Richard Harding

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2228746/publications.pdf

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332 papers 12,403 citations

54 h-index 91 g-index

338 all docs 338 docs citations

338 times ranked

9636 citing authors

#	Article	IF	CITATIONS
1	Preferences for place of death if faced with advanced cancer: a population survey in England, Flanders, Germany, Italy, the Netherlands, Portugal and Spain. Annals of Oncology, 2012, 23, 2006-2015.	1.2	430
2	Redefining Palliative Careâ€"A New Consensus-Based Definition. Journal of Pain and Symptom Management, 2020, 60, 754-764.	1.2	423
3	The escalating global burden of serious health-related suffering: projections to 2060 by world regions, age groups, and health conditions. The Lancet Global Health, 2019, 7, e883-e892.	6.3	395
4	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.	3.1	343
5	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. Journal of Pain and Symptom Management, 2014, 48, 660-677.	1.2	322
6	Implementing patient-reported outcome measures in palliative care clinical practice: A systematic review of facilitators and barriers. Palliative Medicine, 2014, 28, 158-175.	3.1	266
7	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. BMC Medicine, 2013, 11, 111.	5 . 5	256
8	HIV and Depression – a systematic review of interventions. Psychology, Health and Medicine, 2011, 16, 493-527.	2.4	241
9	Short-form Zarit Caregiver Burden Interviews were valid in advanced conditions. Journal of Clinical Epidemiology, 2010, 63, 535-542.	5.0	230
10	Palliative care in sub-Saharan Africa. Lancet, The, 2005, 365, 1971-1977.	13.7	186
11	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". Heart, 2007, 93, 963-967.	2.9	153
12	EAPC White Paper on outcome measurement in palliative care: Improving practice, attaining outcomes and delivering quality services – Recommendations from the European Association for Palliative Care (EAPC) Task Force on Outcome Measurement. Palliative Medicine, 2016, 30, 6-22.	3.1	149
13	Experience of persistent psychological symptoms and perceived stigma among people with HIV on antiretroviral therapy (ART): A systematic review. International Journal of Nursing Studies, 2014, 51, 1171-1189.	5.6	146
14	Validation of a core outcome measure for palliative care in Africa: the APCA African Palliative Outcome Scale. Health and Quality of Life Outcomes, 2010, 8, 10.	2.4	143
15	How can informal caregivers in cancer and palliative care be supported? An updated systematic literature review of interventions and their effectiveness. Palliative Medicine, 2012, 26, 7-22.	3.1	135
16	Benzodiazepines for the relief of breathlessness in advanced malignant and non-malignant diseases in adults. The Cochrane Library, 2016, 2016, CD007354.	2.8	133
17	Meeting the Communication and Information Needs of Chronic Heart Failure Patients. Journal of Pain and Symptom Management, 2008, 36, 149-156.	1.2	131
18	Measuring quality of life among people living with HIV: a systematic review of reviews. Health and Quality of Life Outcomes, 2017, 15, 220.	2.4	130

