Joachim Cohen

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5853304/publications.pdf

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211 papers

8,321 citations

45 h-index 80 g-index

218 all docs

218 docs citations

times ranked

218

5631 citing authors

#	Article	IF	CITATIONS
1	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
2	Increased legalisation of medical assistance in dying: relationship to palliative care. BMJ Supportive and Palliative Care, 2023, 13, 178-180.	0.8	4
3	End-of-life decisions in neonates and infants: a population-level mortality follow-back study. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2022, 107, 340-341.	1.4	3
4	How are family caregivers of people with a serious illness supported by healthcare professionals in their caregiving tasks? A cross-sectional survey of bereaved family caregivers. Palliative Medicine, 2022, 36, 529-539.	1.3	4
5	Barriers and facilitators for parents in end-of-life decision-making for neonates at the Neonatal Intensive Care Unit: A qualitative study. Palliative Medicine, 2022, 36, 730-741.	1.3	4
6	Civic engagement in serious illness, death, and loss: A systematic mixed-methods review. Palliative Medicine, 2022, 36, 625-651.	1.3	13
7	Area-Based Compassionate Communities: A systematic integrative review of existing initiatives worldwide. Palliative Medicine, 2022, 36, 422-442.	1.3	13
8	Engagement of specialized palliative care services with the general public: A population-level survey in three European countries. Palliative Medicine, 2022, , 026921632210795.	1.3	2
9	Use, usability, and impact of a card-based conversation tool to support communication about end-of-life preferences in residential elder care – a qualitative study of staff experiences. BMC Geriatrics, 2022, 22, 274.	1.1	3
10	A multi-stage process to develop quality indicators for community-based palliative care using interRAI data. PLoS ONE, 2022, 17, e0266569.	1.1	4
11	End-of-life decisions in neonates and infants: a nationwide mortality follow-back survey. BMJ Supportive and Palliative Care, 2022, , bmjspcare-2021-003357.	0.8	2
12	Physicians' Experiences and Perceptions of Environmental Factors Affecting Their Practices of Continuous Deep Sedation until Death: A Secondary Qualitative Analysis of an Interview Study. International Journal of Environmental Research and Public Health, 2022, 19, 5472.	1.2	0
13	Place of death and associated factors in 12 Latin American countries: A total population study using death certificate data. Journal of Global Health, 2022, 12, 04031.	1.2	6
14	Healthcare interventions improving and reducing quality of life in children at the end of life: a systematic review. Pediatric Research, 2021, 89, 1065-1077.	1.1	9
15	Assisted dying around the world: a status quaestionis. Annals of Palliative Medicine, 2021, 10, 3540-3553.	0.5	87
16	Are family carers part of the care team providing end-of-life care? A qualitative interview study on the collaboration between family and professional carers. Palliative Medicine, 2021, 35, 109-119.	1.3	25
17	Pilot Study to Develop and Test Palliative Care Quality Indicators for Nursing Homes. International Journal of Environmental Research and Public Health, 2021, 18, 829.	1.2	4
18	Nationwide evaluation of palliative care (Q-PAC study) provided by specialized palliative care teams using quality indicators: Large variations in quality of care. Palliative Medicine, 2021, 35, 1525-1541.	1.3	2

