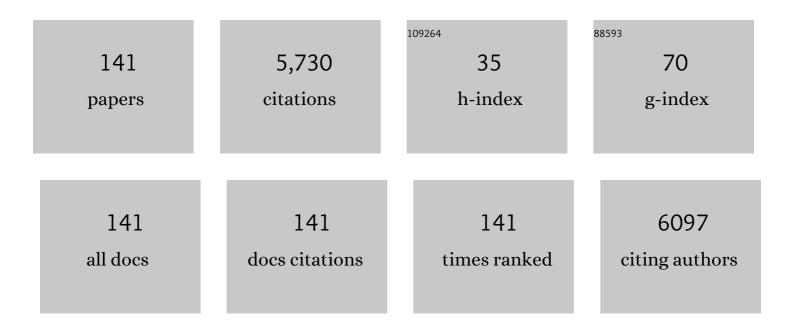
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Appraising the Evidence: Reviewing Disparate Data Systematically. Qualitative Health Research, 2002, 12, 1284-1299.	1.0	803
2	Guidance on Conducting and REporting DElphi Studies (CREDES) in palliative care: Recommendations based on a methodological systematic review. Palliative Medicine, 2017, 31, 684-706.	1.3	775
3	Definition and recommendations for advance care planning: an international consensus supported by the European Association for Palliative Care. Lancet Oncology, The, 2017, 18, e543-e551.	5.1	765
4	Family Caregivers and Palliative Care: Current Status and Agenda for the Future. Journal of Palliative Medicine, 2011, 14, 864-869.	0.6	126
5	Dying trajectories in heart failure. Palliative Medicine, 2007, 21, 95-99.	1.3	123
6	The Carer Support Needs Assessment Tool (CSNAT) for Use in Palliative and End-of-life Care at Home: A Validation Study. Journal of Pain and Symptom Management, 2013, 46, 395-405.	0.6	119
7	Family carers providing support to a person dying in the home setting: A narrative literature review. Palliative Medicine, 2015, 29, 487-495.	1.3	113
8	A systematic review of instruments related to family caregivers of palliative care patients. Palliative Medicine, 2010, 24, 656-668.	1.3	98
9	Using continuous sedation until death for cancer patients: A qualitative interview study of physicians' and nurses' practice in three European countries. Palliative Medicine, 2015, 29, 48-59.	1.3	94
10	Towards integration of palliative care in patients with chronic heart failure and chronic obstructive pulmonary disease: a systematic literature review of European guidelines and pathways. BMC Palliative Care, 2016, 15, 18.	0.8	93
11	A survey of end-of-life care in care homes: issues of definition and practice. Health and Social Care in the Community, 2006, 14, 341-348.	0.7	78
12	How continuity of care is experienced within the context of integrated palliative care: A qualitative study with patients and family caregivers in five European countries. Palliative Medicine, 2017, 31, 946-955.	1.3	76
13	End-of-life issues in acute stroke care: a qualitative study of the experiences and preferences of patients and families. Palliative Medicine, 2010, 24, 146-153.	1.3	75
14	The palliative care knowledge of nursing home staff: The EU FP7 PACE cross-sectional survey in 322 nursing homes in six European countries. Palliative Medicine, 2018, 32, 1487-1497.	1.3	73
15	Consensus Building in Palliative Care: A Europe-Wide Delphi Study on Common Understandings and Conceptual Differences. Journal of Pain and Symptom Management, 2012, 44, 192-205.	0.6	69
16	Palliative Care Development in European Care Homes and Nursing Homes: Application of a Typology of Implementation. Journal of the American Medical Directors Association, 2017, 18, 550.e7-550.e14.	1.2	68
17	Advance care planning in patients with advanced cancer: A 6-country, cluster-randomised clinical trial. PLoS Medicine, 2020, 17, e1003422.	3.9	68
18	Administering anticipatory medications in end-of-life care: A qualitative study of nursing practice in the community and in nursing homes. Palliative Medicine, 2015, 29, 60-70.	1.3	64

SHEILA PAYNE

#	Article	IF	CITATIONS
19	Evaluation of a Palliative Care Program for Nursing Homes in 7 Countries. JAMA Internal Medicine, 2020, 180, 233.	2.6	62
20	A narrative literature review of the contribution of volunteers in end-of-life care services. Palliative Medicine, 2013, 27, 428-436.	1.3	58
21	End-of-life care in general practice: A cross-sectional, retrospective survey of â€~cancer', â€~organ failure' and â€~old-age/dementia' patients. Palliative Medicine, 2014, 28, 965-975.	1.3	53
22	Volunteers in Specialist Palliative Care: A Survey of Adult Services in the United Kingdom. Journal of Palliative Medicine, 2014, 17, 568-574.	0.6	48
23	Comparing Palliative Care in Care Homes Across Europe (PACE): Protocol of a Cross-sectional Study of Deceased Residents in 6 EU Countries. Journal of the American Medical Directors Association, 2016, 17, 566.e1-566.e7.	1.2	47