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19	Advancing Palliative Care as a Human Right. Journal of Pain and Symptom Management, 2009, 38, 767-774.	1.2	128
20	HIV infection associated post-traumatic stress disorder and post-traumatic growth – A systematic review. Psychology, Health and Medicine, 2011, 16, 612-629.	2.4	117
21	Research into palliative care in sub-Saharan Africa. Lancet Oncology, The, 2013, 14, e183-e188.	10.7	113
22	Does palliative care improve outcomes for patients with HIV/AIDS? A systematic review of the evidence. Sexually Transmitted Infections, 2005, 81, 5-14.	1.9	112
23	The Measurement of Spirituality in Palliative Care and the Content of Tools Validated Cross-Culturally: A Systematic Review. Journal of Pain and Symptom Management, 2011, 41, 728-753.	1.2	111
24	Benzodiazepines for the relief of breathlessness in advanced malignant and non-malignant diseases in adults. , 2010, , CD007354.		105
25	Implementing patient reported outcome measures (PROMs) in palliative care - users' cry for help. Health and Quality of Life Outcomes, 2011, 9, 27.	2.4	104
26	Needs, Experiences, and Preferences of Sexual Minorities for End-of-Life Care and Palliative Care: A Systematic Review. Journal of Palliative Medicine, 2012, 15, 602-611.	1.1	101
27	Culture and End of Life Care: A Scoping Exercise in Seven European Countries. PLoS ONE, 2012, 7, e34188.	2.5	97
28	Working with ambivalence: informal caregivers of patients at the end of life. Supportive Care in Cancer, 2001, 9, 642-645.	2.2	96
29	The prevalence and burden of symptoms amongst cancer patients attending palliative care in two African countries. European Journal of Cancer, 2011, 47, 51-56.	2.8	95
30	Recommendations to reduce inequalities for LGBT people facing advanced illness: ACCESSCare national qualitative interview study. Palliative Medicine, 2018, 32, 23-35.	3.1	88
31	Suicidal ideation in UK HIV clinic attenders. Aids, 2008, 22, 1651-1658.	2.2	86
32	Priorities for treatment, care and information if faced with serious illness: A comparative population-based survey in seven European countries. Palliative Medicine, 2014, 28, 101-110.	3.1	86
33	Symptoms are highly prevalent among HIV outpatients and associated with poor adherence and unprotected sexual intercourse. Sexually Transmitted Infections, 2010, 86, 520-524.	1.9	81
34	End-of-life care across Southern Europe: A critical review of cultural similarities and differences between Italy, Spain and Portugal. Critical Reviews in Oncology/Hematology, 2012, 82, 387-401.	4.4	80
35	Comparative Analysis of Informal Caregiver Burden in Advanced Cancer, Dementia, and Acquired Brain Injury. Journal of Pain and Symptom Management, 2015, 50, 445-452.	1.2	79
36	Patient-reported outcomes to enhance person-centred HIV care. Lancet HIV, the, 2020, 7, e59-e68.	4.7	79

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37	Is antiretroviral therapy associated with symptom prevalence and burden?. International Journal of STD and AIDS, 2006, 17, 400-405.	1.1	78
38	The bereavement experiences of lesbian, gay, bisexual and/or trans* people who have lost a partner: A systematic review, thematic synthesis and modelling of the literature. Palliative Medicine, 2016, 30, 730-744.	3.1	77
39	What are the perceived needs and challenges of informal caregivers in home cancer palliative care? Qualitative data to construct a feasible psycho-educational intervention. Supportive Care in Cancer, 2012, 20, 1975-1982.	2.2	73
40	â€~Best practice' in developing and evaluating palliative and end-of-life care services: A meta-synthesis of research methods for the MORECare project. Palliative Medicine, 2013, 27, 885-898.	3.1	73
41	Evaluation of a short-term group intervention for informal carers of patients attending a home palliative care service. Journal of Pain and Symptom Management, 2004, 27, 396-408.	1.2	69
42	Current HIV/AIDS end-of-life care in sub-Saharan Africa: a survey of models, services, challenges and priorities. BMC Public Health, 2003, 3, 33.	2.9	68
43	Development of the APCA African Palliative Outcome Scale. Journal of Pain and Symptom Management, 2007, 33, 229-232.	1.2	67
44	Consensus statement on the role of health systems in advancing the long-term well-being of people living with HIV. Nature Communications, 2021, 12, 4450.	12.8	67
45	â€~My body's falling apart.' Understanding the experiences of patients with advanced multimorbidity to improve care: serial interviews with patients and carers. BMJ Supportive and Palliative Care, 2016, 6, 60-65.	1.6	66
46	Generating an African Palliative Care Evidence Base: The Context, Need, Challenges, and Strategies. Journal of Pain and Symptom Management, 2008, 36, 304-309.	1.2	65
47	Adherence to antiretroviral treatment in patients with HIV in the UK: a study of complexity. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2008, 20, 442-448.	1.2	64
48	Quality of life among patients receiving palliative care in South Africa and Uganda: a multi-centred study. Health and Quality of Life Outcomes, 2011, 9, 21.	2.4	64
49	A critical review of advance directives in Germany: Attitudes, use and healthcare professionals' compliance. Patient Education and Counseling, 2012, 87, 277-288.	2.2	64
50	The relationship between patient characteristics and carer psychological status in home palliative cancer care. Supportive Care in Cancer, 2003, 11, 638-643.	2.2	62
51	An evaluation of a morphine public health programme for cancer and AIDS pain relief in Sub-Saharan Africa. BMC Public Health, 2005, 5, 82.	2.9	62
52	Meeting information needs of patients with incurable progressive disease and their families in South Africa and Uganda: multicentre qualitative study. BMJ: British Medical Journal, 2009, 338, b1326-b1326.	2.3	62
53	Prevalence, Burden, and Correlates of Physical and Psychological Symptoms Among HIV Palliative Care Patients in Sub-Saharan Africa: An International Multicenter Study. Journal of Pain and Symptom Management, 2012, 44, 1-9.	1.2	62
54	A systematic review of Interventions for anxiety in people with HIV. Psychology, Health and Medicine, 2011, 16, 528-547.	2.4	59