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19	Continuums of Change in a Competence-Building Initiative Addressing End-of-Life Communication in Swedish Elder Care. Qualitative Health Research, 2021, 31, 1904-1917.	1.0	5
20	Improved quality of care and reduced healthcare costs at the end-of-life among older people with dementia who received palliative home care: A nationwide propensity score-matched decedent cohort study. Palliative Medicine, 2021, 35, 1701-1712.	1.3	10
21	The Online Representation of Palliative Care by Practice, Policy, and Advocacy Organizations: Definitional Variations and Discursive Tensions. Qualitative Health Research, 2021, 31, 104973232110438.	1.0	1
22	Control Measures for Continuous Deep Sedation Until Death: A Framing Analysis of the Views of Physicians. Qualitative Health Research, 2021, 31, 2390-2402.	1.0	1
23	Prognosis does not change the landscape: palliative home care clients experience high rates of pain and nausea, regardless of prognosis. BMC Palliative Care, 2021, 20, 165.	0.8	4
24	Face-Validated Quality Indicators for Appropriateness Of End-Of-Life Care in Children With Serious Illness: A Study Using the RAND/UCLA Appropriateness Method. Journal of Pediatrics, 2021, , .	0.9	4
25	How can social workers be meaningfully involved in palliative care? A scoping review on the prerequisites and how they can be realised in practice. Palliative Care and Social Practice, 2021, 15, 263235242110588.	0.6	9
26	Feasibility of using death certificates for studying place of death in Latin America. Revista Panamericana De Salud Publica/Pan American Journal of Public Health, 2021, 45, 1.	0.6	5
27	The liminal space palliative care volunteers occupy and their roles within it: a qualitative study. BMJ Supportive and Palliative Care, 2020, 10, e28-e28.	0.8	18
28	Place of death in a small island state: a death certificate population study. BMJ Supportive and Palliative Care, 2020, 10, e30-e30.	0.8	8
29	Neonatologists and neonatal nurses have positive attitudes towards perinatal endâ€ofâ€life decisions, a nationwide survey. Acta Paediatrica, International Journal of Paediatrics, 2020, 109, 494-504.	0.7	13
30	Barriers to and Facilitators of End-of-Life Decision Making by Neonatologists and Neonatal Nurses in Neonates: A Qualitative Study. Journal of Pain and Symptom Management, 2020, 59, 599-608.e2.	0.6	11
31	Family carer support in home and hospital: a cross-sectional survey of specialised palliative care. BMJ Supportive and Palliative Care, 2020, 10, e33-e33.	0.8	5
32	Commonalities and differences in legal euthanasia and physician-assisted suicide in three countries: a population-level comparison. International Journal of Public Health, 2020, 65, 65-73.	1.0	24
33	Psychological support in end-of-life decision-making in neonatal intensive care units: Full population survey among neonatologists and neonatal nurses. Palliative Medicine, 2020, 34, 430-434.	1.3	16
34	The involvement of volunteers in palliative care and their collaboration with healthcare professionals: A crossâ€sectional volunteer survey across the Flemish healthcare system (Belgium). Health and Social Care in the Community, 2020, 28, 747-761.	0.7	9
35	Palliative and End-of-Life Care in a Small Caribbean Country: A Mortality Follow-back Study of Home Deaths. Journal of Pain and Symptom Management, 2020, 60, 1170-1180.	0.6	4
36	Trends in place of death in a small developing country: a population-level study using death certificate data. Journal of Epidemiology and Community Health, 2020, 74, jech-2019-213285.	2.0	0