24	Clinical Aspects of Palliative Sedation in Prospective Studies. A Systematic Review. Journal of Pain and Symptom Management, 2021, 61, 831-844.e10.	0.6	47
25	Exploring the experiences of bereaved families involved in assisted suicide in Southern Switzerland: a qualitative study: TableÂ1. BMJ Supportive and Palliative Care, 2015, 5, 146-152.	0.8	46
26	Integrated palliative care is about professional networking rather than standardisation of care: A qualitative study with healthcare professionals in 19 integrated palliative care initiatives in five European countries. Palliative Medicine, 2018, 32, 1091-1102.	1.3	46
27	Ageing and dying in the contemporary neoliberal prison system: Exploring the â€~double burden' for older prisoners. Social Science and Medicine, 2018, 212, 161-167.	1.8	46
28	Barriers to, and opportunities for, palliative care development in the Eastern Mediterranean Region. Lancet Oncology, The, 2017, 18, e176-e184.	5.1	45
29	Advance care planning – a multi-centre cluster randomised clinical trial: the research protocol of the ACTION study. BMC Cancer, 2016, 16, 264.	1.1	43
30	Cancer survivorship: Advancing the concept in the context of colorectal cancer. European Journal of Oncology Nursing, 2017, 29, 135-147.	0.9	43
31	From â€~conductor' to â€~second fiddle': Older adult care recipients' perspectives on transitions in fa caring at hospital admission. International Journal of Nursing Studies, 2013, 50, 1197-1205.	mily 2.5	42
32	Narrative Research Methods in Palliative Care Contexts: Two Case Studies. Journal of Pain and Symptom Management, 2009, 37, 788-796.	0.6	41
33	Family Caregivers' Reflections on Experiences of Assisted Suicide in Switzerland: A Qualitative Interview Study. Journal of Pain and Symptom Management, 2018, 55, 1085-1094.	0.6	41
34	Integrating palliative care in long-term care facilities across Europe (PACE): protocol of a cluster randomized controlled trial of the â€~PACE Steps to Success' intervention in seven countries. BMC Palliative Care, 2018, 17, 47.	0.8	39
35	Strategies for the implementation of palliative care education and organizational interventions in long-term care facilities: A scoping review. Palliative Medicine, 2020, 34, 558-570.	1.3	39
36	Caring for a dying spouse at the end of life: †lt's one of the things you volunteer for when you get married': a qualitative study of the oldest carers' experiences. Age and Ageing, 2016, 45, 421-426.	0.7	37

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37	Policy analysis: Palliative care in Ireland. Health Policy, 2014, 115, 68-74.	1.4	36
38	Supportive, palliative, and end-of-life care for patients with cancer in Asia: resource-stratified guidelines from the Asian Oncology Summit 2012. Lancet Oncology, The, 2012, 13, e492-e500.	5.1	34
39	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. BMC Medicine, 2015, 13, 263.	2.3	34
40	How effective are volunteers at supporting people in their last year of life? A pragmatic randomised wait-list trial in palliative care (ELSA). BMC Medicine, 2016, 14, 203.	2.3	34
41	Palliative Care Implementation in Long-Term Care Facilities: European Association for Palliative Care White Paper. Journal of the American Medical Directors Association, 2020, 21, 1051-1057.	1.2	34
42	Case study research methods in end-of-life care: reflections on three studies. Journal of Advanced Nursing, 2007, 58, 236-245.	1.5	33
43	Family members' experiences of assisted dying: A systematic literature review with thematic synthesis. Palliative Medicine, 2019, 33, 1091-1105.	1.3	33
44	Older Chinese people's views on food: implications for supportive cancer care. Ethnicity and Health, 2008, 13, 497-514.	1.5	30
45	Advance directives in European long-term care facilities: a cross-sectional survey. BMJ Supportive and Palliative Care, 2022, 12, e393-e402.	0.8	30
46	Revised recommendations on standards and norms for palliative care in Europe from the European Association for Palliative Care (EAPC): A Delphi study. Palliative Medicine, 2022, 36, 680-697.	1.3	30
47	Building a taxonomy of integrated palliative care initiatives: results from a focus group. BMJ Supportive and Palliative Care, 2016, 6, 14-20.	0.8	29