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55	Needs Assessments in Palliative Care: An Appraisal of Definitions and Approaches Used. Journal of Pain and Symptom Management, 2007, 33, 500-505.	1.2	57
56	Systematic Review of the Primary Research on Minority Ethnic Groups and End-of-Life Care From the United Kingdom. Journal of Pain and Symptom Management, 2012, 43, 261-286.	1.2	57
57	Provision of Pain- and Symptom-Relieving Drugs for HIV/AIDS in Sub-Saharan Africa. Journal of Pain and Symptom Management, 2010, 40, 405-415.	1.2	56
58	Pain Control in the African Context: the Ugandan introduction of affordable morphine to relieve suffering at the end of life. Philosophy, Ethics, and Humanities in Medicine, 2010, 5, 10.	1.5	56
59	Service Delivery Models to Maximize Quality of Life for Older People at the End of Life: A Rapid Review. Milbank Quarterly, 2019, 97, 113-175.	4.4	55
60	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.	2.8	54
61	Symptom Burden in HIV-Infected Adults at Time of HIV Diagnosis in Rural Uganda. Journal of Palliative Medicine, 2010, 13, 375-380.	1.1	53
62	End-of-life care in general practice: A cross-sectional, retrospective survey of †cancerâ€, †organ failure†and †old-age/dementia†patients. Palliative Medicine, 2014, 28, 965-975.	3.1	53
63	Towards personâ€centred care for people living with <scp>HIV</scp> : what core outcomes matter, and how might we assess them? A crossâ€national multiâ€centre qualitative study with key stakeholders. HIV Medicine, 2019, 20, 542-554.	2.2	52
64	Multidimensional Problems Among Advanced Cancer Patients in Cuba: Awareness of Diagnosis Is Associated with Better Patient Status. Journal of Pain and Symptom Management, 2009, 37, 325-330.	1.2	51
65	Coordination of care for individuals with advanced progressive conditions: a multi-site ethnographic and serial interview study. British Journal of General Practice, 2013, 63, e580-e588.	1.4	51
66	The conceptual models and mechanisms of action that underpin advance care planning for cancer patients: A systematic review of randomised controlled trials. Palliative Medicine, 2019, 33, 5-23.	3.1	51
67	Psychological, social and spiritual distress at the end of life in heart failure patients. Current Opinion in Supportive and Palliative Care, 2007, 1, 260-266.	1.3	50
68	Symptoms and Concerns Among Children and Young People with Life-Limiting and Life-Threatening Conditions: A Systematic Review Highlighting Meaningful Health Outcomes. Patient, 2019, 12, 15-55.	2.7	50
69	Integration of palliative care throughout HIV disease. Lancet Infectious Diseases, The, 2012, 12, 571-575.	9.1	49
70	A Hospital-Based Palliative Care Service for Patients With Advanced Organ Failure in Sub-Saharan Africa Reduces Admissions and Increases Home Death Rates. Journal of Pain and Symptom Management, 2014, 47, 786-792.	1.2	47
71	Pain in people living with HIV and its association with healthcare resource use, well being and functional status. Aids, 2018, 32, 2697-2706.	2.2	47
72	Brief Report: The Association of Chronic Pain and Long-Term Opioid Therapy With HIV Treatment Outcomes. Journal of Acquired Immune Deficiency Syndromes (1999), 2018, 79, 77-82.	2.1	47

