## Amy E Waller

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2868076/publications.pdf

Version: 2024-02-01

201575 197736 2,651 94 27 49 citations h-index g-index papers 94 94 94 4058 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Advance care planning participation by people with dementia: a cross-sectional survey and medical record audit. BMJ Supportive and Palliative Care, 2022, 12, e464-e468.	0.8	4
2	Barriers to the provision of optimal care to dying patients in hospital: An international cross-sectional comparison study of nurses' perceptions. Collegian, 2022, 29, 62-69.	0.6	0
3	Inadequate completion of advance care directives by individuals with dementia: national audit of health and aged care facilities. BMJ Supportive and Palliative Care, 2022, 12, e319-e328.	0.8	5
4	Prioritising national dementia guidelines for general practice: A Delphi approach. Australasian Journal on Ageing, 2022, 41, 247-257.	0.4	2
5	A Systematic Review of the Development and Implementation of Needs-Based Palliative Care Tools in Heart Failure and Chronic Respiratory Disease. Frontiers in Cardiovascular Medicine, 2022, 9, 878428.	1.1	3
6	Junior medical doctors' decision making when using advance care directives to guide treatment for people with dementia: a cross-sectional vignette study. BMC Medical Ethics, 2022, 23, .	1.0	0
7	Knowledge and confidence of junior medical doctors in discussing and documenting resuscitation plans: a crossâ€sectional survey. Internal Medicine Journal, 2021, 51, 2055-2060.	0.5	2
8	Preferences for life expectancy discussions following diagnosis with a life-threatening illness: a discrete choice experiment. Supportive Care in Cancer, 2021, 29, 417-425.	1.0	3
9	Exploring advance care planning awareness, experiences, and preferences of people with cancer and support people: an Australian online cross-sectional study. Supportive Care in Cancer, 2021, 29, 3677-3688.	1.0	13
10	Gaps in patient-centered follow-up cancer care: a cross sectional study. Journal of Psychosocial Oncology, 2021, 39, 161-172.	0.6	3
11	Oncology patients' and oncology nurses' views on palliative chemotherapy: A cross-sectional comparison. Collegian, 2021, , .	0.6	1
12	Brain cancer patient and support persons' experiences of psychosocial care: a mapping of research outputs. Supportive Care in Cancer, 2021, 29, 5559-5569.	1.0	0
13	Psychosocial wellâ€being of brain cancer patients and support persons: A mapping review of study types over time. European Journal of Cancer Care, 2021, 30, e13446.	0.7	O
14	Participation in future planning by communityâ€dwelling older Australians receiving aged care services: Findings from a crossâ€sectional survey. Australasian Journal on Ageing, 2021, , .	0.4	0
15	Effectiveness of information and communications technology interventions for stroke survivors and their support people: a systematic review. Disability and Rehabilitation, 2021, , 1-16.	0.9	6
16	Oncology nurses' perceptions of advance directives for patients with cancer. Australian Journal of Cancer Nursing, 2021, 23, 439-446.	0.8	3
17	Limitations in health professionals' knowledge of end-of-life law: a cross-sectional survey. BMJ Supportive and Palliative Care, 2021, , bmjspcare-2021-003061.	0.8	8
18	Barriers to the provision of optimal care to dying patients in hospital: a cross-sectional study of nurses' perceptions. Australian Journal of Advanced Nursing, 2021, 38, .	0.4	1