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37	Volunteers in palliative care: A healthcare system-wide cross-sectional survey. BMJ Supportive and Palliative Care, 2020, , bmjspcare-2020-002321.	0.8	7
38	Developing and validating a questionnaire for mortality follow-back studies on end-of-life care and decision-making in a resource-poor Caribbean country. BMC Palliative Care, 2020, 19, 123.	0.8	1
39	Earlier palliative home care is associated with patient-centred medical resource utilisation and lower costs in the last 30 days before death in COPD: a population-level decedent cohort study. European Respiratory Journal, 2020, 55, 1901139.	3.1	9
40	The perception of people with cancer of starting a conversation about palliative care: A qualitative interview study. European Journal of Cancer Care, 2020, 29, e13282.	0.7	4
41	Appropriateness of End-of-Life Care in People Dying With Dementia: Applying Quality Indicators on Linked Administrative Databases. Journal of the American Medical Directors Association, 2020, 21, 1093-1101.e1.	1.2	10
42	Palliative care utilisation: family carers' behaviours and determinants—a qualitative interview study. BMJ Supportive and Palliative Care, 2020, , bmjspcare-2020-002207.	0.8	0
43	Inter-organisational collaboration in palliative care trajectories for nursing home residents: A nation-wide mixed methods study among key persons. International Journal of Care Coordination, 2019, 22, 69-80.	0.3	3
44	Influence of health interventions on quality of life in seriously ill children at the end of life: a systematic review protocol. Systematic Reviews, 2019, 8, 165.	2.5	1
45	Medical assistance in dying: research directions. BMJ Supportive and Palliative Care, 2019, 9, bmjspcare-2018-001727.	0.8	10
46	Educational needs of healthcare professionals and members of the general public in Alberta Canada, 2Âyears after the implementation of medical assistance in dying. Health and Social Care in the Community, 2019, 27, 1295-1302.	0.7	1
47	Is the bereavement grief intensity of survivors linked with their perception of death quality?. International Journal of Palliative Nursing, 2019, 25, 398-405.	0.2	7
48	Integrated Palliative Care for Nursing Home Residents: Exploring the Challenges in the Collaboration between Nursing Homes, Home Care and Hospitals. International Journal of Integrated Care, 2019, 19, 3.	0.1	9
49	Healthcare utilization at the end of life in people dying from amyotrophic lateral sclerosis: A retrospective cohort study using linked administrative data. Journal of the Neurological Sciences, 2019, 406, 116444.	0.3	7
50	Use and timing of referral to specialized palliative care services for people with cancer: A mortality follow-back study among treating physicians in Belgium. PLoS ONE, 2019, 14, e0210056.	1.1	12
51	Impact of palliative home care support on the quality and costs of care at the end of life: a population-level matched cohort study. BMJ Open, 2019, 9, e025180.	0.8	58
52	Discontinuation of medications at the end of life: A population study in Belgium, based on linked administrative databases. British Journal of Clinical Pharmacology, 2019, 85, 827-837.	1.1	15
53	Public Health and Epidemiological Research in Palliative Care. , 2019, , 1651-1668.		0
54	A Public Health Approach to Integrate Palliative Care into a Country's Health-Care System: Guidance as Provided by the WHO., 2019, , 1749-1768.		1

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55	End-of-Life Decision Making for People Who Died of Dementia: A Mortality Follow-Back Study Comparing 1998, 2007, and 2013 in Flanders, Belgium. Journal of the American Medical Directors Association, 2019, 20, 1347-1349.	1.2	1
56	Are We Evolving Toward Greater and Earlier Use of Palliative Home Care Support? A Trend Analysis Using Population-Level Data From 2010 to 2015. Journal of Pain and Symptom Management, 2019, 58, 19-28.e10.	0.6	9
57	Perspectives on Volunteer-Professional Collaboration in Palliative Care: A Qualitative Study Among Volunteers, Patients, Family Carers, and Health Care Professionals. Journal of Pain and Symptom Management, 2019, 58, 198-207.e7.	0.6	13
58	Who finds the road to palliative home care support? A nationwide analysis on the use of supportive measures for palliative home care using linked administrative databases. PLoS ONE, 2019, 14, e0213731.	1.1	4
59	Involvement of palliative care in euthanasia practice: Authors' reply to Regnard and Proffitt. Palliative Medicine, 2019, 33, 117-117.	1.3	0
60	Potential quality indicators for seriously ill home care clients: a cross-sectional analysis using Resident Assessment Instrument for Home Care (RAI-HC) data for Ontario. BMC Palliative Care, 2019, 18, 3.	0.8	9
61	Volunteer involvement in the organisation of palliative care: A survey study of the healthcare system in Flanders and Dutch-speaking Brussels, Belgium. Health and Social Care in the Community, 2019, 27, 459-471.	0.7	11
62	Information provision as evaluated by people with cancer and bereaved relatives: A cross-sectional survey of 34 specialist palliative care teams. Patient Education and Counseling, 2019, 102, 768-775.	1.0	3
63	Resource Use During the Last 6 Months of Life of Individuals Dying with and of Alzheimer's Disease. Journal of the American Geriatrics Society, 2018, 66, 879-885.	1.3	8
64	End-of-life decision-making across cancer types: results from a nationwide retrospective survey among treating physicians. British Journal of Cancer, 2018, 118, 1369-1376.	2.9	7
65	How accurately is euthanasia reported on death certificates in a country with legal euthanasia: a population-based study. European Journal of Epidemiology, 2018, 33, 689-693.	2.5	49
66	The use of behavioural theories in end-of-life care research: A systematic review. Palliative Medicine, 2018, 32, 1055-1077.	1.3	20
67	Do physicians discuss end-of-life decisions with family members? A mortality follow-back study. Patient Education and Counseling, 2018, 101, 1378-1384.	1.0	8
68	Does using the interRAI Palliative Care instrument reduce the needs and symptoms of nursing home residents receiving palliative care?. Palliative and Supportive Care, 2018, 16, 32-40.	0.6	8
69	Information needs about palliative care and euthanasia: A survey of patients in different phases of their cancer trajectory. Patient Education and Counseling, 2018, 101, 132-138.	1.0	14
70	Bereavement grief: A population-based foundational evidence study. Death Studies, 2018, 42, 463-469.	1.8	15
71	Involvement of palliative care in euthanasia practice in a context of legalized euthanasia: A population-based mortality follow-back study. Palliative Medicine, 2018, 32, 114-122.	1.3	41
72	What influences intentions to request physician-assisted euthanasia or continuous deep sedation?. Death Studies, 2018, 42, 491-497.	1.8	7

