

Jane E Seymour

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

151
papers

4,338
citations

36
h-index

59
g-index

161
ext. papers

5,017
ext. citations

3
avg, IF

5.66
L-index

#	Paper	IF	Citations
151	Existential suffering in the day to day lives of those living with palliative care needs arising from chronic obstructive pulmonary disease (COPD): A systematic integrative literature review.. <i>Palliative Medicine</i> , 2022 , 2692163221074539	5.5	1
150	Addressing current challenges in adult nursing: Describing a virtual consensus development project methodology. <i>Nursing Open</i> , 2021 ,	2.1	1
149	What are the psychological effects of mesothelioma on patients and their carers? A scoping review. <i>Psycho-Oncology</i> , 2020 , 29, 1464-1473	3.9	7
148	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial. <i>PLoS Medicine</i> , 2020 , 17, e1003422	11.6	13
147	International transfer and translation of an end of life care intervention: the case of the Liverpool Care Pathway for the dying patient. <i>Wellcome Open Research</i> , 2020 , 5, 256	4.8	2
146	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
145	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
144	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
143	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
142	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
141	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
140	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial 2020 , 17, e1003422		
139	Comparing the predictive ability of the Revised Minimum Dataset Mortality Risk Index (MMRI-R) with nurses' predictions of mortality among frail older people: a cohort study. <i>Age and Ageing</i> , 2019 , 48, 394-400	3	1
138	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	5.5	15
137	Patient and carer involvement in palliative care research: An integrative qualitative evidence synthesis review. <i>Palliative Medicine</i> , 2019 , 33, 969-984	5.5	24
136	The Practice of Mutual Protection in the Care of Children with Palliative Care Needs: A Multiple Qualitative Case Study Approach from Jordan. <i>Journal of Pediatric Nursing</i> , 2019 , 45, e9-e18	2.2	2
135	Do patients with lung cancer recall physician-initiated discussions about planning for end-of-life care following disclosure of a terminal prognosis?. <i>BMJ Supportive and Palliative Care</i> , 2019 , 9, 197-201	2.2	2

134	Supporting family carers in home-based end-of-life care: using participatory action research to develop a training programme for support workers and volunteers. <i>BMJ Supportive and Palliative Care</i> , 2019 , 9, e4	2.2	2
133	Reappraising 'the good death' for populations in the age of ageing. <i>Age and Ageing</i> , 2018 , 47, 328-330	3	15
132	Making sense of frailty: An ethnographic study of the experience of older people living with complex health problems. <i>International Journal of Older People Nursing</i> , 2018 , 13, e12172	2.3	16
131	The involvement of cancer patients in the four stages of decision-making preceding continuous sedation until death: A qualitative study. <i>Palliative Medicine</i> , 2018 , 32, 1198-1207	5.5	6
130	The Impact of Public Health Awareness Campaigns on the Awareness and Quality of Palliative Care. <i>Journal of Palliative Medicine</i> , 2018 , 21, S30-S36	2.2	31
129	Impact of the Macmillan specialist Care at Home service: a mixed methods evaluation across six sites. <i>BMC Palliative Care</i> , 2018 , 17, 36	3	5
128	The costs, resource use and cost-effectiveness of Clinical Nurse Specialist-led interventions for patients with palliative care needs: A systematic review of international evidence. <i>Palliative Medicine</i> , 2018 , 32, 447-465	5.5	28
127	The Liverpool Care Pathway for the Dying Patient: a critical analysis of its rise, demise and legacy in England. <i>Wellcome Open Research</i> , 2018 , 3, 15	4.8	12
126	The Liverpool Care Pathway for the Dying Patient: a critical analysis of its rise, demise and legacy in England. <i>Wellcome Open Research</i> , 2018 , 3, 15	4.8	19
125	Preregistration nursing students' experiences of a palliative care course in a resource-poor setting. <i>International Journal of Palliative Nursing</i> , 2018 , 24, 388-397	0.9	1
124	Palliative care in the USA and England: a critical analysis of meaning and implementation towards a public health approach. <i>Mortality</i> , 2017 , 22, 275-290	1	9
123	Older people's perspectives on dignity: the benefits and challenges of a qualitative longitudinal approach to researching experiences of later life. <i>International Journal of Social Research Methodology: Theory and Practice</i> , 2017 , 20, 647-658	2.8	9
122	Lung cancer diagnosed following an emergency admission: exploring patient and carer perspectives on delay in seeking help. <i>Supportive Care in Cancer</i> , 2017 , 25, 2259-2266	3.9	8
121	The importance of interdisciplinary communication in the process of anticipatory prescribing. <i>International Journal of Palliative Nursing</i> , 2017 , 23, 129-135	0.9	13
120	Definition and recommendations for advance care planning: an international consensus supported by the European Association for Palliative Care. <i>Lancet Oncology</i> , 2017 , 18, e543-e551	21.7	467
119	The Experience of Decision Making in the Care of Children with Palliative Care Needs: The Experiences of Jordanian Mothers. <i>Comprehensive Child and Adolescent Nursing</i> , 2017 , 40, 240-256	0.8	7
118	What do we know about the application of the Mental Capacity Act (2005) in healthcare practice regarding decision-making for frail and older people? A systematic literature review. <i>Health and Social Care in the Community</i> , 2017 , 25, 295-308	2.6	17
117	Advance care planning--a multi-centre cluster randomised clinical trial: the research protocol of the ACTION study. <i>BMC Cancer</i> , 2016 , 16, 264	4.8	34

