

Penny Wright

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

55
papers

951
citations

18
h-index

29
g-index

57
ext. papers

1,160
ext. citations

4.3
avg, IF

3.91
L-index

#	Paper	IF	Citations
55	Patients report improvements in continuity of care when quality of life assessments are used routinely in oncology practice: secondary outcomes of a randomised controlled trial. <i>European Journal of Cancer</i> , 2010 , 46, 2381-8	7.5	125
54	Health-related quality of life after colorectal cancer in England: a patient-reported outcomes study of individuals 12 to 36 months after diagnosis. <i>Journal of Clinical Oncology</i> , 2015 , 33, 616-24	2.2	86
53	Integrating patient reported outcomes with clinical cancer registry data: a feasibility study of the electronic Patient-Reported Outcomes From Cancer Survivors (ePOCS) system. <i>Journal of Medical Internet Research</i> , 2013 , 15, e230	7.6	60
52	Quality of life in men living with advanced and localised prostate cancer in the UK: a population-based study. <i>Lancet Oncology</i> , 2019 , 20, 436-447	21.7	59
51	A Rasch and factor analysis of the Functional Assessment of Cancer Therapy-General (FACT-G). <i>Health and Quality of Life Outcomes</i> , 2007 , 5, 19	3	41
50	The Social Difficulties Inventory (SDI): development of subscales and scoring guidance for staff. <i>Psycho-Oncology</i> , 2011 , 20, 36-43	3.9	40
49	Psychometric evaluation of the revised Illness Perception Questionnaire (IPQ-R) in cancer patients: confirmatory factor analysis and Rasch analysis. <i>Journal of Psychosomatic Research</i> , 2013 , 75, 556-62	4.1	29
48	Illness perceptions within 6 months of cancer diagnosis are an independent prospective predictor of health-related quality of life 15 months post-diagnosis. <i>Psycho-Oncology</i> , 2015 , 24, 1463-70	3.9	29
47	Health-related quality of life after treatment for bladder cancer in England. <i>British Journal of Cancer</i> , 2018 , 118, 1518-1528	8.7	28
46	Ethnicity and the prostate cancer experience: a qualitative metasynthesis. <i>Psycho-Oncology</i> , 2016 , 25, 1147-1156	3.9	27
45	Feasibility test of a UK-scalable electronic system for regular collection of patient-reported outcome measures and linkage with clinical cancer registry data: the electronic Patient-reported Outcomes From Cancer Survivors (ePOCS) system. <i>BMC Medical Informatics and Decision Making</i> , 2014 , 14, 66	3.6	26
44	Life after prostate cancer diagnosis: protocol for a UK-wide patient-reported outcomes study. <i>BMJ Open</i> , 2016 , 6, e013555	3	25
43	Identifying Social Distress: A Cross-Sectional Survey of Social Outcomes 12 to 36 Months After Colorectal Cancer Diagnosis. <i>Journal of Clinical Oncology</i> , 2015 , 33, 3423-30	2.2	23
42	Functional Outcomes and Health-Related Quality of Life After Curative Treatment for Rectal Cancer: A Population-Level Study in England. <i>International Journal of Radiation Oncology Biology Physics</i> , 2019 , 103, 1132-1142	4	22
41	The economic burden of cancer in the UK: a study of survivors treated with curative intent. <i>Psycho-Oncology</i> , 2016 , 25, 77-83	3.9	22
40	Evaluating patient-reported outcome measures (PROMs) for bladder cancer: a systematic review using the COnsensus-based Standards for the selection of health Measurement INstruments (COSMIN) checklist. <i>BJU International</i> , 2018 , 122, 760-773	5.6	21
39	Validation of an item bank for detecting and assessing psychological distress in cancer patients. <i>Psycho-Oncology</i> , 2009 , 18, 195-9	3.9	18

