

Peter Hudson

List of Publications by Year in descending order

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Version: 2024-02-01

84
papers

3,183
citations

147566

31
h-index

168136

53
g-index

87
all docs

87
docs citations

87
times ranked

2844
citing authors

#	ARTICLE	IF	CITATIONS
1	Family carer needs in advanced disease: systematic review of reviews. <i>BMJ Supportive and Palliative Care</i> , 2022, 12, 132-141.	0.8	9
2	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. <i>Palliative Medicine</i> , 2022, 36, 529-539.	1.3	4
3	Implementability of healthcare interventions: an overview of reviews and development of a conceptual framework. <i>Implementation Science</i> , 2022, 17, 10.	2.5	72
4	Family carers research: What progress has been made?. <i>Palliative Medicine</i> , 2022, , 026921632110378.	1.3	5
5	Do family meetings for hospitalised palliative care patients improve outcomes and reduce health care costs? A cluster randomised trial. <i>Palliative Medicine</i> , 2021, 35, 188-199.	1.3	10
6	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. <i>Palliative Medicine</i> , 2021, 35, 109-119.	1.3	25
7	Validation of the responding to urgency of need in palliative care (RUN-PC) triage tool. <i>Palliative Medicine</i> , 2021, 35, 759-767.	1.3	4
8	Developing a model of bereavement care in an adult tertiary hospital. <i>Australian Health Review</i> , 2021, 45, 110.	0.5	8
9	Determining the informational needs of family caregivers of people with intellectual disability who require palliative care: A qualitative study. <i>Palliative and Supportive Care</i> , 2021, 19, 405-414.	0.6	7
10	Reframing palliative care to improve the quality of life of people diagnosed with a serious illness. <i>Medical Journal of Australia</i> , 2021, 215, 443-446.	0.8	10
11	Evaluation of an online toolkit for carers of people with a life-limiting illness at the end-of-life: health professionals'™ perspectives. <i>Australian Journal of Primary Health</i> , 2021, 27, 473-478.	0.4	5
12	Triaging the Terminally Ill™Development of the Responding toUrgency of Need in Palliative Care (RUN-PC) Triage Tool. <i>Journal of Pain and Symptom Management</i> , 2020, 59, 95-104.e11.	0.6	6
13	Characteristics of family meetings for patients with advanced disease in an Australian metropolitan tertiary hospital. <i>BMJ Supportive and Palliative Care</i> , 2020, , bmjspcare-2020-002250.	0.8	0
14	Improving Support for Family Caregivers of People with a Serious Illness in the United States: Strategic Agenda and Call to Action. <i>Palliative Medicine Reports</i> , 2020, 1, 6-17.	0.4	22
15	“œA monster that lives in our lives” experiences of caregivers of people with motor neuron disease and identifying avenues for support: TableA1. <i>BMJ Supportive and Palliative Care</i> , 2019, 9, bmjspcare-2015-001057.	0.8	16
16	Development of an eHealth information resource for family carers supporting a person receiving palliative care on the island of Ireland. <i>BMC Palliative Care</i> , 2019, 18, 74.	0.8	5
17	Informational needs of family caregivers of people with intellectual disability who require palliative care: a two-phase integrative review of the literature. <i>International Journal of Palliative Nursing</i> , 2019, 25, 4-18.	0.2	4
18	Novel application of discrete choice experiment methodology to understand how clinicians around the world triage palliative care needs: A research protocol. <i>Palliative and Supportive Care</i> , 2019, 17, 66-73.	0.6	5