116	Lung cancer diagnosed following an emergency admission: Mixed methods study of the management, outcomes and needs and experiences of patients and carers. <i>Respiratory Medicine</i> , 2016 , 114, 38-45	4.6	9
115	A qualitative evaluation of the impact of a palliative care course on preregistration nursing students' practice in Cameroon. <i>BMC Palliative Care</i> , 2016 , 15, 37	3	9
114	Cardiopulmonary resuscitation decisions in the emergency department: An ethnography of tacit knowledge in practice. <i>Social Science and Medicine</i> , 2016 , 156, 47-54	5.1	12
113	Sharing news of a lung cancer diagnosis with adult family members and friends: a qualitative study to inform a supportive intervention. <i>Patient Education and Counseling</i> , 2016 , 99, 378-385	3.1	15
112	Engaging Terminally Ill Patients in End of Life Talk: How Experienced Palliative Medicine Doctors Navigate the Dilemma of Promoting Discussions about Dying. <i>PLoS ONE</i> , 2016 , 11, e0156174	3.7	37
111	The language of sedation in end-of-life care: The ethical reasoning of care providers in three countries. <i>Health (United Kingdom)</i> , 2015 , 19, 339-54	1.9	19
110	A survey of older peoples' attitudes towards advance care planning. <i>Age and Ageing</i> , 2015 , 44, 371-6	3	84
109	Administering anticipatory medications in end-of-life care: a qualitative study of nursing practice in the community and in nursing homes. <i>Palliative Medicine</i> , 2015 , 29, 60-70	5.5	44
108	Seymour et al. Palliative sedation: Improvement of guidelines necessary, but not sufficient: A reply. <i>Palliative Medicine</i> , 2015 , 29, 481	5.5	1
107	An Educational Intervention to Reduce Pain and Improve Pain Management for Malawian People Living With HIV/AIDS and Their Family Carers: A Randomized Controlled Trial. <i>Journal of Pain and Symptom Management</i> , 2015 , 50, 80-90.e4	4.8	16
106	Descriptions by general practitioners and nurses of their collaboration in continuous sedation until death at home: in-depth qualitative interviews in three European countries. <i>Journal of Pain and Symptom Management</i> , 2015 , 49, 98-109	4.8	13
105	Living with AIDS in Uganda: a qualitative study of patients' and families' experiences following referral to hospice. <i>BMC Palliative Care</i> , 2015 , 14, 67	3	7
104	Using continuous sedation until death for cancer patients: a qualitative interview study of physicians' and nurses' practice in three European countries. <i>Palliative Medicine</i> , 2015 , 29, 48-59	5.5	67
103	Dying at home: a qualitative study of the perspectives of older South Asians living in the United Kingdom. <i>Palliative Medicine</i> , 2014 , 28, 264-72	5.5	23
102	Papavasiliou ES, Brearley SG, Seymour JE, Brown J, Payne SA, on behalf of EUROIMPACT. From sedation to continuous sedation until death: how has the conceptual basis of sedation in end-of-life care changed over time? <i>J Pain Symptom Manage</i> 2013; 5:706-723. <i>Journal of Pain and Symptom Management</i> , 2014 , 47, 370	4.8	2
101	How to communicate with patients about future illness progression and end of life: a systematic review. <i>BMJ Supportive and Palliative Care</i> , 2014 , 4, 331-41	2.2	71
100	Physicians' experiences and perspectives regarding the use of continuous sedation until death for cancer patients in the context of psychological and existential suffering at the end of life. <i>Psycho-Oncology</i> , 2014 , 23, 539-46	3.9	24
99	PRINCIPLES OF SPIRITUAL CARE IN END-OF-LIFE CARE IN INDIA: A HISTORICAL-CULTURAL INVESTIGATION. <i>BMJ Supportive and Palliative Care</i> , 2014 , 4, A18.1-A18	2.2	