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19	Nurses' perceptions, experiences and involvement in the provision of end-of-life care in acute hospitals: A mapping review of research output, quality and effectiveness. International Journal of Nursing Studies, 2021, 122, 104007.	2.5	4
20	Frailty screening among older adults receiving home care packages: a study of feasibility and prevalence. Australian Journal of Primary Health, 2021, , .	0.4	1
21	The System for Patient Assessment of Cancer Experiences (SPACE): a cross-sectional study examining feasibility and acceptability. Supportive Care in Cancer, 2020, 28, 1737-1745.	1.0	2
22	Nurses' perceptions of and barriers to the optimal endâ€ofâ€life care in hospitals: A crossâ€sectional study. Journal of Clinical Nursing, 2020, 29, 1209-1219.	1.4	20
23	Preferences for End-of-Life Care and Decision Making Among Older and Seriously Ill Inpatients: A Cross-Sectional Study. Journal of Pain and Symptom Management, 2020, 59, 187-196.	0.6	19
24	Where would acute care nurses prefer to receive end-of-life care? a cross-sectional survey. International Journal of Nursing Studies, 2020, 109, 103683.	2.5	1
25	Improving depression outcomes among Australian primary care patients: protocol for a cluster randomised controlled trial. BMJ Open, 2020, 10, e032057.	0.8	3
26	Perceptions of optimal endâ€ofâ€life care in hospitals: A crossâ€sectional study of nurses in three locations. Journal of Advanced Nursing, 2020, 76, 3014-3025.	1.5	5
27	Nurses perspectives on healthcare errors in oncology care: A cross-sectional study. European Journal of Oncology Nursing, 2020, 45, 101741.	0.9	2
28	Nurses' perceptions of open disclosure processes in cancer care: a cross-sectional study. Collegian, 2020, 27, 506-511.	0.6	4
29	Prevalence, perceived barriers and sociodemographic correlates of advance care planning in a sample of outpatients. Australian Journal of Primary Health, 2020, 26, 76.	0.4	1
30	"lt sort of hit me like a baseball bat between the eyes― a qualitative study of the psychosocial experiences of mesothelioma patients and carers. Supportive Care in Cancer, 2019, 27, 631-638.	1.0	22
31	Are older and seriously ill inpatients planning ahead for future medical care?. BMC Geriatrics, 2019, 19, 212.	1.1	14
32	Medical oncology outpatients' preferences and experiences with advanced care planning: a cross-sectional study. BMC Cancer, 2019, 19, 63.	1.1	28
33	Assessing patients' experiences of cancer care across the treatment pathway: a mapping review of recent psychosocial cancer care publications. Supportive Care in Cancer, 2019, 27, 1997-2006.	1.0	10
34	Radiation oncology outpatients' patterns of life expectancy discussions. European Journal of Cancer Care, 2019, 28, e13021.	0.7	1
35	Receipt of information about diet by pregnant women: A cross-sectional study. Women and Birth, 2019, 32, e501-e507.	0.9	7
36	Effectiveness of interventions to increase participation in advance care planning for people with a diagnosis of dementia: A systematic review. Palliative Medicine, 2019, 33, 262-273.	1.3	36

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37	Screening for recommended antenatal risk factors: How long does it take?. Women and Birth, 2018, 31, 489-495.	0.9	6
38	The Right Place at the Right Time: Medical Oncology Outpatients' Perceptions of Location of End-of-Life Care. Journal of the National Comprehensive Cancer Network: JNCCN, 2018, 16, 35-41.	2.3	32
39	Quality versus quantity in end-of-life choices of cancer patients and support persons: a discrete choice experiment. Supportive Care in Cancer, 2018, 26, 3593-3599.	1.0	9
40	A discrete choice experiment to assess cancer patients' preferences for when and how to make treatment decisions. Supportive Care in Cancer, 2018, 26, 1215-1220.	1.0	19
41	Do medical oncology patients and their support persons agree about endâ€ofâ€ife issues?. Internal Medicine Journal, 2018, 48, 60-66.	0.5	5
42	Dances With Denial: Have Medical Oncology Outpatients Conveyed Their End-of-Life Wishes and Do They Want To?. Journal of the National Comprehensive Cancer Network: JNCCN, 2018, 16, 498-505.	2.3	4
43	Knowledge of, and participation in, advance care planning: A cross-sectional study of acute and critical care nurses' perceptions. International Journal of Nursing Studies, 2018, 86, 74-81.	2.5	33
44	Methodological quality of teaching communication skills to undergraduate medical students: a mapping review. BMC Medical Education, 2018, 18, 151.	1.0	18
45	Support persons' preferences for the type of consultation and the format of information provided when making a cancer treatment decision. BMC Research Notes, 2018, 11, 456.	0.6	5
46	Increasing advance personal planning: the need for action at the community level. BMC Public Health, 2018, 18, 606.	1.2	21
47	Advance Care Planning. , 2018, , 295-306.		0
48	The first step in ensuring patient-centred quality of care: ask the patient. European Journal of Cancer Care, 2017, 26, e12435.	0.7	32
49	Distress levels in patients with oropharyngeal vs. non-oropharyngeal squamous cell carcinomas of the head and neck over 1Âyear after diagnosis: a retrospective cohort study. Supportive Care in Cancer, 2017, 25, 3225-3233.	1.0	9
50	Improving hospital-based end of life care processes and outcomes: a systematic review of research output, quality and effectiveness. BMC Palliative Care, 2017, 16, 34.	0.8	23
51	Diet during pregnancy: Women's knowledge of and adherence to food safety guidelines. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2017, 57, 315-322.	0.4	8
52	Computer and telephone delivered interventions to support caregivers of people with dementia: a systematic review of research output and quality. BMC Geriatrics, 2017, 17, 265.	1.1	83
53	Access to clinical trials among oncology patients: results of a cross sectional survey. BMC Cancer, 2017, 17, 653.	1.1	14
54	Involvement of informal caregivers in supporting patients with COPD: a review of intervention studies. International Journal of COPD, 2016, Volume 11, 1587-1596.	0.9	39