48	Unpacking the impact of older adults' home death on family care-givers' experiences of home. Health and Place, 2016, 38, 103-111.	1.5	29
49	â€~Being with' or â€~doing for'? How the role of an end-of-life volunteer befriender can impact patient wellbeing: interviews from a multiple qualitative case study (ELSA). Supportive Care in Cancer, 2018, 26, 3163-3172.	1.0	29
50	Enhancing integrated palliative care: what models are appropriate? A cross-case analysis. BMC Palliative Care, 2017, 16, 64.	0.8	27
51	How to Measure the International Development of Palliative Care? A Critique and Discussion of Current Approaches. Journal of Pain and Symptom Management, 2014, 47, 154-165.	0.6	26
52	Evaluating the implementation of the PACE Steps to Success Programme in long-term care facilities in seven countries according to the RE-AIM framework. Implementation Science, 2019, 14, 107.	2.5	26
53	The role of health care assistants in supporting district nurses and family carers to deliver palliative care at home: findings from an evaluation project. Journal of Clinical Nursing, 2011, 20, 2043-2052.	1.4	25
54	To what degree is palliative care integrated in guidelines and pathways for adult cancer patients in Europe: a systematic literature review. BMC Palliative Care, 2016, 15, 26.	0.8	25

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55	Palliative care provision in long-term care facilities differs across Europe: Results of a cross-sectional study in six European countries (PACE). Palliative Medicine, 2019, 33, 1176-1188.	1.3	24
56	Digitisation and the patient–professional relationship in palliative care. Palliative Medicine, 2020, 34, 441-443.	1.3	24
57	Legalisation of assisted suicide: a safeguard to euthanasia?. Lancet, The, 2014, 384, 127.	6.3	23
58	Experiences of end-of-life care in community hospitals. Health and Social Care in the Community, 2007, 15, 494-501.	0.7	21
59	How to measure the effects and potential adverse events of palliative sedation? An integrative review. Palliative Medicine, 2021, 35, 295-314.	1.3	21
60	Protocol for the End-of-Life Social Action Study (ELSA): a randomised wait-list controlled trial and embedded qualitative case study evaluation assessing the causal impact of social action befriending services on end of life experience. BMC Palliative Care, 2016, 15, 60.	0.8	20
61	Identifying associations between quality of life outcomes and healthcare-related variables among colorectal cancer survivors: A cross-sectional survey study. International Journal of Nursing Studies, 2020, 101, 103434.	2.5	20
62	†You can't say, "what about me?―I'm not the one with cancer': information and support needs of relatives. Psycho-Oncology, 2015, 24, 705-711.	1.0	19
63	Recommendations on priorities for integrated palliative care: transparent expert consultation with international leaders for the InSuP-C project. BMC Palliative Care, 2019, 18, 32.	0.8	18
64	Palliative and endâ€ofâ€life educational interventions for staff working in longâ€ŧerm care facilities: An integrative review of the literature. International Journal of Older People Nursing, 2021, 16, e12347.	0.6	18
65	Towards better support for family carers: A richer understanding. Palliative Medicine, 2013, 27, 579-580.	1.3	17
66	The cost of survival: an exploration of colorectal cancer survivors' experiences of pain. Acta Oncológica, 2017, 56, 205-211.	0.8	17
67	The perspectives of bereaved family carers on dying at home: the study protocol of â€~unpacking the home: family carers' reflections on dying at home. BMC Palliative Care, 2012, 11, 23.	0.8	16
68	Research, recruitment and observational data collection in care homes: lessons from the PACE study. BMC Research Notes, 2019, 12, 508.	0.6	16
69	"You get old, you get breathless, and you die― Chronic obstructive pulmonary disease in Barnsley, UK Health and Place, 2012, 18, 1396-1403.	1.5	15
70	Length of stay in long-term care facilities: a comparison of residents in six European countries. Results of the PACE cross-sectional study. BMJ Open, 2020, 10, e033881.	0.8	15
