age and Ageing, 2009, 38, 411-416.	0.7	88
445	Depression assessment and classification in palliative cancer patients: a systematic literature review. Palliative Medicine, 2009, 23, 739-753.	1.3	82
446	Vulnerability in palliative care research: findings from a qualitative study of black Caribbean and white British patients with advanced cancer. Journal of Medical Ethics, 2009, 35, 440-444.	1.0	29
447	Evaluation of hospital palliative care teams: strengths and weaknesses of the before-after study design and strategies to improve it. Palliative Medicine, 2009, 23, 23-28.	1.3	20
448	Effectiveness of Topical Administration of Opioids in Palliative Care: A Systematic Review. Journal of Pain and Symptom Management, 2009, 37, 913-917.	0.6	62
449	Optimal Approaches to the Health Economics of Palliative Care: Report of an International Think Tank. Journal of Pain and Symptom Management, 2009, 38, 4-10.	0.6	62
450	Research Priorities in Health Economics And Funding for Palliative Care: Views of an International Think Tank. Journal of Pain and Symptom Management, 2009, 38, 11-14.	0.6	20

#	ARTICLE	IF	CITATIONS
451	Palliative Care: No Longer a Luxury But a Necessity?. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 1-3.	0.6	26
452	Is Short-Term Palliative Care Cost-Effective in Multiple Sclerosis? A Randomized Phase II Trial. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 816-826.	0.6	166
453	Palliative care: Based on neither diagnosis nor prognosis, but patient and family need. Commentary on Chattoo and Atkin. <i>Social Science and Medicine</i> , 2009, 69, 154-157.	1.8	14
454	Reconciling employment with caring for a husband with an advanced illness. <i>BMC Health Services Research</i> , 2009, 9, 216.	0.9	14
455	Challenges to conducting research with older people living in nursing homes. <i>BMC Geriatrics</i> , 2009, 9, 38.	1.1	85
456	A Phase II randomised controlled trial assessing the feasibility, acceptability and potential effectiveness of Dignity Therapy for older people in care homes: Study protocol. <i>BMC Geriatrics</i> , 2009, 9, 9.	1.1	39
457	Assessing the feasibility, acceptability and potential effectiveness of Dignity Therapy for people with advanced cancer referred to a hospital-based palliative care team: Study protocol. <i>BMC Palliative Care</i> , 2009, 8, 5.	0.8	35
458	Provision of palliative care for chronic heart failure inpatients: how much do we need?. <i>BMC Palliative Care</i> , 2009, 8, 8.	0.8	17
459	The feasibility of a single-blinded fast-track pragmatic randomised controlled trial of a complex intervention for breathlessness in advanced disease. <i>BMC Palliative Care</i> , 2009, 8, 9.	0.8	42
460	Researching breathlessness in palliative care: consensus statement of the National Cancer Research Institute Palliative Care Breathlessness Subgroup. <i>Palliative Medicine</i> , 2009, 23, 213-227.	1.3	84
461	Pharmacological treatment of refractory breathlessness. <i>Expert Review of Respiratory Medicine</i> , 2009, 3, 21-36.	1.0	44
462	The detection of depression in palliative care. <i>Current Opinion in Supportive and Palliative Care</i> , 2009, 3, 55-60.	0.5	43
463	Meeting information needs of patients with incurable progressive disease and their families in South Africa and Uganda: multicentre qualitative study. <i>BMJ: British Medical Journal</i> , 2009, 338, b1326-b1326.	2.4	62
464	General practitioners' use and experiences of palliative care services: a survey in south east England. <i>BMC Palliative Care</i> , 2008, 7, 18.	0.8	16
465	Randomised controlled trial of a new palliative care service: Compliance, recruitment and completeness of follow-up. <i>BMC Palliative Care</i> , 2008, 7, 7.	0.8	49
466	Is the qualitative research interview an acceptable medium for research with palliative care patients and carers?. <i>BMC Medical Ethics</i> , 2008, 9, 7.	1.0	82
467	Cancer Trajectories at the End of Life: is there an effect of age and gender?. <i>BMC Cancer</i> , 2008, 8, 127.	1.1	20
468	Use of the Palliative Outcome Scale in Argentina: A Cross-Cultural Adaptation and Validation Study. <i>Journal of Pain and Symptom Management</i> , 2008, 35, 188-202.	0.6	29

#	ARTICLE	IF	CITATIONS
469	“œl Will Do It If It Will Help Others:” Motivations Among Patients Taking Part in Qualitative Studies in Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2008, 35, 347-355.	0.6	72
