

Katherine J Hunt

List of Publications by Year in descending order

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Version: 2024-02-01

30
papers

780
citations

567281

15
h-index

526287

27
g-index

30
all docs

30
docs citations

30
times ranked

1314
citing authors

#	ARTICLE	IF	CITATIONS
1	Living With, Managing and Minimising Treatment Burden in Long Term Conditions: A Systematic Review of Qualitative Research. PLoS ONE, 2015, 10, e0125457.	2.5	104
2	Work of being an adult patient with chronic kidney disease: a systematic review of qualitative studies. BMJ Open, 2018, 8, e023507.	1.9	83
3	The nursing work of hospital-based clinical practice guideline implementation: An explanatory systematic review using Normalisation Process Theory. International Journal of Nursing Studies, 2014, 51, 289-299.	5.6	70
4	End-of-life care and achieving preferences for place of death in England: Results of a population-based survey using the VOICES-SF questionnaire. Palliative Medicine, 2014, 28, 412-421.	3.1	65
5	Palliative and end-of-life care conversations in COPD: a systematic literature review. ERJ Open Research, 2017, 3, 00068-2016.	2.6	56
6	Participant recruitment in sensitive surveys: a comparative trial of "opt in" versus "opt out" approaches. BMC Medical Research Methodology, 2013, 13, 3.	3.1	51
7	Development and testing of a text-mining approach to analyse patients' comments on their experiences of colorectal cancer care. BMJ Quality and Safety, 2016, 25, 604-614.	3.7	43
8	Inflammation in Aging Part 1: Physiology and Immunological Mechanisms. Biological Research for Nursing, 2010, 11, 245-252.	1.9	40
9	End-of-Life Care and Preferences for Place of Death among the Oldest Old: Results of a Population-Based Survey Using VOICES' Short Form. Journal of Palliative Medicine, 2014, 17, 176-182.	1.1	37
10	Inflammation in Aging Part 2: Implications for the Health of Older People and Recommendations for Nursing Practice. Biological Research for Nursing, 2010, 11, 253-260.	1.9	28
11	The palliative care needs of adults with intellectual disabilities and their access to palliative care services: A systematic review. Palliative Medicine, 2020, 34, 1006-1018.	3.1	28
12	Developing the methods and questionnaire (VOICES-SF) for a national retrospective mortality follow-back survey of palliative and end-of-life care in England. BMJ Supportive and Palliative Care, 2019, 9, e5-e5.	1.6	27
13	End-of-life care in intellectual disability: a retrospective cross-sectional study. BMJ Supportive and Palliative Care, 2020, 10, 469-477.	1.6	19
14	Last months of life of people with intellectual disabilities: A UK population-based study of death and dying in intellectual disability community services. Journal of Applied Research in Intellectual Disabilities, 2020, 33, 1245-1258.	2.0	19
15	Managing expectations: cognitive authority and experienced control in complex healthcare processes. BMC Health Services Research, 2017, 17, 459.	2.2	18
16	The preferences of patients with chronic obstructive pulmonary disease are to discuss palliative care plans with familiar respiratory clinicians, but to delay conversations until their condition deteriorates: A study guided by interpretative phenomenological analysis. Palliative Medicine, 2020, 34, 1361-1373.	3.1	14
17	Development and initial validation of a new outcome measure for hospice and palliative care: the St Christopher's Index of Patient Priorities (SKIPP). BMJ Supportive and Palliative Care, 2014, 4, 175-181.	1.6	12
18	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.	2.9	12

#	ARTICLE	IF	CITATIONS
19	Reducing avoidable hospital admission in older people: Health status, frailty and predicting risk of ill-defined conditions diagnoses in older people admitted with collapse. Archives of Gerontology and Geriatrics, 2013, 57, 172-176.	3.0	9
20	EXPERTS 1–experiences of long-term life-limiting conditions among patients and carers: protocol for a qualitative meta-synthesis and conceptual modelling study. BMJ Open, 2015, 5, e007372-e007372.	1.9	9
21	Cross-cultural adaptation and measurement properties of generic and cancer-related patient-reported outcome measures (PROMs) for use with cancer patients in Brazil: a systematic review. Quality of Life Research, 2018, 27, 857-870.	3.1	9
22	Nursing care at end of life: a UK-based survey of the deaths of people living in care settings for people with intellectual disability. Journal of Research in Nursing, 2019, 24, 366-382.	0.9	8
23	Burden of treatment in the light of the international classification of functioning, disability and health: a “best fit” framework synthesis. Disability and Rehabilitation, 2017, 39, 1253-1261.	1.8	6
24	Expecting the unexpected: measures, outcomes and dying trajectories for expected and unexpected death in adults with intellectual disabilities in social care settings in the UK. Journal of Applied Research in Intellectual Disabilities, 2021, 34, 594-605.	2.0	5
25	Prognostic disclosure and quality of life in palliative care: a systematic review. BMJ Supportive and Palliative Care, 2020, 11, bmjspcare-2020-002460.	1.6	4
26	Hidden lives and deaths: the last months of life of people with intellectual disabilities living in long-term, generic care settings in the UK. Journal of Applied Research in Intellectual Disabilities, 2021, 34, 1489-1498.	2.0	2
27	Clinician Perspectives on How to Hold Earlier Discussions About Palliative and End-of-Life Care With Chronic Obstructive Pulmonary Disease Patients. Journal of Hospice and Palliative Nursing, 2022, Publish Ahead of Print, .	0.9	2
28	Achieving preferred place of care and death: Results from a post bereavement survey. BMJ Supportive and Palliative Care, 2012, 2, A4.3-A5.	1.6	0
29	Planning the first national end of life care survey. BMJ Supportive and Palliative Care, 2012, 2, A8.1-A8.	1.6	0
30	98â€¦Palliative and advance care planning discussions with copd patients. , 2018, , .		0