71	Undertaking Research Using Online Nominal Group Technique: Lessons from an International Study (RESPACC). Journal of Palliative Medicine, 2021, 24, 1867-1871.	0.6	15
72	A Systematic Content Analysis of Policy Barriers Impeding Access to Opioid Medication in Central and Eastern Europe: Results of ATOME. Journal of Pain and Symptom Management, 2016, 51, 99-107.	0.6	14

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73	Benefits and challenges of collaborative research: lessons from supportive and palliative care. BMJ Supportive and Palliative Care, 2011, 1, 5-11.	0.8	13
74	Legal Barriers in Accessing Opioid Medicines: Results of the ATOME Quick Scan of National Legislation of Eastern European Countries. Journal of Pain and Symptom Management, 2014, 48, 1135-1144.	0.6	13
75	Physical restraining of nursing home residents in the last week of life: An epidemiological study in six European countries. International Journal of Nursing Studies, 2020, 104, 103511.	2.5	13
76	What it means to be a palliative care volunteer in eight European countries: a qualitative analysis of accounts of volunteering. Scandinavian Journal of Caring Sciences, 2021, 35, 170-177.	1.0	13
77	Improving regional care in the last year of life by setting up a pragmatic evidence-based Plan–Do–Study–Act cycle: results from a cross-sectional survey. BMJ Open, 2020, 10, e035988.	0.8	13
78	Coping strategies of patients with advanced lung or colorectal cancer in six European countries: Insights from the ACTION Study. Psycho-Oncology, 2020, 29, 347-355.	1.0	12
79	Colorectal cancer survivors' quality of life: a qualitative study of unmet need. BMJ Supportive and Palliative Care, 2020, , bmjspcare-2020-002190.	0.8	12
80	Review of European Guidelines on Palliative Sedation: A Foundation for the Updating of the European Association for Palliative Care Framework. Journal of Palliative Medicine, 2022, 25, 1721-1731.	0.6	12
81	The roles, responsibilities and practices of healthcare assistants in out-of-hours community palliative care: A systematic scoping review. Palliative Medicine, 2020, 34, 976-988.	1.3	11
82	Opioid underuse in terminal care of long-term care facility residents with pain and/or dyspnoea: A cross-sectional PACE-survey in six European countries. Palliative Medicine, 2020, 34, 784-794.	1.3	11
83	Prevalence vs impact: a mixed methods study of survivorship issues in colorectal cancer. Quality of Life Research, 2022, 31, 1117-1134.	1.5	11
84	Prior Study of Cross-Cultural Validation of McGill Quality-of-Life Questionnaire in Mainland Mandarin Chinese Patients With Cancer. American Journal of Hospice and Palliative Medicine, 2015, 32, 709-714.	0.8	10
85	Moving the integration of palliative care from idea to practice. Palliative Medicine, 2016, 30, 197-199.	1.3	10
86	Family carers' experiences of coping with the deaths of adults in home settings: A narrative analysis of carers' relevant background worries. Palliative Medicine, 2018, 32, 950-959.	1.3	9
87	Action learning: an effective way to improve cancerâ€related pain management. Journal of Clinical Nursing, 2017, 26, 3430-3441.	1.4	8
88	Integrated palliative care networks from the perspectives of patients: A cross-sectional explorative study in five European countries. Palliative Medicine, 2018, 32, 1103-1113.	1.3	8
89	Swiss Families' Experiences of Interactions with Providers during Assisted Suicide: A Secondary Data Analysis of an Interview Study. Journal of Palliative Medicine, 2020, 23, 506-512.	0.6	8
90	Potentially Inappropriate Treatments at the End of Life in Nursing Home Residents: Findings From the PACE Cross-Sectional Study in Six European Countries. Journal of Pain and Symptom Management, 2021, 61, 732-742.e1.	0.6	8

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91	An evaluation of research capacity building from the Cancer Experiences Collaborative. BMJ Supportive and Palliative Care, 2012, 2, 280-285.	0.8	7
92	Quality of Life Trends in People With and Without Cancer Referred to Volunteer-Provided Palliative Care Services (ELSA): A Longitudinal Study. Journal of Pain and Symptom Management, 2018, 56, 689-698.	0.6	7