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73	What Palliative Care-Related Problems Do Patients Experience at HIV Diagnosis? A Systematic Review of the Evidence. Journal of Pain and Symptom Management, 2011, 42, 734-753.	1.2	46
74	Pediatric Palliative Care in Sub-Saharan Africa: A Systematic Review of the Evidence for Care Models, Interventions, and Outcomes. Journal of Pain and Symptom Management, 2014, 47, 642-651.	1.2	46
75	Children's palliative care in low- and middle-income countries. Archives of Disease in Childhood, 2016, 101, 85-90.	1.9	46
76	Enhancing validity, reliability and participation in self-reported health outcome measurement for children and young people: a systematic review of recall period, response scale format, and administration modality. Quality of Life Research, 2021, 30, 1803-1832.	3.1	46
77	Improving HIV management in sub-Saharan Africa: How much palliative care is needed?. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2007, 19, 1304-1306.	1.2	45
78	Physical and Psychological Symptoms and Risk of Virologic Rebound Among Patients With Virologic Suppression on Antiretroviral Therapy. Journal of Acquired Immune Deficiency Syndromes (1999), 2010, 54, 500-505.	2.1	45
79	What factors are associated with patient self-reported health status among HIV outpatients? A multi-centre UK study of biomedical and psychosocial factors. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2012, 24, 963-971.	1.2	45
80	Nurse-led palliative care for HIV-positive patients taking antiretroviral therapy in Kenya: a randomised controlled trial. Lancet HIV,the, 2015, 2, e328-e334.	4.7	45
81	The PRISMA Symposium 1: Outcome Tool Use. Disharmony in European Outcomes Research for Palliative and Advanced Disease Care: Too Many Tools in Practice. Journal of Pain and Symptom Management, 2011, 42, 493-500.	1.2	43
82	Pain Among Ambulatory HIV/AIDS Patients: Multicenter Study of Prevalence, Intensity, Associated Factors, and Effect. Journal of Pain, 2012, 13, 704-713.	1.4	43
83	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. Palliative Medicine, 2014, 28, 293-301.	3.1	43
84	Modelling services to meet the palliative care needs of chronic heart failure patients and their families: current practice in the UK. Palliative Medicine, 2007, 21, 385-390.	3.1	42
85	To be involved or not to be involved: A survey of public preferences for self-involvement in decision-making involving mental capacity (competency) within Europe. Palliative Medicine, 2013, 27, 418-427.	3.1	41
86	Public preferences and priorities for end-of-life care in Kenya: a population-based street survey. BMC Palliative Care, 2014, 13, 4.	1.8	41
87	Targeting smoking cessation to high prevalence communities: outcomes from a pilot intervention for gay men. BMC Public Health, 2004, 4, 43.	2.9	40
88	A Phase II randomised controlled trial assessing the feasibility, acceptability and potential effectiveness of Dignity Therapy for older people in care homes: Study protocol. BMC Geriatrics, 2009, 9, 9.	2.7	39
89	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. BMC Health Services Research, 2011, 11, 141.	2.2	39
90	Outcomes 'out of africa': the selection and implementation of outcome measures for palliative care in Africa. BMC Palliative Care, 2012, 11 , 1 .	1.8	38