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73	A Public Health Approach to Integrate Palliative Care into a Country's Health-Care System: Guidance as Provided by the WHO. , 2018, , 1-20.		O
74	A post-mortem population survey on foetal-infantile end-of-life decisions: a research protocol. BMC Pediatrics, 2018, 18, 260.	0.7	5
75	Drugs Used for Euthanasia: A Repeated Population-Based Mortality Follow-Back Study in Flanders, Belgium, 1998–2013. Journal of Pain and Symptom Management, 2018, 56, 551-559.	0.6	8
76	Palliative care volunteerism across the healthcare system: A survey study. Palliative Medicine, 2018, 32, 1233-1245.	1.3	23
77	Comparing the use and timing of palliative care services in COPD and lung cancer: a population-based survey. European Respiratory Journal, 2018, 51, 1702405.	3.1	12
78	Appropriateness of End-of-Life Care in People Dying From COPD. Applying Quality Indicators on Linked Administrative Databases. Journal of Pain and Symptom Management, 2018, 56, 541-550.e6.	0.6	13
79	Place of death for people with HIV: a population-level comparison of eleven countries across three continents using death certificate data. BMC Infectious Diseases, 2018, 18, 55.	1.3	12
80	Resource Use During the Last SixÂMonths of Life Among COPD Patients: A Population-Level Study. Journal of Pain and Symptom Management, 2018, 56, 318-326.e7.	0.6	12
81	Main themes, barriers, and solutions to palliative and end-of-life care in the English-speaking Caribbean: a scoping review. Revista Panamericana De Salud Publica/Pan American Journal of Public Health, 2018, 42, 1-9.	0.6	3
82	Public Health and Epidemiological Research in Palliative Care. , 2018, , 1-18.		0
83	Systematic Quality Monitoring For Specialized Palliative Care Services: Development of a Minimal Set of Quality Indicators for Palliative Care Study (QPAC). American Journal of Hospice and Palliative Medicine, 2017, 34, 532-546.	0.8	16
84	Appropriateness and avoidability of terminal hospital admissions: Results of a survey among family physicians. Palliative Medicine, 2017, 31, 456-464.	1.3	17
85	Place of death of children with complex chronic conditions: cross-national study of 11 countries. European Journal of Pediatrics, 2017, 176, 327-335.	1.3	52
86	Developing indicators of appropriate and inappropriate end-of-life care in people with Alzheimer's disease, cancer or chronic obstructive pulmonary disease for population-level administrative databases: A RAND/UCLA appropriateness study. Palliative Medicine, 2017, 31, 932-945.	1.3	60
87	International variations in clinical practice guidelines for palliative sedation: a systematic review. BMJ Supportive and Palliative Care, 2017, 7, bmjspcare-2016-001159.	0.8	79
88	Improving comfort around dying in elderly people: a cluster randomised controlled trial. Lancet, The, 2017, 390, 125-134.	6.3	28
89	Differences in place of death between lung cancer and COPD patients: a 14-country study using death certificate data. Npj Primary Care Respiratory Medicine, 2017, 27, 14.	1.1	52
90	Determinants in the place of death for people with different cancer types: a national population-based study. Acta Oncol \tilde{A}^3 gica, 2017, 56, 455-461.	0.8	23