470	Meeting the Communication and Information Needs of Chronic Heart Failure Patients. <i>Journal of Pain and Symptom Management</i> , 2008, 36, 149-156.	0.6	131
471	Access to Services for Patients with Chronic Obstructive Pulmonary Disease: The Invisibility of Breathlessness. <i>Journal of Pain and Symptom Management</i> , 2008, 36, 451-460.	0.6	164
472	“œl know he controls cancer” The meanings of religion among Black Caribbean and White British patients with advanced cancer. <i>Social Science and Medicine</i> , 2008, 67, 780-789.	1.8	68
473	Caregiver assessment of patients with advanced cancer: concordance with patients, effect of burden and positivity. <i>Health and Quality of Life Outcomes</i> , 2008, 6, 42.	1.0	116
474	Where people die (1974”2030): past trends, future projections and implications for care. <i>Palliative Medicine</i> , 2008, 22, 33-41.	1.3	398
475	Non-pharmacological interventions for breathlessness in advanced stages of malignant and non-malignant diseases. , 2008, , CD005623.		180
476	Dying with cancer, living well with advanced cancer. <i>European Journal of Cancer</i> , 2008, 44, 1414-1424.	1.3	144
477	The etiology and management of intractable breathlessness in patients with advanced cancer: a systematic review of pharmacological therapy. <i>Nature Clinical Practice Oncology</i> , 2008, 5, 90-100.	4.3	112
478	The views of patients with advanced cancer regarding participation in serial questionnaire studies. <i>Palliative Medicine</i> , 2008, 22, 913-920.	1.3	11
479	Symptom Assessment in Palliative Care: A Need for International Collaboration. <i>Journal of Clinical Oncology</i> , 2008, 26, 3867-3873.	0.8	102
480	Cultural meanings of pain: a qualitative study of Black Caribbean and White British patients with advanced cancer. <i>Palliative Medicine</i> , 2008, 22, 350-359.	1.3	74
481	Improving generalist end of life care: national consultation with practitioners, commissioners, academics, and service user groups. <i>BMJ: British Medical Journal</i> , 2008, 337, a1720-a1720.	2.4	174
482	Measurement of dyspnoea in the clinical rather than the research setting. <i>Current Opinion in Supportive and Palliative Care</i> , 2008, 2, 95-99.	0.5	42
483	Methodological considerations for end-of-life research in patients with chronic kidney disease. <i>Journal of Nephrology</i> , 2008, 21, 268-82.	0.9	10
484	Death from Renal Failure Eighty Years On: How Far Have We Come?. <i>Journal of Palliative Medicine</i> , 2007, 10, 1236-1238.	0.6	9
485	The Use of Opioid Analgesia in End-Stage Renal Disease Patients Managed Without Dialysis. <i>Journal of Pain and Palliative Care Pharmacotherapy</i> , 2007, 21, 5-16.	0.5	4
486	Loss and change: experiences of people severely affected by multiple sclerosis. <i>Palliative Medicine</i> , 2007, 21, 101-107.	1.3	61

#	ARTICLE	IF	CITATIONS
487	The value of cognitive interviewing techniques in palliative care research. <i>Palliative Medicine</i> , 2007, 21, 87-93.	1.3	77
488	The end of life: unknown and unplanned?. <i>European Journal of Public Health</i> , 2007, 17, 331-332.	0.1	14
489	A tale of two cities: Factors affecting place of cancer death in London and New York. <i>European Journal of Public Health</i> , 2007, 17, 285-290.	0.1	49
490	Symptoms in Advanced Renal Disease: A Cross-Sectional Survey of Symptom Prevalence in Stage 5 Chronic Kidney Disease Managed without Dialysis. <i>Journal of Palliative Medicine</i> , 2007, 10, 1266-1276.	0.6	229
491	Do religious or spiritual beliefs influence bereavement? A systematic review. <i>Palliative Medicine</i> , 2007, 21, 207-217.	1.3	103
492	'Fighting for everything': service experiences of people severely affected by multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2007, 13, 660-667.	1.4	66
493	Improving end-of-life care for patients with chronic heart failure: "Let's hope it'll get better, when I know in my heart of hearts it won't". <i>Heart</i> , 2007, 93, 963-967.	1.2	153
494	Modelling services to meet the palliative care needs of chronic heart failure patients and their families: current practice in the UK. <i>Palliative Medicine</i> , 2007, 21, 385-390.	1.3	42
495	Rediscovering dignity at the bedside. <i>BMJ: British Medical Journal</i> , 2007, 335, 167-168.	2.4	5