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91	Coordination of end-of-life care for patients with lung cancer and those with advanced COPD: are there transferable lessons? A longitudinal qualitative study. Primary Care Respiratory Journal: Journal of the General Practice Airways Group, 2014, 23, 46-51.	2.3	38
92	Place of death of people living with Parkinson's disease: a population-level study in 11 countries. BMC Palliative Care, 2015, 14, 28.	1.8	38
93	Access and equity in HIV/AIDS palliative care: a review of the evidence and responses. Palliative Medicine, 2005, 19, 251-258.	3.1	37
94	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.	2.8	37
95	â€~Peace' and â€~life worthwhile' as measures of spiritual well-being in African palliative care: a mixed-methods study. Health and Quality of Life Outcomes, 2013, 11, 94.	2.4	37
96	If you had less than a year to live, would you want to know? A sevenâ€country European population survey of public preferences for disclosure of poor prognosis. Psycho-Oncology, 2013, 22, 2298-2305.	2.3	37
97	The Real-World Problem of Care Coordination: A Longitudinal Qualitative Study with Patients Living with Advanced Progressive Illness and Their Unpaid Caregivers. PLoS ONE, 2014, 9, e95523.	2.5	37
98	Palliative care in South Asia: a systematic review of the evidence for care models, interventions, and outcomes. BMC Research Notes, 2015, 8, 172.	1.4	37
99	What influences patients' decisions regarding palliative care in advance care planning discussions? Perspectives from a qualitative study conducted with advanced cancer patients, families and healthcare professionals. Palliative Medicine, 2019, 33, 1299-1309.	3.1	37
100	The empirical evidence underpinning the concept and practice of person-centred care for serious illness: a systematic review. BMJ Global Health, 2020, 5, e003330.	4.7	37
101	A Psychometric Evaluation of Measures of Spirituality Validated in Culturally Diverse Palliative Care Populations. Journal of Pain and Symptom Management, 2011, 42, 604-622.	1.2	36
102	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. BMC Palliative Care, 2009, 8, 5.	1.8	35
103	â€~Burden to others' as a public concern in advanced cancer: a comparative survey in seven European countries. BMC Cancer, 2013, 13, 105.	2.6	35
104	What are the supportive and palliative care needs of patients with cutaneous T-cell lymphoma and their caregivers? A systematic review of the evidence. British Journal of Dermatology, 2014, 170, 599-608.	1.5	35
105	Palliative Care Research in Africa: Consensus Building for a Prioritized Agenda. Journal of Pain and Symptom Management, 2014, 47, 315-324.	1.2	35
106	Home care by general practitioners for cancer patients in the last 3 months of life: An epidemiological study of quality and associated factors. Palliative Medicine, 2016, 30, 64-74.	3.1	35
107	Doctor-patient concordance during HIV treatment switching decision-making. HIV Medicine, 2011, 12, 87-96.	2.2	34
108	Stressors and Resources of Caregivers of Patients With Incurable Progressive Illness in Sub-Saharan Africa. Qualitative Health Research, 2014, 24, 317-328.	2.1	34