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91	End-of-Life Care and Quality of Dying in 23 Acute Geriatric Hospital Wards in Flanders, Belgium. Journal of Pain and Symptom Management, 2017, 53, 693-702.	0.6	17
92	Degeneracy-Preserving Quantum Nondemolition Measurement of Parity-Type Observables for Cat Qubits. Physical Review Letters, 2017, 119, 060503.	2.9	27
93	Applying Quality Indicators For Administrative Databases To Evaluate End-Of-Life Care For Cancer Patients In Belgium. Health Affairs, 2017, 36, 1234-1243.	2.5	32
94	Euthanasia for people with psychiatric disorders or dementia in Belgium: analysis of officially reported cases. BMC Psychiatry, 2017, 17, 203.	1.1	74
95	Estimating the need for palliative care at the population level: A cross-national study in 12 countries. Palliative Medicine, 2017, 31, 526-536.	1.3	111
96	Palliative care needs and symptoms of nursing home residents with and without dementia: A crossâ€sectional study. Geriatrics and Gerontology International, 2017, 17, 1501-1507.	0.7	24
97	Euthanasia and Public Health. , 2017, , 46-56.		2
98	Policy Measures to Support Palliative Care at Home: A Cross-Country Case Comparison in Three European Countries. Journal of Pain and Symptom Management, 2017, 54, 523-529.e5.	0.6	23
99	Trends in Continuous Deep Sedation until Death between 2007 and 2013: A Repeated Nationwide Survey. PLoS ONE, 2016, 11, e0158188.	1.1	49
100	Palliative care needs at different phases in the illness trajectory: a survey study in patients with cancer. European Journal of Cancer Care, 2016, 25, 534-543.	0.7	37
101	Attitudes and Practices of Euthanasia and Physician-Assisted Suicide in the United States, Canada, and Europe. JAMA - Journal of the American Medical Association, 2016, 316, 79.	3.8	462
102	The European Association for Palliative Care White Paper on euthanasia and physician-assisted suicide: Dodging responsibility. Palliative Medicine, 2016, 30, 893-894.	1.3	7
103	Reasons for End-of-Life Hospital Admissions: Results of a Survey Among Family Physicians. Journal of Pain and Symptom Management, 2016, 52, 498-506.	0.6	21
104	Usefulness, feasibility and face validity of the interRAI Palliative Care instrument according to care professionals in nursing homes: A qualitative study. International Journal of Nursing Studies, 2016, 62, 90-99.	2.5	8
105	Euthanasia in Belgium: trends in reported cases between 2003 and 2013. Cmaj, 2016, 188, E407-E414.	0.9	48
106	Resource Use and Health Care Costs of COPD Patients at the End of Life: A Systematic Review. Journal of Pain and Symptom Management, 2016, 52, 588-599.	0.6	37
107	Euthanasia and Physician-Assisted Suicide—In Reply. JAMA - Journal of the American Medical Association, 2016, 316, 1600.	3.8	6
108	Using linked administrative and disease-specific databases to study end-of-life care on a population level. BMC Palliative Care, 2016, 15, 86.	0.8	40