496	Psychological, social and spiritual distress at the end of life in heart failure patients. <i>Current Opinion in Supportive and Palliative Care</i> , 2007, 1, 260-266.	0.5	50
497	Quality criteria valuable with slight modification. <i>Journal of Clinical Epidemiology</i> , 2007, 60, 1315.	2.4	13
498	Health Status and Service Needs of Male Inmates Seriously Ill With HIV/AIDS at Two Large Urban Jails. <i>American Journal of Men's Health</i> , 2007, 1, 213-223.	0.7	0
499	The Prevalence of Symptoms in End-Stage Renal Disease: A Systematic Review. <i>Advances in Chronic Kidney Disease</i> , 2007, 14, 82-99.	0.6	703
500	Experiences of breathlessness: A systematic review of the qualitative literature. <i>Palliative and Supportive Care</i> , 2007, 5, 281-302.	0.6	80
501	Does the patient-held record improve continuity and related outcomes in cancer care: a systematic review. <i>Health Expectations</i> , 2007, 10, 75-91.	1.1	76
502	Electroacupuncture as an Adjunctive Treatment to Control Neuropathic Pain in Patients with Cancer. <i>Journal of Pain and Symptom Management</i> , 2007, 33, 115-117.	0.6	29
503	Needs Assessments in Palliative Care: An Appraisal of Definitions and Approaches Used. <i>Journal of Pain and Symptom Management</i> , 2007, 33, 500-505.	0.6	57
504	Interactive technologies and videotapes for patient education in cancer care: systematic review and meta-analysis of randomised trials. <i>Supportive Care in Cancer</i> , 2007, 15, 7-20.	1.0	81

#	ARTICLE	IF	CITATIONS
505	Experimental and quasi-experimental designs. , 2007, , 85-98.		54
506	Outcome measurement. , 2007, , 99-114.		6
507	Systematic reviews. , 2007, , 115-136.		7
508	The use of opioid analgesia in end-stage renal disease patients managed without dialysis: recommendations for practice. <i>Journal of Pain and Palliative Care Pharmacotherapy</i> , 2007, 21, 5-16.	0.5	30
509	Symptom prevalence among people affected by advanced and progressive neurological conditions—a systematic review. <i>Journal of Palliative Care</i> , 2007, 23, 291-9.	0.4	9
510	Symptom Prevalence and Severity in People Severely Affected by Multiple Sclerosis. <i>Journal of Palliative Care</i> , 2006, 22, 158-165.	0.4	88
511	The impact of a breathlessness intervention service (BIS) on the lives of patients with intractable dyspnea: A qualitative phase 1 study. <i>Palliative and Supportive Care</i> , 2006, 4, 287-293.	0.6	83
512	End of life content in geriatric textbooks: what is the current situation?. <i>BMC Palliative Care</i> , 2006, 5, 5.	0.8	6
513	Study protocol: delayed intervention randomised controlled trial within the Medical Research Council (MRC) Framework to assess the effectiveness of a new palliative care service. <i>BMC Palliative Care</i> , 2006, 5, 7.	0.8	43
514	How Should Nations Measure the Quality of End-of-Life Care for Older Adults? Recommendations for an International Minimum Data Set. <i>Journal of the American Geriatrics Society</i> , 2006, 54, 1765-1771.	1.3	41
515	Chest Pain: Physician Perceptions and Decisionmaking in a London Emergency Department. <i>Annals of Emergency Medicine</i> , 2006, 48, 77-85.	0.3	22
516	Chest pain: Communication of symptoms and history in a London emergency department. <i>Patient Education and Counseling</i> , 2006, 63, 138-144.	1.0	23
517	A Comparison of Symptom Prevalence in Far Advanced Cancer, AIDS, Heart Disease, Chronic Obstructive Pulmonary Disease and Renal Disease. <i>Journal of Pain and Symptom Management</i> , 2006, 31, 58-69.	0.6	966
518	Factors influencing death at home in terminally ill patients with cancer: systematic review. <i>BMJ: British Medical Journal</i> , 2006, 332, 515-521.	2.4	796
519	Psychological services in hospices in the UK and Republic of Ireland. <i>Journal of the Royal Society of Medicine</i> , 2006, 99, 637-639.	1.1	19
520	Palliative care and the need for education - Do we know what makes a difference? A limited systematic review. <i>Health Education Journal</i> , 2006, 65, 101-125.	0.6	9
521	Is antiretroviral therapy associated with symptom prevalence and burden?. <i>International Journal of STD and AIDS</i> , 2006, 17, 400-405.	0.5	78
522	Discussion of Quality and Audit in Health. <i>Social Work in Public Health</i> , 2006, 22, 29-38.	0.2	1

#	ARTICLE	IF	CITATIONS
523	SYMPTOM MANAGEMENT IN PATIENTS WITH ESTABLISHED RENAL FAILURE MANAGED WITHOUT DIALYSIS. <i>Journal of Renal Care</i> , 2006, 32, 93-98.	0.2	34