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55	What is generated and what is used: a description of public health research output and citation: Table 1. European Journal of Public Health, 2016, 26, 523-525.	0.1	19
56	Women's perceptions of antenatal care: are we following guideline recommended care?. BMC Pregnancy and Childbirth, 2016, 16, 191.	0.9	22
57	The Perspectives of Haematological Cancer Patients on Tissue Banking. Oncology and Therapy, 2016, 4, 91-102.	1.0	0
58	Does screening for physical and psychosocial symptoms vary between medical oncology treatment centres?. Psycho-Oncology, 2016, 25, 521-528.	1.0	8
59	The psychosocial outcomes of individuals with hematological cancers: Are we doing enough high quality research, and what is it telling us?. Critical Reviews in Oncology/Hematology, 2016, 101, 21-31.	2.0	9
60	Burden and Unmet Needs of Caregivers of Chronic Obstructive Pulmonary Disease Patients: A Systematic Review of the Volume and Focus of Research Output. COPD: Journal of Chronic Obstructive Pulmonary Disease, 2016, 13, 662-667.	0.7	30
61	Validation of the PHQ-2 against the PHQ-9 for detecting depression in a large sample of Australian general practice patients. Australian Journal of Primary Health, 2016, 22, 262.	0.4	41
62	Assisting the bereaved: A systematic review of the evidence for grief counselling. Palliative Medicine, 2016, 30, 132-148.	1.3	56
63	A race against time: The dementia epidemic. Archives of Medicine and Health Sciences, 2016, 4, 127.	0.0	2
64	Life expectancy discussions in a multisite sample of Australian medical oncology outpatients. Medical Journal of Australia, 2015, 203, 405-405.	0.8	7
65	Prevalence and correlates of current smoking among medical oncology outpatients. Psycho-Oncology, 2015, 24, 1258-1264.	1.0	5
66	Prevalence and correlates of the unmet supportive care needs of individuals diagnosed with a haematological malignancy. Acta Oncol $\tilde{A}^3$ gica, 2015, 54, 507-514.	0.8	65
67	Preparatory education for cancer patients undergoing surgery: A systematic review of volume and quality of research output over time. Patient Education and Counseling, 2015, 98, 1540-1549.	1.0	70
68	Patient-reported areas for quality improvement in general practice: a cross-sectional survey. British Journal of General Practice, 2015, 65, e312-e318.	0.7	4
69	Medical Oncology Patients: Are They Offered Help and Does ItÂProvide Relief?. Journal of Pain and Symptom Management, 2015, 50, 436-444.	0.6	17
70	The Role of eHealth in Optimizing Preventive Care in the Primary Care Setting. Journal of Medical Internet Research, 2015, 17, e126.	2.1	41
71	Optimizing Patient Preparation and Surgical Experience Using eHealth Technology. JMIR Medical Informatics, 2015, 3, e29.	1.3	32
72	Patient-centred care: making cancer treatment centres accountable. Supportive Care in Cancer, 2014, 22, 1989-1997.	1.0	34