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109	Implementing the care programme for the last days of life in an acute geriatric hospital ward: a phase 2 mixed method study. BMC Palliative Care, 2016, 15, 27.	0.8	9
110	Is There a Need for Early Palliative Care in Patients With Life-Limiting Illnesses? Interview Study With Patients About Experienced Care Needs From Diagnosis Onward. American Journal of Hospice and Palliative Medicine, 2016, 33, 489-497.	0.8	44
111	Comparison of Site of Death, Health Care Utilization, and Hospital Expenditures for Patients Dying With Cancer in 7 Developed Countries. JAMA - Journal of the American Medical Association, 2016, 315, 272.	3.8	388
112	International comparison of death place for suicide; a population-level eight country death certificate study. Social Psychiatry and Psychiatric Epidemiology, 2016, 51, 101-106.	1.6	9
113	Place of death in the population dying from diseases indicative of palliative care need: a cross-national population-level study in 14 countries. Journal of Epidemiology and Community Health, 2016, 70, 17-24.	2.0	150
114	Research in public health and end-of-life care – Building on the past and developing the new. Progress in Palliative Care, 2016, 24, 25-30.	0.7	28
115	Prescription and Deprescription of Medication During the Last 48 Hours of Life: Multicenter Study in 23 Acute Geriatric Wards in Flanders, Belgium. Journal of Pain and Symptom Management, 2016, 51, 1020-1026.	0.6	19
116	Surveying End-of-Life Medical Decisions in France: Evaluation of an Innovative Mixed-Mode Data Collection Strategy. Interactive Journal of Medical Research, 2016, 5, e8.	0.6	0
117	Family physicians' role in palliative care throughout the care continuum: stakeholder perspectives. Family Practice, 2015, 32, cmv072.	0.8	27
118	Development of the care programme for the last days of life for older patients in acute geriatric hospital wards: a phase $0\hat{a}\in^{c}1$ study according to the Medical Research Council Framework. BMC Palliative Care, 2015, 14, 24.	0.8	7
119	A population-level study of place of death and associated factors in Sweden. Scandinavian Journal of Public Health, 2015, 43, 744-751.	1.2	89
120	Comparison of the Expression and Granting of Requests for Euthanasia in Belgium in 2007 vs 2013. JAMA Internal Medicine, 2015, 175, 1703.	2.6	84
121	International Variation in Place of Death of Older People Who Died From Dementia in 14 European and non-European Countries. Journal of the American Medical Directors Association, 2015, 16, 165-171.	1.2	105
122	Attitudes of Belgian Students of Medicine, Philosophy, and Law Toward Euthanasia and the Conditions for Its Acceptance. Death Studies, 2015, 39, 139-150.	1.8	17
123	Physical and Psychological Distress Are Related to Dying Peacefully in Residents With Dementia in Long-Term Care Facilities. Journal of Pain and Symptom Management, 2015, 50, 1-8.	0.6	13
124	Patient Characteristics Associated With Prognostic Awareness: A Study of a Canadian Palliative Care Population Using the InterRAI Palliative Care Instrument. Journal of Pain and Symptom Management, 2015, 49, 716-725.	0.6	24
125	How to implement quality indicators successfully in palliative care services: perceptions of team members about facilitators of and barriers to implementation. Supportive Care in Cancer, 2015, 23, 3503-3511.	1.0	19
126	Improving end-of-life care in acute geriatric hospital wards using the Care Programme for the Last Days of Life: study protocol for a phase 3 cluster randomized controlled trial. BMC Geriatrics, 2015, 15, 13.	1.1	8

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127	Endâ€ofâ€Life Decisions in Individuals Dying with Dementia in Belgium. Journal of the American Geriatrics Society, 2015, 63, 290-296.	1.3	24
128	Recent Trends in Euthanasia and Other End-of-Life Practices in Belgium. New England Journal of Medicine, 2015, 372, 1179-1181.	13.9	141
129	Place of death of people living with Parkinson's disease: a population-level study in 11 countries. BMC Palliative Care, 2015, 14, 28.	0.8	38
130	International study of the place of death of people with cancer: a population-level comparison of 14 countries across 4 continents using death certificate data. British Journal of Cancer, 2015, 113, 1397-1404.	2.9	158
131	Quality indicators for palliative care services: Mixed-method study testing for face validity, feasibility, discriminative power and usefulness. Palliative Medicine, 2015, 29, 71-82.	1.3	19
132	What Are Physicians' Reasons for Not Referring People with Life-Limiting Illnesses to Specialist Palliative Care Services? A Nationwide Survey. PLoS ONE, 2015, 10, e0137251.	1.1	49
133	Informed palliative care in nursing homes through the interRAI Palliative Care instrument: a study protocol based on the Medical Research Council framework. BMC Geriatrics, 2014, 14, 132.	1.1	10
134	Early identification of palliative care needs by family physicians: A qualitative study of barriers and facilitators from the perspective of family physicians, community nurses, and patients. Palliative Medicine, 2014, 28, 480-490.	1.3	64
135	What justifies a hospital admission at the end of life? A focus group study on perspectives of family physicians and nurses. Palliative Medicine, 2014, 28, 941-948.	1.3	45
136	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
137	Dying in hospital: a study of incidence and factors related to hospital death using death certificate data. European Journal of Public Health, 2014, 24, 751-756.	0.1	28
138	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.	1.3	86
139	How do general end-of-life treatment goals and values relate to specific treatment preferences? A population-based study. Palliative Medicine, 2014, 28, 1206-1212.	1.3	16
140	Media Coverage of Medical Decision Making at the End of Life: A Belgian Case Study. Death Studies, 2014, 38, 125-135.	1.8	7
141	Descriptions of euthanasia as social representations: comparing the views of Finnish physicians and religious professionals. Sociology of Health and Illness, 2014, 36, 354-368.	1.1	8
142	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.	0.8	26
143	Forgoing artificial nutrition or hydration at the end of life: a large cross-sectional survey in Belgium. Journal of Medical Ethics, 2014, 40, 501-504.	1.0	11
144	The Family Physician's Perceived Role in Preventing and Guiding Hospital Admissions at the End of Life: A Focus Group Study. Annals of Family Medicine, 2014, 12, 441-446.	0.9	21