98	A modified systematic review of research evidence about education for pre-registration nurses in palliative care. <i>BMC Palliative Care</i> , 2014 , 13, 56	3	38
97	Identity in the fourth age: perseverance, adaptation and maintaining dignity. <i>Ageing and Society</i> , 2014 , 34, 1-19	1.7	93
96	INFORMATION PROVISION AND THE WIDER EXPERIENCE OF CARE: A QUALITATIVE STUDY OF PATIENTS DIAGNOSED WITH LUNG CANCER FOLLOWING EMERGENCY ADMISSION AND THEIR CARERS. <i>BMJ Supportive and Palliative Care</i> , 2014 , 4, A25.3-A25	2.2	
95	Transitions at the end of life for older adults [patient, carer and professional perspectives: a mixed-methods study. <i>Health Services and Delivery Research</i> , 2014 , 2, 1-102	1.5	73
94	Peer education for advance care planning: volunteers' perspectives on training and community engagement activities. <i>Health Expectations</i> , 2013 , 16, 43-55	3.7	46
93	An educational intervention to reduce pain and improve pain management for Malawian people living with HIV/AIDS and their family carers: study protocol for a randomised controlled trial. <i>Trials</i> , 2013 , 14, 216	2.8	8
92	What is different about living alone with cancer in older age? A qualitative study of experiences and preferences for care. <i>BMC Family Practice</i> , 2013 , 14, 22	2.6	26
91	Lung cancer diagnosed following emergency admission: a mixed methods study protocol to improve understanding of patients' characteristics, needs, experiences and outcomes. <i>BMC Palliative Care</i> , 2013 , 12, 24	3	2
90	What is the extent of potentially avoidable admissions amongst hospital inpatients with palliative care needs?. <i>BMC Palliative Care</i> , 2013 , 12, 9	3	28
89	From 'conductor' to 'second fiddle': older adult care recipients' perspectives on transitions in family caring at hospital admission. <i>International Journal of Nursing Studies</i> , 2013 , 50, 1197-205	5.8	36
88	Living into old age with the consequences of breast cancer. <i>European Journal of Oncology Nursing</i> , 2013 , 17, 311-6	2.8	24
87	From sedation to continuous sedation until death: how has the conceptual basis of sedation in end-of-life care changed over time?. <i>Journal of Pain and Symptom Management</i> , 2013 , 46, 691-706	4.8	46
86	Continuous sedation (CS) until death: mapping the literature by bibliometric analysis. <i>Journal of Pain and Symptom Management</i> , 2013 , 45, 1073-1082.e10	4.8	22
85	The withdrawal of the Liverpool Care Pathway in England: implications for clinical practice and policy. <i>International Journal of Palliative Nursing</i> , 2013 , 19, 369-71	0.9	4
84	Extent of palliative care need in the acute hospital setting: a survey of two acute hospitals in the UK. <i>Palliative Medicine</i> , 2013 , 27, 76-83	5.5	67
83	Public attitudes to death and dying in the UK: a review of published literature. <i>BMJ Supportive and Palliative Care</i> , 2013 , 3, 37-45	2.2	34
82	Talking about end-of-life care: the perspectives of older South Asians living in East London. <i>Journal of Research in Nursing</i> , 2013 , 18, 394-406	0.9	16
81	Maintaining integrity in the face of death: a grounded theory to explain the perspectives of people affected by lung cancer about the expression of wishes for end of life care. <i>International Journal of Nursing Studies</i> , 2012 , 49, 718-26	5.8	34