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73	Interventions for preparing patients for chemotherapy and radiotherapy: a systematic review. Supportive Care in Cancer, 2014, 22, 2297-2308.	1.0	54
74	A Longitudinal Analysis of Symptom Clusters in Cancer Patients and Their Sociodemographic Predictors. Journal of Pain and Symptom Management, 2014, 47, 566-578.	0.6	43
75	Screening for distress, the sixth vital sign: examining selfâ€referral in people with cancer over a oneâ€year period. Psycho-Oncology, 2013, 22, 388-395.	1.0	43
76	What goes up does not always come down: patterns of distress, physical and psychosocial morbidity in people with cancer over a one year period. Psycho-Oncology, 2013, 22, 168-176.	1.0	124
77	Risk Factors for Continuous Distress Over a 12-Month Period in Newly Diagnosed Cancer Outpatients. Journal of Psychosocial Oncology, 2013, 31, 489-506.	0.6	28
78	Screening for distress, the sixth vital sign, in lung cancer patients: effects on pain, fatigue, and common problemsâ€"secondary outcomes of a randomized controlled trial. Psycho-Oncology, 2013, 22, 1880-1888.	1.0	46
79	Facilitating Needs-Based Support and Palliative Care for People With Chronic Heart Failure: Preliminary Evidence for the Acceptability, Inter-Rater Reliability, and Validity of a Needs Assessment Tool. Journal of Pain and Symptom Management, 2013, 45, 912-925.	0.6	51
80	Physical, Psychosocial, Relationship, and Economic Burden of Caring for People With Cancer: A Review. Journal of Oncology Practice, 2013, 9, 197-202.	2.5	346
81	Reply: benefits of screening cancer patients for distress still not demonstrated. British Journal of Cancer, 2013, 108, 738-739.	2.9	4
82	Reply: Comment on †Online screening for distress, the 6th vital sign, in newly diagnosed oncology outpatients: randomised controlled trial of computerised vs personalised triage' – Psychological distress in patients with cancer: is screening the effective solution?. British Journal of Cancer, 2013, 108, 2631-2632.	2.9	2
83	Implementing Routine Screening for Distress, the Sixth Vital Sign, for Patients With Head and Neck and Neurologic Cancers. Journal of the National Comprehensive Cancer Network: JNCCN, 2013, 11, 1249-1261.	2.3	34
84	Implementar un programa de screening del malestar emocional asociado al c $\tilde{A}_i$ ncer: ciencia y pr $\tilde{A}_i$ ctica. Psicooncologia, 2013, 9, .	0.1	2
85	Online screening for distress, the 6th vital sign, in newly diagnosed oncology outpatients: randomised controlled trial of computerised vs personalised triage. British Journal of Cancer, 2012, 107, 617-625.	2.9	78
86	Screening for Distress and Unmet Needs in Patients With Cancer: Review and Recommendations. Journal of Clinical Oncology, 2012, 30, 1160-1177.	0.8	456
87	Using Screening for Distress, the Sixth Vital Sign, to advance patient care with assessment and targeted interventions. Supportive Care in Cancer, 2012, 20, 2241-2246.	1.0	8
88	Screening for distress, the 6th vital sign: common problems in cancer outpatients over one year in usual care: associations with marital status, sex, and age. BMC Cancer, 2012, 12, 441.	1.1	69
89	Implications of a needs assessment intervention for people with progressive cancer: Impact on clinical assessment, response and service utilisation. Psycho-Oncology, 2012, 21, 550-557.	1.0	18
90	Improving Outcomes for People With Progressive Cancer: Interrupted Time Series Trial of a Needs Assessment Intervention. Journal of Pain and Symptom Management, 2012, 43, 569-581.	0.6	42

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#	Article	IF	CITATION
91	Reply to S. Palmer et al. Journal of Clinical Oncology, 2011, 29, e279-e280.	0.8	3
92	Validity, reliability and clinical feasibility of a Needs Assessment Tool for people with progressive cancer. Psycho-Oncology, 2010, 19, 726-733.	1.0	44
93	Facilitating needs based cancer care for people with a chronic disease: Evaluation of an intervention using a multi-centre interrupted time series design. BMC Palliative Care, 2010, 9, 2.	0.8	21
94	Development of the Palliative Care Needs Assessment Tool (PC-NAT) for use by multi-disciplinary health professionals. Palliative Medicine, 2008, 22, 956-964.	1,3	79