93	A framework for cross-cultural development and implementation of complex interventions to improve palliative care in nursing homes: the PACE steps to success programme. BMC Health Services Research, 2019, 19, 745.	0.9	7
94	Perception of the Quality of Communication With Physicians Among Relatives of Dying Residents of Long-term Care Facilities in 6 European Countries: PACE Cross-Sectional Study. Journal of the American Medical Directors Association, 2020, 21, 331-337.	1.2	7
95	Ethical and research governance approval across Europe: Experiences from three European palliative care studies. Palliative Medicine, 2020, 34, 817-821.	1.3	7
96	Associations between Length of Stay in Long Term Care Facilities and End of Life Care. Analysis of the PACE Cross-Sectional Study. International Journal of Environmental Research and Public Health, 2020, 17, 2742.	1.2	7
97	The Decision-Making Process for Palliative Sedation for Patients with Advanced Cancer–Analysis from a Systematic Review of Prospective Studies. Cancers, 2022, 14, 301.	1.7	7
98	The impact of covid-19 on out-of-hours adult hospice care: an online survey. BMC Palliative Care, 2022, 21, .	0.8	7
99	A sociological commentary on the refusal of treatment by patients with cancer. Mortality, 2009, 14, 309-324.	0.3	6
100	Pain Intervention for people with Dementia in nursing homes (PID): study protocol for a quasi-experimental nurse intervention. BMC Palliative Care, 2017, 16, 27.	0.8	6
101	European Palliative Sedation Project. Journal of Palliative Medicine, 2020, 23, 154-155.	0.6	6
102	Lack of Effect of a Multicomponent Palliative Care Program for Nursing Home Residents on Hospital Use in the Last Month of Life and on Place of Death: A Secondary Analysis of a Multicountry Cluster Randomized Control Trial. Journal of the American Medical Directors Association, 2020, 21, 1973-1978.e2.	1.2	6
103	Is the content of guidelines/pathways a barrier for the integration of palliative Care in Chronic Heart Failure (CHF) and chronic pulmonary obstructive disease (COPD)? A comparison with the case of cancer in Europe. BMC Palliative Care, 2017, 16, 62.	0.8	5
104	Do patients with lung cancer recall physician-initiated discussions about planning for end-of-life care following disclosure of a terminal prognosis?. BMJ Supportive and Palliative Care, 2019, 9, 197-201.	0.8	5
105	MyPal: Designing and Evaluating Digital Patient-Reported Outcome Systems for Cancer Palliative Care in Europe. Journal of Palliative Medicine, 2021, 24, 962-964.	0.6	5
106	Family carers research: What progress has been made?. Palliative Medicine, 2022, , 026921632110378.	1.3	5
107	Agreement of Nursing Home Staff With Palliative Care Principles: A PACE Cross-sectional Study Among Nurses and Care Assistants in Five European Countries. Journal of Pain and Symptom Management, 2019, 58, 824-834.	0.6	4
108	The perception of barriers concerning opioid medicines: A survey examining differences between policy makers, healthcare professionals and other stakeholders. Palliative Medicine, 2020, 34, 493-503.	1.3	4

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109	Understanding the Outcomes of Supplementary Support Services in Palliative Care for Older People. A Scoping Review and Mapping Exercise. Journal of Pain and Symptom Management, 2020, 60, 449-459.e21.	0.6	4
110	No difference in effects of â€~PACE steps to success' palliative care program for nursing home residents with and without dementia: a pre-planned subgroup analysis of the seven-country PACE trial. BMC Palliative Care, 2021, 20, 39.	0.8	4
111	Large differences in the organization of palliative care in nursing homes in six European countries: findings from the PACE cross-sectional study. BMC Palliative Care, 2021, 20, 131.	0.8	4
112	The study protocol of: †Initiating end of life care in stroke: clinical decision-making around prognosis'. BMC Palliative Care, 2014, 13, 55.	0.8	3
113	Assessing Emotional Functioning with Increased Power: Relative Validity of a Customized EORTC Short Form in the International ACTION Trial. Value in Health, 2019, 22, 92-98.	0.1	3