38	Measuring social difficulties in routine patient-centred assessment: a Rasch analysis of the social difficulties inventory. <i>Quality of Life Research</i> , 2007 , 16, 823-31	3.7	18
37	A Needs-led Framework for Understanding the Impact of Caring for a Family Member With Dementia. <i>Gerontologist, The</i> , 2018 , 58, e68-e77	5	17
36	Factors influencing job loss and early retirement in working men with prostate cancer-findings from the population-based Life After Prostate Cancer Diagnosis (LAPCD) study. <i>Journal of Cancer Survivorship</i> , 2018 , 12, 669-678	5.1	17
35	Developing a useful, user-friendly website for cancer patient follow-up: users' perspectives on ease of access and usefulness. <i>European Journal of Cancer Care</i> , 2012 , 21, 747-57	2.4	15
34	Measurement and interpretation of social distress using the social difficulties inventory (SDI). <i>European Journal of Cancer</i> , 2008 , 44, 1529-35	7.5	15
33	Cancer-related symptoms, mental well-being, and psychological distress in men diagnosed with prostate cancer treated with androgen deprivation therapy. <i>Quality of Life Research</i> , 2019 , 28, 2741-2751	3.7	14
32	Decision regret in men living with and beyond nonmetastatic prostate cancer in the United Kingdom: A population-based patient-reported outcome study. <i>Psycho-Oncology</i> , 2020 , 29, 886-893	3.9	13
31	Traditional and Rasch psychometric analyses of the Quality of Life in Adult Cancer Survivors (QLACS) questionnaire in shorter-term cancer survivors 15 months post-diagnosis. <i>Journal of Psychosomatic Research</i> , 2014 , 77, 322-9	4.1	13
30	Quality of Life After Bladder Cancer: A Cross-sectional Survey of Patient-reported Outcomes. <i>European Urology</i> , 2021 , 79, 621-632	10.2	13
29	Key factors associated with social distress after prostate cancer: Results from the United Kingdom Life after Prostate Cancer diagnosis study. <i>Cancer Epidemiology</i> , 2019 , 60, 201-207	2.8	12
28	Cancer patients' and clinicians' opinions on the best time in secondary care to approach patients for recruitment to longitudinal questionnaire-based research. <i>Supportive Care in Cancer</i> , 2012 , 20, 3365-72	3.9	12
27	Managing social difficulties: roles and responsibilities of patients and staff. <i>Psycho-Oncology</i> , 2012 , 21, 20-8	3.9	12
26	Routine assessment of social difficulties in cancer patients: are we opening Pandora's box?. <i>Supportive Care in Cancer</i> , 2009 , 17, 1425-32	3.9	11
25	Urinary, bowel and sexual health in older men from Northern Ireland. <i>BJU International</i> , 2018 , 122, 845-857	5.7	10
24	Score equivalence of electronic and paper versions of the Social Difficulties Inventory (SDI-21): a randomised crossover trial in cancer patients. <i>Quality of Life Research</i> , 2013 , 22, 1435-40	3.7	8
23	Regional multidisciplinary team intervention programme to improve colorectal cancer outcomes: study protocol for the Yorkshire Cancer Research Bowel Cancer Improvement Programme (YCR BCIP). <i>BMJ Open</i> , 2019 , 9, e030618	3	8
22	Development of an Item Pool for a Needs-Based Measure of Quality of Life of Carers of a Family Member with Dementia. <i>Patient</i> , 2019 , 12, 125-136	3.7	7
21	Transforming discrete choice experiment latent scale values for EQ-5D-3L using the visual analogue scale. <i>European Journal of Health Economics</i> , 2020 , 21, 787-800	3.6	6

20	Regional Variations in Quality of Survival Among Men with Prostate Cancer Across the United Kingdom. <i>European Urology</i> , 2019 , 76, 228-237	10.2	5
19	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	1.7	5
18	Health-related quality of life in cancer survivorship: Predictive power of the Social Difficulties Inventory. <i>Psycho-Oncology</i> , 2017 , 26, 1994-1997	3.9	4
17	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	3.9	4
16	Measuring Quality of Life in Carers of People With Dementia: Development and Psychometric Evaluation of Scales measuring the Impact of DEmentia on CARers (SIDE CAR). <i>Gerontologist, The</i> , 2021 , 61, e1-e11	5	4
15	Translation of the Social Difficulties Inventory (SDI-21) into three South Asian languages and preliminary evaluation of SDI-21(Urdu). <i>Quality of Life Research</i> , 2011 , 20, 431-8	3.7	3
14	Life and bladder cancer: protocol for a longitudinal and cross-sectional patient-reported outcomes study of Yorkshire (UK) patients. <i>BMJ Open</i> , 2019 , 9, e030850	3	3
13	Quality of life among symptomatic compared to PSA-detected prostate cancer survivors - results from a UK wide patient-reported outcomes study. <i>BMC Cancer</i> , 2019 , 19, 947	4.8	2
12	Measurement invariance of the 16-item social distress scale. <i>Quality of Life Research</i> , 2011 , 20, 507-11	3.7	2
11	Influence of deprivation and rurality on patient-reported outcomes of men living with and beyond prostate cancer diagnosis in the UK: A population-based study. <i>Cancer Epidemiology</i> , 2020 , 69, 101830	2.8	2
10	Improvements in measuring the health-related quality of life of cancer patients. <i>Expert Review of Pharmacoeconomics and Outcomes Research</i> , 2006 , 6, 97-105	2.2	1
9	Using international web sites as a cancer resource. <i>Cancer Practice</i> , 2002 , 10, 167-9		1
8	A data flow process for confidential data and its application in a health research project.. <i>PLoS ONE</i> , 2022 , 17, e0262609	3.7	1
7	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	1.1	1
6	Strategies for living well with hormone-responsive advanced prostate cancer-a qualitative exploration. <i>Supportive Care in Cancer</i> , 2021 , 29, 1317-1325	3.9	0
5	Stability of health-related quality of life and morbidity burden from 18 months after diagnosis of prostate cancer: results of a UK-wide population-based outcome cohort.. <i>Supportive Care in Cancer</i> , 2021 , 30, 3151	3.9	0
4	UK General Population Utility Values for the SIDE CAR-D Instrument Measuring the Impact of Caring for People With Dementia. <i>Value in Health</i> , 2020 , 23, 1079-1086	3.3	
3	Using the Clinical Research Network for psychosocial cancer research: lessons learned from two observational studies. <i>BMJ Supportive and Palliative Care</i> , 2014 , 4, 202-211	2.2	

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| 2 | The social difficulties of cancer patients of South Asian Indian and Pakistani origin: a cross-sectional questionnaire and interview study. <i>BMJ Supportive and Palliative Care</i> , 2011 , 1, 154-61 | 2.2 |
| 1 | Collection of cancer Patient Reported Outcome Measures (PROMS) to link with primary and secondary electronic care records to understand and improve long term cancer outcomes: A protocol paper.. <i>PLoS ONE</i> , 2022 , 17, e0266804 | 3.7 |