524	Symptom prevalence and severity in people severely affected by multiple sclerosis. <i>Journal of Palliative Care</i> , 2006, 22, 158-65.	0.4	38
525	Does ethnicity matter? Bereavement outcomes in two ethnic groups living in the United Kingdom. <i>Palliative and Supportive Care</i> , 2005, 3, 183-190.	0.6	18
526	Living with motor neurone disease: lives, experiences of services and suggestions for change. <i>Health and Social Care in the Community</i> , 2005, 13, 64-74.	0.7	81
527	A New Approach to Eliciting Patients' Preferences for Palliative Day Care: The Choice Experiment Method. <i>Journal of Pain and Symptom Management</i> , 2005, 29, 435-445.	0.6	42
528	Validation and Clinical Application of the German Version of the Palliative Care Outcome Scale. <i>Journal of Pain and Symptom Management</i> , 2005, 30, 51-62.	0.6	120
529	Patient Reports of Symptoms and Their Treatment at Three Palliative Care Projects Servicing Individuals with HIV/AIDS. <i>Journal of Pain and Symptom Management</i> , 2005, 30, 408-417.	0.6	48
530	Communication training for health professionals who care for patients with cancer: a systematic review of training methods. <i>Supportive Care in Cancer</i> , 2005, 13, 356-366.	1.0	142
531	Systematic review of specialist palliative day-care for adults with cancer. <i>Supportive Care in Cancer</i> , 2005, 13, 607-627.	1.0	50
532	Patterns and predictors of place of cancer death for the oldest old. <i>BMC Palliative Care</i> , 2005, 4, 6.	0.8	39
533	Assessing the Effectiveness and Acceptability of Interprofessional Palliative Care Education. <i>Journal of Palliative Care</i> , 2005, 21, 262-269.	0.4	19
534	Access and equity in HIV/AIDS palliative care: a review of the evidence and responses. <i>Palliative Medicine</i> , 2005, 19, 251-258.	1.3	37
535	Pain and Symptom Control in HIV Disease: Under-Researched and Poorly Managed. <i>Clinical Infectious Diseases</i> , 2005, 40, 491-492.	2.9	20
536	Systematic review of hypnotherapy for treating symptoms in terminally ill adult cancer patients. <i>Palliative Medicine</i> , 2005, 19, 418-426.	1.3	65
537	A research study to identify facilitators and barriers to outcome measure implementation. <i>International Journal of Palliative Nursing</i> , 2005, 11, 218-225.	0.2	50
538	Palliative care in sub-Saharan Africa. <i>Lancet</i> , The, 2005, 365, 1971-1977.	6.3	186
539	Public Health and Palliative Care. <i>Clinics in Geriatric Medicine</i> , 2005, 21, 45-55.	1.0	37
540	End-of-Life Care: Lessons from Other Nations. <i>Journal of Palliative Medicine</i> , 2005, 8, s-161-s-173.	0.6	48

#	ARTICLE	IF	CITATIONS
541	Does palliative care improve outcomes for patients with HIV/AIDS? A systematic review of the evidence. <i>Sexually Transmitted Infections</i> , 2005, 81, 5-14.	0.8	112
542	Assessing palliative care outcomes for people with motor neurone disease living at home. <i>International Journal of Palliative Nursing</i> , 2004, 10, 449-453.	0.2	13
543	Developing methods to improve the quality of end-of-life care. <i>Journal of Interprofessional Care</i> , 2004, 18, 200-201.	0.8	7
544	Appropriate Methods to Assess the Effectiveness and Efficacy of Treatments or Interventions to Control Cancer Pain. <i>Journal of Palliative Medicine</i> , 2004, 7, 423-430.	0.6	13
545	It would be NICE to have more evidence?. <i>Palliative Medicine</i> , 2004, 18, 85-86.	1.3	15
546	Prevalence, severity, duration and measurement of pain in sub-Saharan Africa. <i>Palliative Medicine</i> , 2004, 18, 737-738.	1.3	7
547	Quality of life measures for the palliative care of people severely affected by multiple sclerosis: a systematic review. <i>Multiple Sclerosis Journal</i> , 2004, 10, 690-725.	1.4	74
548	Patterns of dying: palliative care for non-malignant disease. <i>Clinical Medicine</i> , 2004, 4, 39-44.	0.8	169
549	Spiritual needs in health care. <i>BMJ: British Medical Journal</i> , 2004, 329, 123-124.	2.4	80
550	Developing primary palliative care. <i>BMJ: British Medical Journal</i> , 2004, 329, 1056-1057.	2.4	106
551	Home or hospital? Choices at the end of life. <i>Journal of the Royal Society of Medicine</i> , 2004, 97, 413-414.	1.1	28
552	What methods do stakeholders prefer for feeding back performance data: a qualitative study in palliative care. <i>International Journal for Quality in Health Care</i> , 2004, 16, 375-381.	0.9	5
