

Jo Brett

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4845150/publications.pdf>

Version: 2024-02-01

28
papers

2,634
citations

567281

15
h-index

526287

27
g-index

28
all docs

28
docs citations

28
times ranked

4084
citing authors

#	ARTICLE	IF	CITATIONS
1	Mapping the impact of patient and public involvement on health and social care research: a systematic review. <i>Health Expectations</i> , 2014, 17, 637-650.	2.6	986
2	A Systematic Review of the Impact of Patient and Public Involvement on Service Users, Researchers and Communities. <i>Patient</i> , 2014, 7, 387-395.	2.7	471
3	The GRIPP checklist: Strengthening the quality of patient and public involvement reporting in research. <i>International Journal of Technology Assessment in Health Care</i> , 2011, 27, 391-399.	0.5	253
4	Patient and Public Involvement in Patient-Reported Outcome Measures. <i>Patient</i> , 2012, 5, 79-87.	2.7	147
5	Developing the evidence base of patient and public involvement in health and social care research: the case for measuring impact. <i>International Journal of Consumer Studies</i> , 2011, 35, 628-632.	11.6	118
6	A systematic mapping review of effective interventions for communicating with, supporting and providing information to parents of preterm infants. <i>BMJ Open</i> , 2011, 1, e000023-e000023.	1.9	109
7	Breast Screening: Adverse Psychological Consequences One Month after Placing Women on Early Recall Because of a Diagnostic Uncertainty. A Multicentre Study. <i>Journal of Medical Screening</i> , 1997, 4, 158-168.	2.3	94
8	Factors associated with intentional and unintentional non-adherence to adjuvant endocrine therapy following breast cancer. <i>European Journal of Cancer Care</i> , 2018, 27, e12601.	1.5	70
9	The Warwick Patient Experiences Framework: patient-based evidence in clinical guidelines. <i>International Journal for Quality in Health Care</i> , 2014, 26, 151-157.	1.8	66
10	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. <i>Quality of Life Research</i> , 2015, 24, 1069-1076.	3.1	65
11	The POPPY Study: Developing a Model of Family-Centred Care for Neonatal Units. <i>Worldviews on Evidence-Based Nursing</i> , 2012, 9, 243-255.	2.9	47
12	Adjuvant endocrine therapy after breast cancer: a qualitative study of factors associated with adherence. <i>Patient Preference and Adherence</i> , 2018, Volume 12, 291-300.	1.8	36
13	Reaching consensus on reporting patient and public involvement (PPI) in research: methods and lessons learned from the development of reporting guidelines. <i>BMJ Open</i> , 2017, 7, e016948.	1.9	35
14	"It's a maybe test": men's experiences of prostate specific antigen testing in primary care. <i>British Journal of General Practice</i> , 2007, 57, 303-10.	1.4	24
15	Experiences and supportive care needs of UK patients with pancreatic cancer: a cross-sectional questionnaire survey. <i>BMJ Open</i> , 2019, 9, e032681.	1.9	20
16	Development of an e-health app to support women prescribed adjuvant endocrine therapy after treatment for breast cancer. <i>Patient Preference and Adherence</i> , 2018, Volume 12, 2639-2647.	1.8	19
17	"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. <i>Patient Education and Counseling</i> , 2019, 102, 797-803.	2.2	12
18	Psychometric properties of the Beliefs about Medicine Questionnaire "adjuvant endocrine therapy (BMQ-AET) for women taking AETs following early-stage breast cancer. <i>Health Psychology Open</i> , 2017, 4, 205510291774046.	1.4	10

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19	What if something goes wrong? A grounded theory study of parents' decision-making processes around mode of breech birth at term gestation. <i>Midwifery</i> , 2019, 78, 114-122.	2.3	10
20	A Qualitative Exploration of Prostate Cancer Survivors Experiencing Psychological Distress: Loss of Self, Function, Connection, and Control. <i>Oncology Nursing Forum</i> , 2020, 47, 318-330.	1.2	9
21	Electronic cigarettes as a smoking cessation aid for patients with cancer: beliefs and behaviours of clinicians in the UK. <i>BMJ Open</i> , 2020, 10, e037637.	1.9	9
22	Adjustment strategies amongst black African and black Caribbean men following treatment for prostate cancer: Findings from the Life After Prostate Cancer Diagnosis (LAPCD) study. <i>European Journal of Cancer Care</i> , 2020, 29, e13183.	1.5	8
23	The psychological impact of being on a monitoring pathway for localised prostate cancer: A UK-wide mixed methods study. <i>Psycho-Oncology</i> , 2019, 28, 1567-1575.	2.3	6
24	Experiences of Support for Sexual Dysfunction in Men with Prostate Cancer: Findings from a U.K.-Wide Mixed Methods Study. <i>Journal of Sexual Medicine</i> , 2021, 18, 515-525.	0.6	4
25	Availability and quality assessment of online nutrition information materials for pelvic cancer patients in the UK. <i>European Journal of Cancer Care</i> , 2019, 28, e13039.	1.5	2
26	Strategies for living well with hormone-responsive advanced prostate cancer—a qualitative exploration. <i>Supportive Care in Cancer</i> , 2021, 29, 1317-1325.	2.2	2
27	Visualizing community networks to recruit South Asian participants for interviews about bowel cancer screening. <i>Journal of Cancer Policy</i> , 2022, 32, 100333.	1.4	2
28	Lifestage differences in young UK women's reasons for research participation. <i>Health Promotion International</i> , 2021, 36, 132-142.	1.8	0