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19	Palliative care for people with non-malignant respiratory disease and their carers: a review of the current evidence. <i>Journal of Research in Nursing</i> , 2019, 24, 420-430.	0.3	2
20	A randomised phase II trial to examine feasibility of standardised, early palliative (STEP) care for patients with advanced cancer and their families [ACTRN12617000534381]: a research protocol. <i>Pilot and Feasibility Studies</i> , 2019, 5, 44.	0.5	10
21	The Ethics of Prioritizing Access to Palliative Care: A Qualitative Study. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 577-582.	0.8	7
22	Long-term prevalence and predictors of prolonged grief disorder amongst bereaved cancer caregivers: A cohort study. <i>Palliative and Supportive Care</i> , 2019, 17, 507-514.	0.6	24
23	Patterns of end-of-life hospital care for patients with non-Hodgkin lymphoma: exploring the landscape. <i>Leukemia and Lymphoma</i> , 2019, 60, 1908-1916.	0.6	3
24	Using a six-step co-design model to develop and test a peer-led web-based resource (PLWR) to support informal carers of cancer patients. <i>Psycho-Oncology</i> , 2019, 28, 518-524.	1.0	31
25	The trajectory of patients who die from metastatic prostate cancer: a population-based study. <i>BJU International</i> , 2019, 123, 19-26.	1.3	13
26	Transition Points for the Routine Integration of Palliative Care in Patients With Advanced Cancer. <i>Journal of Pain and Symptom Management</i> , 2018, 56, 185-194.	0.6	31
27	The experiences and preparedness of family carers for best interest decision-making of a relative living with advanced dementia: A qualitative study. <i>Journal of Advanced Nursing</i> , 2018, 74, 1595-1604.	1.5	35
28	A self-care, problem-solving and mindfulness intervention for informal caregivers of people with motor neurone disease: A pilot study. <i>Palliative Medicine</i> , 2018, 32, 726-732.	1.3	7
29	A conceptual model: Redesigning how we provide palliative care for patients with chronic obstructive pulmonary disease. <i>Palliative and Supportive Care</i> , 2018, 16, 452-460.	0.6	20
30	Bereavement support standards and bereavement care pathway for quality palliative care. <i>Palliative and Supportive Care</i> , 2018, 16, 375-387.	0.6	52
31	The experience of palliative care service provision for people with non-malignant respiratory disease and their family carers: An all-Ireland qualitative study. <i>Journal of Advanced Nursing</i> , 2018, 74, 383-394.	1.5	18
32	Effectiveness of advance care planning with family carers in dementia nursing homes: A paired cluster randomized controlled trial. <i>Palliative Medicine</i> , 2018, 32, 603-612.	1.3	116
33	Responding to urgency of need: Initial qualitative stage in the development of a triage tool for use in palliative care services. <i>Palliative Medicine</i> , 2018, 32, 1246-1254.	1.3	8
34	A randomized open-label study of guideline-driven antiemetic therapy versus single agent antiemetic therapy in patients with advanced cancer and nausea not related to anticancer treatment. <i>BMC Cancer</i> , 2018, 18, 510.	1.1	22
35	The provision of generalist and specialist palliative care for patients with non-malignant respiratory disease in the North and Republic of Ireland: a qualitative study. <i>BMC Palliative Care</i> , 2018, 17, 6.	0.8	20
36	A metasynthesis study of family caregivers' transition experiences caring for community-dwelling persons with advanced cancer at the end of life. <i>Palliative Medicine</i> , 2017, 31, 602-616.	1.3	35

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37	Honoring the voices of bereaved caregivers: a Metasummary of qualitative research. <i>BMC Palliative Care</i> , 2017, 16, 48.	0.8	28
38	Toward a systematic approach to assessment and care planning in palliative care: A practical review of clinical tools. <i>Palliative and Supportive Care</i> , 2016, 14, 161-173.	0.6	13
39	Establishment and preliminary outcomes of a palliative care research network. <i>Palliative and Supportive Care</i> , 2016, 14, 52-59.	0.6	2
40	Music's Relevance for People Affected by Cancer: A Meta-Ethnography and Implications for Music Therapists. <i>Journal of Music Therapy</i> , 2016, 53, 398-429.	0.6	25
41	Health care professionals' experience, understanding and perception of need of advanced cancer patients with cachexia and their families: The benefits of a dedicated clinic. <i>BMC Palliative Care</i> , 2016, 15, 100.	0.8	21
42	Insights into hospitalisation of advanced cancer patients: a study of medical records. <i>European Journal of Cancer Care</i> , 2016, 25, 190-201.	0.7	8
43	Legalizing physician-assisted suicide and/or euthanasia: Pragmatic implications. <i>Palliative and Supportive Care</i> , 2015, 13, 1399-1409.	0.6	19
44	Benefits and resource implications of family meetings for hospitalized palliative care patients: research protocol. <i>BMC Palliative Care</i> , 2015, 14, 73.	0.8	11
45	Metastatic non-small cell lung cancer: a benchmark for quality end-of-life cancer care?. <i>Medical Journal of Australia</i> , 2015, 202, 139-143.	0.8	24
46	Reducing the psychological distress of family caregivers of home based palliative care patients: longer term effects from a randomised controlled trial. <i>Psycho-Oncology</i> , 2015, 24, 19-24.	1.0	102
47	Protocol for the Care-IS Trial: a randomised controlled trial of a supportive educational intervention for carers of patients with high-grade glioma (HGG). <i>BMJ Open</i> , 2015, 5, e009477.	0.8	19
48	Use of the Preparedness for Caregiving Scale in Palliative Care: A Rasch Evaluation Study. <i>Journal of Pain and Symptom Management</i> , 2015, 50, 533-541.	0.6	22
49	The Melbourne Family Support Program: evidence-based strategies that prepare family caregivers for supporting palliative care patients: Table A1. <i>BMJ Supportive and Palliative Care</i> , 2014, 4, 231-237.	0.8	34
50	Establishing a nurse practitioner collaborative: evolution, development, and outcomes. <i>International Journal of Palliative Nursing</i> , 2014, 20, 457-461.	0.2	2
51	The experiences of palliative care health service provision for people with non-malignant respiratory disease and their caregivers: an all-Ireland study. <i>Journal of Advanced Nursing</i> , 2014, 70, 687-697.	1.5	2
52	Risk Factors for Developing Prolonged Grief During Bereavement in Family Carers of Cancer Patients in Palliative Care: A Longitudinal Study. <i>Journal of Pain and Symptom Management</i> , 2014, 47, 531-541.	0.6	109
53	Sound Continuing Bonds with the Deceased: The Relevance of Music, Including Preloss Music Therapy, for Eight Bereaved Caregivers. <i>Death Studies</i> , 2013, 37, 101-125.	1.8	34
54	Improving support for family carers: Key implications for research, policy and practice. <i>Palliative Medicine</i> , 2013, 27, 581-582.	1.3	39