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109	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.	5.5	34
110	Factors Associated with Transition from Community Settings to Hospital as Place of Death for Adults Aged 75 and Older: A Populationâ€Based Mortality Followâ€Back Survey. Journal of the American Geriatrics Society, 2016, 64, 2210-2217.	2.6	34
111	The burden of serious health-related suffering among cancer decedents: Global projections study to 2060. Palliative Medicine, 2021, 35, 231-235.	3.1	34
112	'My dreams are shuttered down and it hurts lots'–a qualitative study of palliative care needs and their management by HIV outpatient services in Kenya and Uganda. BMC Palliative Care, 2013, 12, 35.	1.8	33
113	The needs, models of care, interventions and outcomes of palliative care in the Caribbean: a systematic review of the evidence. BMC Palliative Care, 2016, 15, 9.	1.8	33
114	â€They will be afraid to touch you': LGBTI people and sex workers' experiences of accessing healthcare in Zimbabweâ€"an in-depth qualitative study. BMJ Global Health, 2017, 2, e000168.	4.7	33
115	Successive switching of antiretroviral therapy is associated with high psychological and physical burden. International Journal of STD and AIDS, 2007, 18, 700-704.	1.1	32
116	Advancing research and practice in HIV and rehabilitation: a framework of research priorities in HIV, disability and rehabilitation. BMC Infectious Diseases, 2014, 14, 724.	2.9	32
117	â€We're all carrying a burden that we're not sharing': a qualitative study of the impact of cutaneous T-cell lymphoma on the family. British Journal of Dermatology, 2015, 172, 1581-1592.	1.5	32
118	Positive futures? The impact of HIV infection on achieving health, wealth and future planning. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2008, 20, 565-570.	1.2	31
119	Cultural Competence in End-of-Life Care: Terms, Definitions, and Conceptual Models from the British Literature. Journal of Palliative Medicine, 2012, 15, 812-820.	1.1	31
120	\hat{a} € We had to change to single beds because I itch in the night \hat{a} € a qualitative study of the experiences, attitudes and approaches to coping of patients with cutaneous T-cell lymphoma. British Journal of Dermatology, 2015, 173, 83-92.	1.5	31
121	Diversity in Defining End of Life Care: An Obstacle or the Way Forward?. PLoS ONE, 2013, 8, e68002.	2.5	30
122	Use of the Palliative Outcome Scale in Argentina: A Cross-Cultural Adaptation and Validation Study. Journal of Pain and Symptom Management, 2008, 35, 188-202.	1.2	29
123	Enhancing patient-reported outcome measurement in research and practice of palliative and end-of-life care. Supportive Care in Cancer, 2012, 20, 1573-1578.	2.2	29
124	Evaluation of a physiotherapy-led group rehabilitation intervention for adults living with HIV: referrals, adherence and outcomes. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2016, 28, 1495-1505.	1.2	29
125	Provision of palliative care for life-limiting disease in a low income country national hospital setting: how much is needed?. BMJ Supportive and Palliative Care, 2012, 2, 140-144.	1.6	28
126	The Prevalence and Associated Distress of Physical and Psychological Symptoms in Patients With Advanced Heart Failure Attending a South African Medical Center. Journal of Cardiovascular Nursing, 2016, 31, 313-322.	1.1	28

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127	Are we heading in the same direction? European and African doctors' and nurses' views and experiences regarding outcome measurement in palliative care. Palliative Medicine, 2012, 26, 242-249.	3.1	27
128	Culture Is a Priority for Research in End-of-Life Care in Europe: A Research Agenda. Journal of Pain and Symptom Management, 2012, 44, 285-294.	1.2	27
129	The "Spirit 8―successfully captured spiritual well-being in African palliative care: factor and Rasch analysis. Journal of Clinical Epidemiology, 2012, 65, 434-443.	5.0	27
130	Inclusion of end-of-life care in the global health agenda. The Lancet Global Health, 2014, 2, e375-e376.	6.3	27
131	Palliative Care in the Greater China Region: A Systematic Review of Needs, Models, and Outcomes. Journal of Pain and Symptom Management, 2021, 61, 585-612.	1.2	27
132	Palliative and end-of-life care in the global response to multidrug-resistant tuberculosis. Lancet Infectious Diseases, The, 2012, 12, 643-646.	9.1	26
133	Choosing care homes as the least preferred place to die: a cross-national survey of public preferences in seven European countries. BMC Palliative Care, 2014, 13, 48.	1.8	26
134	What will it really take to end the HIV epidemic?. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2018, 30, 1-4.	1.2	26
135	Cost of Medical Care of Patients with Advanced Serious Illness in Singapore (COMPASS): prospective cohort study protocol. BMC Cancer, 2018, 18, 459.	2.6	26
136	Multilevel model of stigma and barriers to cancer palliative care in India: a qualitative study. BMJ Open, 2019, 9, e024248.	1.9	26
137	A Multi-Professional Short-Term Group Intervention for Informal Caregivers of Patients Using a Home Palliative Care Service. Journal of Palliative Care, 2002, 18, 275-281.	1.0	25
138	How to Analyze Palliative Care Outcome Data for Patients in Sub-Saharan Africa: An International, Multicenter, Factor Analytic Examination of the APCA African POS. Journal of Pain and Symptom Management, 2013, 45, 746-752.	1.2	25
139	Traditional Healers' Views of the Required Processes for a "Good Death―Among Xhosa Patients Preand Post-Death. Journal of Pain and Symptom Management, 2013, 46, 386-394.	1.2	25
140	Palliative care as an essential component of the HIV care continuum. Lancet HIV, the, 2018, 5, e524-e530.	4.7	25
141	The Role of Palliative Care in Burns: A Scoping Review. Journal of Pain and Symptom Management, 2020, 59, 1089-1108.	1.2	25
142	Financial difficulties are associated with greater total pain and suffering among patients with advanced cancer: results from the COMPASS study. Supportive Care in Cancer, 2020, 28, 3781-3789.	2.2	25
143	Preferences and Experiences of Muslim Patients and Their Families in Muslim-Majority Countries for End-of-Life Care: A Systematic Review and Thematic Analysis. Journal of Pain and Symptom Management, 2020, 60, 1223-1238.e4.	1,2	25
144	Outcome measurement in paediatric palliative care: lessons from the past and future developments. Annals of Palliative Medicine, 2018, 7, S151-S163.	1,2	25