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145	The acute hospital setting as a place of death and final care: A qualitative study on perspectives of family physicians, nurses and family carers. Health and Place, 2014, 27, 77-83.	1.5	58
146	Public acceptance of euthanasia in Europe: a survey study in 47 countries. International Journal of Public Health, 2014, 59, 143-156.	1.0	86
147	Labelling of end-of-life decisions by physicians. Journal of Medical Ethics, 2014, 40, 505-507.	1.0	55
148	Nationwide survey to evaluate the decision-making process in euthanasia requests in Belgium: do specifically trained 2nd physicians improve quality of consultation?. BMC Health Services Research, 2014, 14, 307.	0.9	15
149	Factors associated with the goal of treatment in the last week of life in old compared to very old patients: a population-based death certificate survey. BMC Geriatrics, 2014, 14, 61.	1.1	6
150	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.	0.6	322
151	A Comparative Analysis of Comprehensive Geriatric Assessments forÂNursing Home Residents Receiving Palliative Care: AÂSystematicÂReview. Journal of the American Medical Directors Association, 2014, 15, 467-476.	1.2	19
152	Nurses' involvement in physician-assisted dying under the euthanasia law in Belgium. International Journal of Nursing Studies, 2014, 51, 1696-1697.	2.5	13
153	Dying in Hospital with Dementia and Pneumonia: A Nationwide Study Using Death Certificate Data. Gerontology, 2014, 60, 31-37.	1.4	10
154	Afterword: The Social Construction of Death: Reflections from a Quantitative Public Health Researcher., 2014,, 259-270.		1
155	Towards a standardized method of developing quality indicators for palliative care: protocol of the Quality indicators for Palliative Care (Q-PAC) study. BMC Palliative Care, 2013, 12, 6.	0.8	33
156	Is educational attainment related to end-of-life decision-making? A large post-mortem survey in Belgium. BMC Public Health, 2013, 13, 1055.	1.2	23
157	Referral to palliative care in COPD and other chronic diseases: A population-based study. Respiratory Medicine, 2013, 107, 1731-1739.	1.3	160
158	Quality Indicators for Palliative Care: Update of a Systematic Review. Journal of Pain and Symptom Management, 2013, 46, 556-572.	0.6	158
159	The public's viewpoint on the right to hastened death in Alberta, Canada: findings from a population survey study. Health and Social Care in the Community, 2013, 21, 200-208.	0.7	8
160	Different trends in euthanasia acceptance across Europe. A study of 13 western and 10 central and eastern European countries, 1981–2008. European Journal of Public Health, 2013, 23, 378-380.	0.1	36
161	Social-cultural factors in end-of-life care in Belgium: A scoping of the research literature. Palliative Medicine, 2013, 27, 131-143.	1.3	14
162	The Preferred Place of Last Days: Results of a Representative Population-Based Public Survey. Journal of Palliative Medicine, 2013, 16, 502-508.	0.6	81

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