80	The practice of continuous deep sedation until death in Flanders (Belgium), the Netherlands, and the U.K.: a comparative study. <i>Journal of Pain and Symptom Management</i> , 2012 , 44, 33-43	4.8	55
79	Older adults' experiences of transitions between care settings at the end of life in England: a qualitative interview study. <i>Journal of Pain and Symptom Management</i> , 2012 , 44, 74-83	4.8	47
78	The experiences of relatives with the practice of palliative sedation: a systematic review. <i>Journal of Pain and Symptom Management</i> , 2012 , 44, 431-45	4.8	49
77	A comparison of strategies to recruit older patients and carers to end-of-life research in primary care. <i>BMC Health Services Research</i> , 2012 , 12, 342	2.9	18
76	Why is the Liverpool care pathway used for some dying cancer patients and not others? Healthcare professionals' perspectives. <i>BMC Research Notes</i> , 2012 , 5, 524	2.3	9
75	Liminality as a framework for understanding the experience of cancer survivorship: a literature review. <i>Journal of Advanced Nursing</i> , 2012 , 68, 2155-64	3.1	63
74	Older patients' attitudes towards and experiences of patient-physician end-of-life communication: a secondary analysis of interviews from British, Dutch and Belgian patients. <i>BMC Palliative Care</i> , 2012 , 11, 24	3	14
73	After you: conversations between patients and healthcare professionals in planning for end of life care. <i>BMC Palliative Care</i> , 2012 , 11, 15	3	90
72	Muslim women's narratives about bodily change and care during critical illness: a qualitative study. <i>Journal of Nursing Scholarship</i> , 2012 , 44, 99-107	3.6	16
71	Funding health and social services for older people - a qualitative study of care recipients in the last year of life. <i>Journal of the Royal Society of Medicine</i> , 2012 , 105, 201-7	2.3	10
70	Combined qualitative and quantitative research designs. <i>Current Opinion in Supportive and Palliative Care</i> , 2012 , 6, 514-24	2.6	17
69	An evaluation of research capacity building from the Cancer Experiences Collaborative. <i>BMJ Supportive and Palliative Care</i> , 2012 , 2, 280-5	2.2	5
68	Breaking bad news sensitively: what is important to patients in their last year of life?. <i>BMJ Supportive and Palliative Care</i> , 2012 , 2, 24-8	2.2	26
67	'That's part of everybody's job': the perspectives of health care staff in England and New Zealand on the meaning and remit of palliative care. <i>Palliative Medicine</i> , 2012 , 26, 232-41	5.5	80
66	Looking back, looking forward: the evolution of palliative and end-of-life care in England. <i>Mortality</i> , 2012 , 17, 1-17	1	27
65	Using Focus Groups for Researching End-of-Life Care Issues with Older People 2012 , 127-145		
64	The role of health care assistants in supporting district nurses and family carers to deliver palliative care at home: findings from an evaluation project. <i>Journal of Clinical Nursing</i> , 2011 , 20, 2043-52	3.2	20
63	Is it recorded in the notes? Documentation of end-of-life care and preferred place to die discussions in the final weeks of life. <i>BMC Palliative Care</i> , 2011 , 10, 18	3	29