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145	What is the evidence for effectiveness of interventions to enhance coping among people living with HIV disease? A systematic review. Psychology, Health and Medicine, 2011, 16, 564-587.	2.4	24
146	Adjusting to the caregiving role: the importance of coping and support. International Journal of Palliative Nursing, 2012, 18, 541-545.	0.5	24
147	Public Priorities and Preferences for End-of-Life Care in Namibia. Journal of Pain and Symptom Management, 2014, 47, 620-630.	1.2	24
148	Behavioral Surveillance Study: Sexual Risk Taking Behaviour in UK HIV Outpatient Attendees. AIDS and Behavior, 2012, 16, 1708-1715.	2.7	23
149	Models of delivering palliative and end-of-life care in sub-Saharan Africa. Current Opinion in Supportive and Palliative Care, 2013, 7, 223-228.	1.3	23
150	Can palliative care integrated within HIV outpatient settings improve pain and symptom control in a low-income country? A prospective, longitudinal, controlled intervention evaluation. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2013, 25, 795-804.	1.2	23
151	Out of the shadows: non-communicable diseases and palliative care in Africa. BMJ Supportive and Palliative Care, 2017, 7, 128-132.	1.6	23
152	Evaluation of the psychometric properties of the HIV Disability Questionnaire among adults living with HIV in the United Kingdom: A cross-sectional self-report measurement study. PLoS ONE, 2019, 14, e0213222.	2.5	22
153	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. BMC Infectious Diseases, 2012, 12, 288.	2.9	21
154	A pan-European survey of research in end-of-life cancer care. Supportive Care in Cancer, 2012, 20, 39-48.	2.2	21
155	Research priorities for rehabilitation and aging with HIV: a framework from the Canada-International HIV and Rehabilitation Research Collaborative (CIHRRC). AIDS Research and Therapy, 2020, 17, 21.	1.7	21
156	The development and cognitive testing of the positive outcomes HIV PROM: a brief novel patient-reported outcome measure for adults living with HIV. Health and Quality of Life Outcomes, 2020, 18, 214.	2.4	21
157	Pain and Symptom Control in HIV Disease: Under-Researched and Poorly Managed. Clinical Infectious Diseases, 2005, 40, 491-492.	5.8	20
158	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.	1,2	20
159	The MVQOLI successfully captured quality of life in African palliative care: a factor analysis. Journal of Clinical Epidemiology, 2011, 64, 913-924.	5.0	20
160	Self-report measurement of pain & Damp; symptoms in palliative care patients: a comparison of verbal, visual and hand scoring methods in Sub-Saharan Africa. Health and Quality of Life Outcomes, 2014, 12, 118.	2.4	20
161	Conversations Worth Having: The Perceived Relevance of Advance Care Planning among Teachers, Hospice Staff, and Pastors in Knysna, South Africa. Journal of Palliative Medicine, 2013, 16, 762-767.	1.1	19
162	Understanding data and information needs for palliative cancer care to inform digital health intervention development in Nigeria, Uganda and Zimbabwe: protocol for a multicountry qualitative study. BMJ Open, 2019, 9, e032166.	1.9	19

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