553	Service Needs of Patients with Advanced HIV Disease: A Comparison of Client and Staff Reports at Three Palliative Care Projects. <i>AIDS Patient Care and STDs</i> , 2004, 18, 145-158.	1.1	12
554	Dying to be Home? Preferred Location of Death of First-Generation Black Caribbean and Native-Born White Patients in the United Kingdom. <i>Journal of Palliative Medicine</i> , 2004, 7, 628-636.	0.6	36
555	Study of white coat effect on the prognosis of women with breast cancer. <i>European Journal of Cancer Care</i> , 2004, 13, 193-193.	0.7	0
556	Evaluation of a short-term group intervention for informal carers of patients attending a home palliative care service. <i>Journal of Pain and Symptom Management</i> , 2004, 27, 396-408.	0.6	69
557	It just didn't work: the realities of quality assessment in the English health care context. <i>International Journal of Nursing Studies</i> , 2004, 41, 705-712.	2.5	33
558	Communication training for health professionals who care for patients with cancer: a systematic review of effectiveness. <i>Supportive Care in Cancer</i> , 2004, 12, 692-700.	1.0	188



#	ARTICLE	IF	CITATIONS
559	Quality of life and survival prediction in terminal cancer patients. <i>Cancer</i> , 2004, 101, 1090-1098.	2.0	116
560	Pain experienced by lung cancer patients: a review of prevalence, causes and pathophysiology. <i>Lung Cancer</i> , 2004, 43, 247-257.	0.9	93
561	Relationship between three palliative care outcome scales. <i>Health and Quality of Life Outcomes</i> , 2004, 2, 68.	1.0	56
562	What is the potential for the use of clinical outcome measures to be computerised? Findings from a qualitative research study. <i>International Journal of Health Care Quality Assurance</i> , 2004, 17, 47-52.	0.2	4
563	Mental health status of clients from three HIV/AIDS palliative care projects. <i>Palliative and Supportive Care</i> , 2004, 2, 125-138.	0.6	11
564	Building user involvement in motor neurone disease: key lessons. <i>Journal of Interprofessional Care</i> , 2004, 18, 80-1.	0.8	1
565	Communication, information and support for adults with malignant cerebral glioma : a systematic literature review. <i>Supportive Care in Cancer</i> , 2003, 11, 21-29.	1.0	73
566	The relationship between patient characteristics and carer psychological status in home palliative cancer care. <i>Supportive Care in Cancer</i> , 2003, 11, 638-643.	1.0	62
567	Is There Evidence That Palliative Care Teams Alter End-of-Life Experiences of Patients and Their Caregivers?. <i>Journal of Pain and Symptom Management</i> , 2003, 25, 150-168.	0.6	447
568	Effectiveness of Palliative Day Care in Improving Pain, Symptom Control, and Quality of Life. <i>Journal of Pain and Symptom Management</i> , 2003, 25, 202-212.	0.6	74
569	Current HIV/AIDS end-of-life care in sub-Saharan Africa: a survey of models, services, challenges and priorities. <i>BMC Public Health</i> , 2003, 3, 33.	1.2	68
570	Support in the community for people with dementia and their carers: a comparative outcome study of specialist mental health service interventions. <i>International Journal of Geriatric Psychiatry</i> , 2003, 18, 298-307.	1.3	56
571	Translating clinical tools in nursing practice. <i>Journal of Advanced Nursing</i> , 2003, 44, 420-426.	1.5	18
572	Fit to care? A comparison of informal caregivers of first-generation Black Caribbeans and White dependants with advanced progressive disease in the UK. <i>Health and Social Care in the Community</i> , 2003, 11, 528-536.	0.7	14
573	Using satisfaction to measure the quality of palliative care: a review of the literature. <i>Journal of Advanced Nursing</i> , 2003, 42, 324-339.	1.5	127
574	JAN Forum: your views and letters. <i>Journal of Advanced Nursing</i> , 2003, 44, 653-653.	1.5	1
575	Improving palliative care for cancer. <i>Lancet Oncology</i> , The, 2003, 4, 73-74.	5.1	7
576	A virtual user involvement forum for people living with motor neuron disease. <i>Journal of Telemedicine and Telecare</i> , 2003, 9, 352-353.	1.4	2

#	ARTICLE	IF	CITATIONS
577	Effect of a palliative home care team on hospital admissions among patients with advanced cancer. Palliative Medicine, 2003, 17, 315-321.	1.3	74
578	What is the best way to help caregivers in cancer and palliative care? A systematic literature review of interventions and their effectiveness. Palliative Medicine, 2003, 17, 63-74.	1.3	343