62	The perspectives of clinical staff and bereaved informal care-givers on the use of continuous sedation until death for cancer patients: The study protocol of the UNBIASED study. <i>BMC Palliative Care</i> , 2011 , 10, 5	3	23
61	Factors influencing the referral of non-cancer patients to community specialist palliative care nurses. <i>International Journal of Palliative Nursing</i> , 2011 , 17, 35-41	0.9	13
60	Perspectives of staff providing care at the end of life for people with progressive long-term neurological conditions. <i>Palliative and Supportive Care</i> , 2011 , 9, 377-85	2.5	18
59	Older people living alone at the end of life in the U.K.: research and policy challenges. <i>Palliative Medicine</i> , 2011 , 25, 650-7	5.5	44
58	Jacqueline H. Watts, <i>Death, Dying and Bereavement: Issues for Practice</i> , Dunedin Academic Press, Edinburgh, UK, 2010, 124 pp., pbk £14.50, ISBN 13: 978 1 906716 08 0.. <i>Ageing and Society</i> , 2011 , 31, 173-174	1.7	17
57	Do nursing homes for older people have the support they need to provide end-of-life care? A mixed methods enquiry in England. <i>Palliative Medicine</i> , 2011 , 25, 125-38	5.5	63
56	Changing times: preparing to meet palliative needs in the 21st Century. <i>British Journal of Community Nursing</i> , 2011 , 16, 18	0.6	4
55	Interviews on end-of-life care with older people: reflections on six european studies. <i>Qualitative Health Research</i> , 2011 , 21, 1588-600	3.9	25
54	Death and Dying in America. <i>Sociology of Health and Illness</i> , 2010 , 32, 509-510	3	
53	The Mental Capacity Act 2005: Making decisions on capacity. <i>British Journal of Neuroscience Nursing</i> , 2010 , 6, 182-185	0.1	1
52	Working with the Mental Capacity Act: findings from specialist palliative and neurological care settings. <i>Palliative Medicine</i> , 2010 , 24, 396-402	5.5	25
51	Exploring the Impact of Sexual Orientation on Experiences and Concerns about End of Life Care and on Bereavement for Lesbian, Gay and Bisexual Older People. <i>Sociology</i> , 2010 , 44, 908-924	2.6	90
50	"At the foot of a very long ladder": discussing the end of life with older people and informal caregivers. <i>Journal of Pain and Symptom Management</i> , 2010 , 40, 857-69	4.8	48
49	Muslim women's experiences of suffering in Jordanian intensive care units: a narrative study. <i>Intensive and Critical Care Nursing</i> , 2010 , 26, 175-84	3.1	27
48	Implementing advance care planning: a qualitative study of community nurses' views and experiences. <i>BMC Palliative Care</i> , 2010 , 9, 4	3	97
47	Dying matters: let's talk about it. <i>BMJ, The</i> , 2010 , 341, c4860	5.9	31
46	Advance care planning in patients with end-stage renal disease 2010 , 49-74		2
45	Advance Care Planning for the end of life: an overview 2010 , 16-27		3

44	The role of the nurse in palliative care settings in a global context. <i>Cancer Nursing Practice</i> , 2009 , 8, 21-26.6		8
43	How to improve end of life care in acute hospitals. <i>Nursing Older People</i> , 2009 , 21, 26-9	0.6	18
42	Evaluating a peer education programme for advance end-of-life care planning for older adults: The peer educators' perspective. <i>International Journal on Disability and Human Development</i> , 2009 , 8,		9
41	Advance care planning: evidence and implications for practice 2009 , 3, 58-64		7
40	Older Chinese people's views on food: implications for supportive cancer care. <i>Ethnicity and Health</i> , 2008 , 13, 497-514	2.2	26
39	A UK literature review of progressive long-term neurological conditions. <i>British Journal of Community Nursing</i> , 2008 , 13, 206, 208-12	0.6	7
38	Exploring the palliative and end-of-life care needs of those affected by progressive long-term neurological conditions. <i>Journal of Care Services Management</i> , 2007 , 2, 92-102		3
37	Hospice or home? Expectations of end-of-life care among white and Chinese older people in the UK. <i>Sociology of Health and Illness</i> , 2007 , 29, 872-90	3	53
36	Relieving suffering at the end of life: practitioners' perspectives on palliative sedation from three European countries. <i>Social Science and Medicine</i> , 2007 , 64, 1679-91	5.1	46
35	Needs and preferences of patients with diseases other than cancer: the UK experience. <i>Progress in Palliative Care</i> , 2007 , 15, 108-112	1	3
34	Commentary 1. 'Quality of life to the end'. <i>Communications: the European Journal of Communication Research</i> , 2007 , 4, 117-8; discussion 125	0.5	2
33	Making sense of dying: a review of narratives written since 1950 by people facing death from cancer and other diseases. <i>Palliative Medicine</i> , 2006 , 20, 183-95	5.5	36
32	Advance care planning for patients with inoperable lung cancer. <i>International Journal of Palliative Nursing</i> , 2006 , 12, 172-8	0.9	28
31	Development of a peer education programme for advance end-of-life care planning. <i>International Journal of Palliative Nursing</i> , 2006 , 12, 216-223	0.9	32
30	The Cancer Experiences Research Collaborative (CECo): building research capacity in supportive and palliative care. <i>Progress in Palliative Care</i> , 2006 , 14, 265-270	1	7
29	Development of a peer education programme for advance end-of-life care planning. <i>International Journal of Palliative Nursing</i> , 2006 , 12, 214, 216-23	0.9	13
28	Mesothelioma from the patient's perspective. <i>Hematology/Oncology Clinics of North America</i> , 2005 , 19, 1175-90, viii	3.1	40
27	Stories of cancer pain: a historical perspective. <i>Journal of Pain and Symptom Management</i> , 2005 , 29, 22-31.8		14