114	What should we report? Lessons learnt from the development and implementation of serious adverse event reporting procedures in non-pharmacological trials in palliative care. BMC Palliative Care, 2021, 20, 19.	0.8	3
115	The crossover artist: consensus methods in health research. , 2020, , .		3
116	Ethical Principles in Digital Palliative Care for Children: The MyPal Project and Experiences Made in Designing a Trustworthy Approach. Frontiers in Digital Health, 2022, 4, 730430.	1.5	3
117	Instruments to assess the burden of care for family caregivers of adult palliative care patients. International Journal of Palliative Nursing, 2022, 28, 80-99.	0.2	3
118	Nursing and palliative care – better practices within the team. Progress in Palliative Care, 2010, 18, 328-329.	0.7	2
119	The European Association for Palliative Care: forging a vision of excellence in palliative care. Progress in Palliative Care, 2012, 20, 223-226.	0.7	2
120	Recruitment into palliative care research studies: a common challenge. Progress in Palliative Care, 2013, 21, 263-264.	0.7	2
121	Unaddressed biases in reported prevalence of continuous deep sedation until death. Journal of Clinical Epidemiology, 2015, 68, 1241-1242.	2.4	2
122	Factors Associated with Perception of the Quality of Physicians' End-of-life Communication in Long-Term Care Facilities: PACE Cross-Sectional Study. Journal of the American Medical Directors Association, 2020, 21, 439.e1-439.e8.	1.2	2
123	Disseminating research findings using a massive online open course for maximising impact and developing recommendations for practice. BMC Palliative Care, 2020, 19, 54.	0.8	2
124	Exploring a New Model of End-of-Life Care for Older People That Operates in the Space Between the Life World and the Healthcare System: A Qualitative Case Study. International Journal of Health Policy and Management, 2020, 9, 344-351.	0.5	2
125	Integrating Palliative Care by Virtue of Diplomacy; A Cross-sectional Group Interview Study of the Roles and Attitudes of Palliative Care Professionals to Further Integrate Palliative Care in Europe. International Journal of Health Policy and Management, 2020, , .	0.5	1
126	Advance care planning—family carer psychological distress and involvement in decision making: the ACTION trial. BMJ Supportive and Palliative Care, 2023, 13, e807-e811.	0.8	1

#	Article	IF	CITATIONS
127	An exploration of the views of staff on cultural aspects of end-of-life care in Japanese long-term care facilities: a qualitative study. Ageing and Society, 0, , 1-23.	1.2	1
128	Following bereavement, poor health is more likely in carers who perceived that their support from health services was insufficient or whose family member did not die in the carer's preferred place of death. Evidence-based Nursing, 2011, 13, 94-95.	0.1	0
129	Self-reported knowledge, attitudes, and behaviour towards hospice care and how are these related to training in palliative care: An online survey among oncologists in the Czech Republic and Slovakia. Progress in Palliative Care, 2015, 23, 1-8.	0.7	0
130	Protocol for a mixed methods exploratory investigation into the role and contribution of the healthcare assistant in out-of-hours palliative care. BMC Nursing, 2021, 20, 55.	0.9	0
131	Developing a decision support tool for responding to patients' reported levels of information needs, family anxiety, depression, and breathlessness Journal of Clinical Oncology, 2014, 32, 173-173.	0.8	0
132	Palliative Care for Older People. , 2019, , 8-8.		0
133	Title is missing!. , 2020, 17, e1003422.		0
134	Title is missing!. , 2020, 17, e1003422.		0
135	Title is missing!. , 2020, 17, e1003422.		0
136	Title is missing!. , 2020, 17, e1003422.		0
137	Title is missing!. , 2020, 17, e1003422.		0
138	Title is missing!. , 2020, 17, e1003422.		0
139	Title is missing!. , 2020, 17, e1003422.		0
140	Information and support: implications for developing palliative care programs. Annals of Palliative Medicine, 2013, 2, 43-4.	0.5	0
141	International Leadership in Palliative Care: New Leaders Required. Journal of Palliative Medicine, 2022, 25, 696-697.	0.6	Ο