579	Attitudes to Timeliness of Death and Euthanasia Among First Generation Black Caribbean and White Patients and Their Families Living in the United Kingdom. Journal of Palliative Medicine, 2003, 6, 245-249.	0.6	4
580	Palliative day care: what does it cost to run a centre and does attendance affect use of other services?. Palliative Medicine, 2003, 17, 628-637.	1.3	23
581	Children and young people who die from cancer: epidemiology and place of death in England (1995-9). BMJ: British Medical Journal, 2003, 327, 478-479.	2.4	45
582	Language translation of outcome measurement tools: views of health professionals. International Journal of Palliative Nursing, 2003, 9, 49-55.	0.2	11
583	Professionals's views and experiences of using outcome measures in palliative care. International Journal of Palliative Nursing, 2003, 9, 234-238.	0.2	34
584	MANAGING BEREAVEMENT IN THE CLASSROOM: A CONSPIRACY OF SILENCE?. Death Studies, 2003, 27, 717-741.	1.8	45
585	Complementary evidence?. Journal of the Royal Society of Medicine, 2003, 96, 427-428.	1.1	2
586	Identifying neuropathic pain in patients with head and neck cancer: use of the Leeds Assessment of Neuropathic Symptoms and Signs Scale. Journal of the Royal Society of Medicine, 2003, 96, 379-383.	1.1	61
587	Symptom Severity in Advanced Cancer, Assessed in Two Ethnic Groups by Interviews with Bereaved Family Members and Friends. Journal of the Royal Society of Medicine, 2003, 96, 10-16.	1.1	8
588	Complementary Evidence?. Journal of the Royal Society of Medicine, 2003, 96, 427-428.	1.1	1
589	Identifying Neuropathic Pain in Patients with Head and Neck Cancer: Use of the Leeds Assessment of Neuropathic Symptoms and Signs Scale. Journal of the Royal Society of Medicine, 2003, 96, 379-383.	1.1	44
590	Doctors should not discuss resuscitation with terminally ill patients: AGAINST. BMJ: British Medical Journal, 2003, 327, 615-616.	2.4	14
591	Symptom severity in advanced cancer, assessed in two ethnic groups by interviews with bereaved family members and friends. Journal of the Royal Society of Medicine, 2003, 96, 10-16.	1.1	15
592	Efficacy of acupuncture in asthma: systematic review and meta-analysis of published data from 11 randomised controlled trials. European Respiratory Journal, 2002, 20, 846-852.	3.1	73
593	Do we need palliative care audit in developing countries?. Palliative Medicine, 2002, 16, 546-547.	1.3	19
594	Communication in End-of-Life Cancer Care: A Comparison of Team Assessments in Three European Countries. Journal of Clinical Oncology, 2002, 20, 3674-3682.	0.8	61

#	ARTICLE	IF	CITATIONS
595	Religious faith and support at the end of life: a comparison of first generation black Caribbean and white populations. <i>Palliative Medicine</i> , 2002, 16, 540-541.	1.3	21
596	Marie Curie nurses: enabling patients with cancer to die at home. <i>British Journal of Community Nursing</i> , 2002, 7, 240-244.	0.2	12
597	Reports. <i>Journal of Interprofessional Care</i> , 2002, 16, 289-291.	0.8	6
598	An Evaluation of Systematic Reviews of Palliative Care Services. <i>Journal of Palliative Care</i> , 2002, 18, 77-83.	0.4	32
599	A Multi-Professional Short-Term Group Intervention for Informal Caregivers of Patients Using a Home Palliative Care Service. <i>Journal of Palliative Care</i> , 2002, 18, 275-281.	0.4	25
600	Do Hospital-Based Palliative Teams Improve Care for Patients or Families at the End of Life?. <i>Journal of Pain and Symptom Management</i> , 2002, 23, 96-106.	0.6	314
601	Accuracy of prognosis estimates by four palliative care teams: a prospective cohort study. <i>BMC Palliative Care</i> , 2002, 1, 1.	0.8	41
602	What is palliative day care?. <i>Supportive Care in Cancer</i> , 2002, 10, 556-562.	1.0	38
603	An evaluation of systematic reviews of palliative care services. <i>Journal of Palliative Care</i> , 2002, 18, 77-83.	0.4	11
604	A multi-professional short-term group intervention for informal caregivers of patients using a home palliative care service. <i>Journal of Palliative Care</i> , 2002, 18, 275-81.	0.4	11
605	Cancer pain epidemiology. , 2001, , 37-52.		3
606	Working with ambivalence: informal caregivers of patients at the end of life. <i>Supportive Care in Cancer</i> , 2001, 9, 642-645.	1.0	96