26	The measurement of pain, 1945-2000. <i>Journal of Pain and Symptom Management</i> , 2005 , 29, 14-21	4.8	68
25	Pain and palliative care: the emergence of new specialties. <i>Journal of Pain and Symptom Management</i> , 2005 , 29, 2-13	4.8	29
24	Chinese Community Views: Promoting Cultural Competence in Palliative Care. <i>Journal of Palliative Care</i> , 2005 , 21, 111-116	1.8	31
23	Ethical and methodological issues in palliative care studies: The experiences of a research group. <i>Journal of Research in Nursing</i> , 2005 , 10, 169-188	0.9	20
22	Older people's views about home as a place of care at the end of life. <i>Palliative Medicine</i> , 2004 , 18, 460-75	5	254
21	Planning for the end of life: the views of older people about advance care statements. <i>Social Science and Medicine</i> , 2004 , 59, 57-68	5.1	131
20	Changing technologies of cancer pain relief: case studies of innovation. <i>Progress in Palliative Care</i> , 2004 , 12, 123-133	1	2
19	Economic evaluation of specialist cancer and palliative nursing: Macmillan evaluation study findings. <i>International Journal of Palliative Nursing</i> , 2003 , 9, 429-38	0.9	20
18	Changing technologies of cancer pain relief: themes from the twentieth century. <i>Progress in Palliative Care</i> , 2003 , 11, 256-260	1	2
17	Economic evaluation of specialist cancer and palliative nursing: a literature review. <i>International Journal of Palliative Nursing</i> , 2003 , 9, 424-8	0.9	7
16	Exploring nursing outcomes for patients with advanced cancer following intervention by Macmillan specialist palliative care nurses. <i>Journal of Advanced Nursing</i> , 2003 , 41, 561-74	3.1	52
15	Specialist palliative care: patients' experiences. <i>Journal of Advanced Nursing</i> , 2003 , 44, 24-33	3.1	20
14	JAN Forum: your views and letters. Response to: Watson's guest editorial 'Scientific methods are the only credible way forward for nursing research', <i>Journal of advanced Nursing</i> 43, 219-220. <i>Journal of Advanced Nursing</i> , 2003 , 44, 547-8	3.1	3
13	Clinical nurse specialists in palliative care. Part 1. A description of the MacMillan Nurse caseload. <i>Palliative Medicine</i> , 2002 , 16, 285-96	5.5	48
12	Clinical nurse specialists in palliative care. Part 2. Explaining diversity in the organization and costs of Macmillan nursing services. <i>Palliative Medicine</i> , 2002 , 16, 375-85	5.5	28
11	Clinical nurse specialists in palliative care. Part 3. Issues for the Macmillan Nurse role. <i>Palliative Medicine</i> , 2002 , 16, 386-94	5.5	46
10	Using focus groups to explore older people's attitudes to end of life care. <i>Ageing and Society</i> , 2002 , 22, 517-526	1.7	35
9	Meeting complex needs: an analysis of Macmillan nurses' work with patients. <i>International Journal of Palliative Nursing</i> , 2002 , 8, 574-82	0.9	28

8	Good deaths, bad deaths: Older people's assessments of the risks and benefits of morphine and terminal sedation in end-of-life care. <i>Health, Risk and Society</i> , 2002 , 4, 287-303	2	20
7	Analysing qualitative data: examples from two studies of end-of-life care. <i>International Journal of Palliative Nursing</i> , 2001 , 7, 227-33	0.9	10
6	Negotiating natural death in intensive care. <i>Social Science and Medicine</i> , 2000 , 51, 1241-52	5.1	80
5	Support and supervision in palliative care research. <i>Palliative Medicine</i> , 2000 , 14, 441-6	5.5	15
4	Revisiting medicalisation and 'natural' death. <i>Social Science and Medicine</i> , 1999 , 49, 691-704	5.1	64
3	Phenomenological approaches to palliative care research. <i>Palliative Medicine</i> , 1998 , 12, 127-31	5.5	26
2	Understanding the role of nurses in the management of symptoms and distress in the last days of life	100-115	4
1	Hospice or Home? Expectations of End-of-Life Care Among White and Chinese Older People in the UK	74-91	