Jo Brett

List of Publications by Year in descending order

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567281 526287 2,634 28 15 27 citations h-index g-index papers 28 28 28 4084 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Visualizing community networks to recruit South Asian participants for interviews about bowel cancer screening. Journal of Cancer Policy, 2022, 32, 100333.	1.4	2
2	Lifestage differences in young UK women's reasons for research participation. Health Promotion International, 2021, 36, 132-142.	1.8	O
3	Strategies for living well with hormone-responsive advanced prostate cancer—a qualitative exploration. Supportive Care in Cancer, 2021, 29, 1317-1325.	2.2	2
4	Experiences of Support for Sexual Dysfunction in Men with Prostate Cancer: Findings from a U.KWide Mixed Methods Study. Journal of Sexual Medicine, 2021, 18, 515-525.	0.6	4
5	Adjustment strategies amongst black African and black Caribbean men following treatment for prostate cancer: Findings from the Life After Prostate Cancer Diagnosis (LAPCD) study. European Journal of Cancer Care, 2020, 29, e13183.	1.5	8
6	A Qualitative Exploration of Prostate Cancer Survivors Experiencing Psychological Distress: Loss of Self, Function, Connection, and Control. Oncology Nursing Forum, 2020, 47, 318-330.	1.2	9
7	Electronic cigarettes as a smoking cessation aid for patients with cancer: beliefs and behaviours of clinicians in the UK. BMJ Open, 2020, 10, e037637.	1.9	9
8	What if something goes wrong? A grounded theory study of parents' decision-making processes around mode of breech birth at term gestation. Midwifery, 2019, 78, 114-122.	2.3	10
9	The psychological impact of being on a monitoring pathway for localised prostate cancer: A UKâ€wide mixed methods study. Psycho-Oncology, 2019, 28, 1567-1575.	2.3	6
10	Availability and quality assessment of online nutrition information materials for pelvic cancer patients in the UK. European Journal of Cancer Care, 2019, 28, e13039.	1.5	2
11	â€`Very difficult for an ordinary guy': Factors influencing the quality of treatment decision-making amongst men diagnosed with localised and locally advanced prostate cancer: Findings from a UK-wide mixed methods study. Patient Education and Counseling, 2019, 102, 797-803.	2.2	12
12	Experiences and supportive care needs of UK patients with pancreatic cancer: a cross-sectional questionnaire survey. BMJ Open, 2019, 9, e032681.	1.9	20
13	Factors associated with intentional and unintentional non-adherence to adjuvant endocrine therapy following breast cancer. European Journal of Cancer Care, 2018, 27, e12601.	1.5	70
14	Development of an e-health app to support women prescribed adjuvant endocrine therapy after treatment for breast cancer. Patient Preference and Adherence, 2018, Volume 12, 2639-2647.	1.8	19
15	Adjuvant endocrine therapy after breast cancer: a qualitative study of factors associated with adherence. Patient Preference and Adherence, 2018, Volume 12, 291-300.	1.8	36
16	Reaching consensus on reporting patient and public involvement (PPI) in research: methods and lessons learned from the development of reporting guidelines. BMJ Open, 2017, 7, e016948.	1.9	35
17	Psychometric properties of the Beliefs about Medicine Questionnaire–adjuvant endocrine therapy (BMQ-AET) for women taking AETs following early-stage breast cancer. Health Psychology Open, 2017, 4, 205510291774046.	1.4	10
18	Patient and public engagement in health-related quality of life and patient-reported outcomes research: what is important and why should we care? Findings from the first ISOQOL patient engagement symposium. Quality of Life Research, 2015, 24, 1069-1076.	3.1	65

#	Article	IF	CITATION
19	The Warwick Patient Experiences Framework: patient-based evidence in clinical guidelines. International Journal for Quality in Health Care, 2014, 26, 151-157.	1.8	66
20	Mapping the impact of patient and public involvement on health and social care research: a systematic review. Health Expectations, 2014, 17, 637-650.	2.6	986
21	A Systematic Review of the Impact of Patient and Public Involvement on Service Users, Researchers and Communities. Patient, 2014, 7, 387-395.	2.7	471
22	Patient and Public Involvement in Patient-Reported Outcome Measures. Patient, 2012, 5, 79-87.	2.7	147
23	The POPPY Study: Developing a Model of Familyâ€Centred Care for Neonatal Units. Worldviews on Evidence-Based Nursing, 2012, 9, 243-255.	2.9	47
24	Developing the evidence base of patient and public involvement in health and social care research: the case for measuring impact. International Journal of Consumer Studies, 2011, 35, 628-632.	11.6	118
25	The GRIPP checklist: Strengthening the quality of patient and public involvement reporting in research. International Journal of Technology Assessment in Health Care, 2011, 27, 391-399.	0.5	253
26	A systematic mapping review of effective interventions for communicating with, supporting and providing information to parents of preterm infants. BMJ Open, 2011, 1, e000023-e000023.	1.9	109
27	"It's a maybe test": men's experiences of prostate specific antigen testing in primary care. British Journal of General Practice, 2007, 57, 303-10.	1.4	24
28	Breast Screening: Adverse Psychological Consequences One Month after Placing Women on Early Recall Because of a Diagnostic Uncertainty. A Multicentre Study. Journal of Medical Screening, 1997, 4, 158-168	2.3	94