607	Quality of Life Assessment and Outcome of Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2001, 21, 179-188.	0.6	31
608	Efficiency of Searching the Grey Literature in Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2001, 22, 797-801.	0.6	42
609	Accounts of carers' satisfaction with health care at the end of life: A comparison of first generation black Caribbeans and white patients with advanced disease. <i>Palliative Medicine</i> , 2001, 15, 337-345.	1.3	73
610	Effective methods of giving information in cancer: a systematic literature review of randomized controlled trials. <i>Journal of Public Health</i> , 2001, 23, 227-234.	1.0	224
611	Measuring quality of life: Using quality of life measures in the clinical setting. <i>BMJ: British Medical Journal</i> , 2001, 322, 1297-1300.	2.4	737
612	The need for palliative care for patients with non-cancer diseases: a review of the evidence. <i>International Journal of Palliative Nursing</i> , 2001, 7, 221-226.	0.2	91

#	ARTICLE	IF	CITATIONS
613	Palliative care in the age of HIV / AIDS. Conclusions from the meeting. Journal of the Royal Society of Medicine, 2001, 94, 496-8.	1.1	0
614	Project to impROve management of terminal illnEss (PROMOTE). Journal of Interprofessional Care, 2001, 15, 398-9.	0.8	7
615	Music to be born to, music to die to. BMJ: British Medical Journal, 2000, 321, 1577-1579.	2.4	2
616	Assessing structure, process and outcome in palliative day care: a pilot study for a multicentre trial. Health and Social Care in the Community, 2000, 8, 336-344.	0.7	25
617	Is the Presence of Dyspnea a Risk Factor for Morbidity in Cancer Patients?. Journal of Pain and Symptom Management, 2000, 19, 15-22.	0.6	44
618	The quality of expectation: Healing, palliation or disappointment. Journal of the Royal Society of Medicine, 2000, 93, 609-610.	1.1	5
619	Palliative day care: what do services do?. Palliative Medicine, 2000, 14, 277-286.	1.3	54
620	Child and adolescent mental health service use. British Journal of Psychiatry, 2000, 177, 52-58.	1.7	130
621	Place of Care in Advanced Cancer: A Qualitative Systematic Literature Review of Patient Preferences. Journal of Palliative Medicine, 2000, 3, 287-300.	0.6	795
622	Correspondence. Minority ethnic groups and our healthier nation. Journal of Public Health, 2000, 22, 245-245.	1.0	4
623	Minimum dataset activity for hospice and hospital palliative care services in the UK 1997/98. Palliative Medicine, 2000, 14, 395-404.	1.3	41
624	The level of need for palliative care: a systematic review of the literature. Palliative Medicine, 2000, 14, 93-104.	1.3	126
625	Managing patients with lung cancer. BMJ: British Medical Journal, 2000, 320, 379-379.	2.4	4
626	Services, costs and appropriate outcomes in end of life care. Annals of Oncology, 1999, 10, 135-136.	0.6	9
627	Palliative Care Services in the Community: What Do Family Doctors Want?. Journal of Palliative Care, 1999, 15, 21-25.	0.4	42
628	What Constitutes High-Quality HIV/AIDS Palliative Care?. Journal of Palliative Care, 1999, 15, 5-12.	0.4	13
629	The impact of palliative medicine?. Palliative Medicine, 1999, 13, 273-274.	1.3	4
630	Do social factors affect where patients die: an analysis of 10 years of cancer deaths in England. Journal of Public Health, 1999, 21, 22-28.	1.0	105

#	ARTICLE	IF	CITATIONS
631	Home palliative care for terminal cancer patients: a survey on the final week of life. <i>Palliative Medicine</i> , 1999, 13, 233-241.	1.3	94
632	Systematic reviews. <i>Palliative Medicine</i> , 1999, 13, 75-80.	1.3	18
633	Terminal Cancer Patients and Timing of Referral to Palliative Care A Multicenter Prospective Cohort Study. <i>Journal of Pain and Symptom Management</i> , 1999, 18, 243-252.	0.6	82
634	Paddington Complexity Scale and Health of the Nation Outcome Scales for Children and Adolescents. <i>British Journal of Psychiatry</i> , 1999, 174, 417-423.	1.7	69
635	Palliative care needs to be provided on basis of need rather than diagnosis. <i>BMJ: British Medical Journal</i> , 1999, 318, 123-123.	2.4	32
636	Who needs palliative care?. <i>Journal of the Royal Society of Medicine</i> , 1998, 91, 563-564.	1.1	12
637	Do specialist palliative care teams improve outcomes for cancer patients? A systematic literature review. <i>Palliative Medicine</i> , 1998, 12, 317-332.	1.3	361