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55	Experiences of carers supporting dying renal patients managed without dialysis. <i>Journal of Advanced Nursing</i> , 2013, 69, 1829-1839.	1.5	27
56	Reducing the psychological distress of family caregivers of home-based palliative care patients: short-term effects from a randomised controlled trial. <i>Psycho-Oncology</i> , 2013, 22, 1987-1993.	1.0	84
57	International Association for Hospice and Palliative Care (IAHPC) List of Essential Practices in Palliative Care. <i>Journal of Pain and Palliative Care Pharmacotherapy</i> , 2012, 26, 118-122.	0.5	42
58	Palliative care provision for those with non-malignant respiratory disease: a literature review. <i>BMJ Supportive and Palliative Care</i> , 2012, 2, A6.2-A7.	0.8	0
59	Guidelines for the Psychosocial and Bereavement Support of Family Caregivers of Palliative Care Patients. <i>Journal of Palliative Medicine</i> , 2012, 15, 696-702.	0.6	141
60	Psycho-Educational Group Intervention for Family Caregivers of Hospitalized Palliative Care Patients: Pilot Study. <i>Journal of Palliative Medicine</i> , 2012, 15, 277-281.	0.6	14
61	Psychological and Social Profile of Family Caregivers on Commencement of Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2011, 41, 522-534.	0.6	118
62	Research Priorities Associated with Family Caregivers in Palliative Care: International Perspectives. <i>Journal of Palliative Medicine</i> , 2011, 14, 397-401.	0.6	34
63	Family Caregivers and Palliative Care: Current Status and Agenda for the Future. <i>Journal of Palliative Medicine</i> , 2011, 14, 864-869.	0.6	126
64	A systematic review of psychosocial interventions for family carers of palliative care patients. <i>BMC Palliative Care</i> , 2010, 9, 17.	0.8	147
65	Meeting the needs of family carers: an evaluation of three home-based palliative care services in Australia. <i>Palliative Medicine</i> , 2010, 24, 183-191.	1.3	34
66	Psychometric Properties of the Hospital Anxiety and Depression Scale in Family Caregivers of Palliative Care Patients. <i>Journal of Pain and Symptom Management</i> , 2009, 37, 797-806.	0.6	90
67	Teaching Family Carers About Home-Based Palliative Care: Final Results from a Group Education Program. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 299-308.	0.6	64
68	Assessing the Family and Caregivers. , 2009, , 320-325.		5
69	Family meetings in palliative care: Multidisciplinary clinical practice guidelines. <i>BMC Palliative Care</i> , 2008, 7, 12.	0.8	120
70	“Palliative Care: The Essentials” Evaluation of a Multidisciplinary Education Program. <i>Journal of Palliative Medicine</i> , 2008, 11, 1122-1129.	0.6	38
71	The future of family caregiving: research, social policy and clinical practice. , 2008, , 277-304.		4
72	Australia and New Zealand. , 2007, , 59-68.		3

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73	Diabetes Management in Patients Receiving Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2006, 32, 275-286.	0.6	57
74	Would people with Parkinson's disease benefit from palliative care?. <i>Palliative Medicine</i> , 2006, 20, 87-94.	1.3	82
75	How Well Do Family Caregivers Cope after Caring for a Relative with Advanced Disease and How Can Health Professionals Enhance their Support?. <i>Journal of Palliative Medicine</i> , 2006, 9, 694-703.	0.6	64
76	Positive aspects and challenges associated with caring for a dying relative at home. <i>International Journal of Palliative Nursing</i> , 2004, 10, 58-65.	0.2	151
77	Meeting the Supportive Needs of Family Caregivers in Palliative Care: Challenges for Health Professionals. <i>Journal of Palliative Medicine</i> , 2004, 7, 19-25.	0.6	162
78	A Critical Review of Supportive Interventions for Family Caregivers of Patients with Palliative-Stage Cancer. <i>Journal of Psychosocial Oncology</i> , 2004, 22, 77-92.	0.6	42
79	Focus group interviews: a guide for palliative care researchers and clinicians. <i>International Journal of Palliative Nursing</i> , 2003, 9, 202-207.	0.2	40
80	A conceptual model and key variables for guiding supportive interventions for family caregivers of people receiving palliative care. <i>Palliative and Supportive Care</i> , 2003, 1, 353-365.	0.6	70
81	The experience of research participation for family caregivers of palliative care cancer patients. <i>International Journal of Palliative Nursing</i> , 2003, 9, 120-123.	0.2	40
82	Home-based support for palliative care families: challenges and recommendations. <i>Medical Journal of Australia</i> , 2003, 179, S35-7.	0.8	36
83	Intervention development for enhanced lay palliative caregiver support - the use of focus groups. <i>European Journal of Cancer Care</i> , 2002, 11, 262-270.	0.7	74
84	Randomized controlled trials in palliative care: overcoming the obstacles. <i>International Journal of Palliative Nursing</i> , 2001, 7, 427-434.	0.2	60