638	Accreditation of specialist palliative care: minimum standards or improved care?. <i>Palliative Medicine</i> , 1998, 12, 73-74.	1.3	0
639	Where do cancer patients die? Ten-year trends in the place of death of cancer patients in England. <i>Palliative Medicine</i> , 1998, 12, 353-363.	1.3	216
640	Outcome measures in palliative care for advanced cancer patients: a review. <i>Journal of Public Health</i> , 1997, 19, 193-199.	1.0	160
641	Family anxiety in advanced cancer: a multicentre prospective study in Ireland. <i>British Journal of Cancer</i> , 1997, 76, 1211-1214.	2.9	40
642	A multicenter evaluation of cancer pain control by Palliative Care Teams. <i>Journal of Pain and Symptom Management</i> , 1997, 14, 29-35.	0.6	78
643	Bereavement Follow-Up: What do Palliative Support Teams Actually Do?. <i>Journal of Palliative Care</i> , 1996, 12, 12-17.	0.4	33
644	Predictors of family anxiety in the weeks before bereavement. <i>Social Science and Medicine</i> , 1996, 43, 1621-1625.	1.8	31
645	Audit in palliative care: does practice change?. <i>European Journal of Cancer Care</i> , 1996, 5, 233-236.	0.7	17
646	Postgraduate research training: the PhD and MD thesis. <i>Palliative Medicine</i> , 1996, 10, 113-118.	1.3	14
647	Patients' Views Towards Care Received from Psychiatrists. <i>British Journal of Psychiatry</i> , 1996, 168, 641-646.	1.7	98
648	Review: Measures to determine the Outcome of Community Services for People with Dementia. <i>Age and Ageing</i> , 1995, 24, 73-83.	0.7	28

#	ARTICLE	IF	CITATIONS
649	Psychosocial issues in palliative care: The patient, the family, and the process and outcome of care. <i>Journal of Pain and Symptom Management</i> , 1995, 10, 142-150.	0.6	68
650	What do Palliative Staff think about Audit?. <i>Journal of Palliative Care</i> , 1995, 11, 17-19.	0.4	7
651	Two HIV/AIDS community support teams: Patient characteristics, problems at referral and during the last 6 weeks of life. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 1995, 7, 593-604.	0.6	14
652	A comparison of two measures of quality of life: their sensitivity and validity for patients with advanced cancer. <i>Palliative Medicine</i> , 1994, 8, 282-290.	1.3	46
653	Are bereaved family members a valid proxy for a patient's assessment of dying?. <i>Social Science and Medicine</i> , 1994, 38, 553-557.	1.8	185
654	Reducing hospital beds for patients with advanced cancer. <i>Lancet, The</i> , 1994, 344, 409.	6.3	25
655	Quality of care and evaluating services. <i>International Review of Psychiatry</i> , 1994, 6, 5-14.	1.4	11
656	Palliative care for people with HIV/AIDS: Views of patients, carers and providers. <i>AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV</i> , 1993, 5, 105-116.	0.6	32
657	Audit in palliative care. <i>Critical Public Health</i> , 1993, 4, 26-32.	1.4	8
658	Validity of the support team assessment schedule: do staffs' ratings reflect those made by patients or their families?. <i>Palliative Medicine</i> , 1993, 7, 219-228.	1.3	166
659	Palliative care: a review of past changes and future trends. <i>Journal of Public Health</i> , 1993, 15, 3-8.	1.0	89
660	Computer database for palliative care. <i>Lancet, The</i> , 1992, 340, 243.	6.3	2
661	Assessing the symptoms, anxiety and practical needs of HIV/AIDS patients receiving palliative care. <i>Quality of Life Research</i> , 1992, 1, 47-51.	1.5	31
662	Clinical audit by a palliative care team. <i>Palliative Medicine</i> , 1991, 5, 215-221.	1.3	37
663	Research degree supervision: Lottery or life belt. <i>Critical Public Health</i> , 1990, 1, 42-47.	1.4	1
664	Financial help for terminally ill patients. <i>Lancet, The</i> , 1990, 335, 172-172.	6.3	3
665	A Comparison of Four Outcome Measures of Terminal Care. , 1988, , 205-211.		6
666	Respiratory interventions for breathlessness in adults with advanced diseases. <i>The Cochrane Library</i> , 0, , .	1.5	1

#	ARTICLE	IF	CITATIONS
667	Physical interventions for breathlessness in adults with advanced diseases. The Cochrane Library, 0, , .	1.5	0
668	Better Breathlessness Services for Older People with Advanced Diseases: A Multicentre Discrete Choice Experiment and Economic Modelling (OPRBreathe). SSRN Electronic Journal, 0, , .	0.4	0
669	Meeting information needs of patients with incurable progressive disease and their families in South Africa and Uganda: multicentre qualitative study. , 0, .		3