

# Penny Wright

## List of Publications by Year in Descending Order

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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	A data flow process for confidential data and its application in a health research project.. <i>PLoS ONE</i> , <b>2022</b> , 17, e0262609	3.7	1
54	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> , <b>2022</b> , 17, e0266804	3.7	
53	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> , <b>2021</b> , 18, 515-525	1.1	1
52	Quality of Life After Bladder Cancer: A Cross-sectional Survey of Patient-reported Outcomes. <i>European Urology</i> , <b>2021</b> , 79, 621-632	10.2	13
51	Strategies for living well with hormone-responsive advanced prostate cancer-a qualitative exploration. <i>Supportive Care in Cancer</i> , <b>2021</b> , 29, 1317-1325	3.9	0
50	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> , <b>2021</b> , 61, e1-e11	5	4
49	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> , <b>2021</b> , 30, 3151	3.9	0
48	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> , <b>2020</b> , 23, 1079-1086	3.3	
47	A Qualitative Exploration of Prostate Cancer Survivors Experiencing Psychological Distress: Loss of Self, Function, Connection, and Control. <i>Oncology Nursing Forum</i> , <b>2020</b> , 47, 318-330	1.7	5
46	Transforming discrete choice experiment latent scale values for EQ-5D-3L using the visual analogue scale. <i>European Journal of Health Economics</i> , <b>2020</b> , 21, 787-800	3.6	6
45	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> , <b>2020</b> , 29, 886-893	3.9	13
44	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> , <b>2020</b> , 69, 101830	2.8	2
43	Quality of life in men living with advanced and localised prostate cancer in the UK: a population-based study. <i>Lancet Oncology, The</i> , <b>2019</b> , 20, 436-447	21.7	59
42	The psychological impact of being on a monitoring pathway for localised prostate cancer: A UK-wide mixed methods study. <i>Psycho-Oncology</i> , <b>2019</b> , 28, 1567-1575	3.9	4
41	Regional Variations in Quality of Survival Among Men with Prostate Cancer Across the United Kingdom. <i>European Urology</i> , <b>2019</b> , 76, 228-237	10.2	5
40	Key factors associated with social distress after prostate cancer: Results from the United Kingdom Life after Prostate Cancer diagnosis study. <i>Cancer Epidemiology</i> , <b>2019</b> , 60, 201-207	2.8	12
39	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> , <b>2019</b> , 103, 1132-1142	4	22

38	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> , <b>2019</b> , 28, 2741-2751	3.7	14
37	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> , <b>2019</b> , 19, 947	4.8	2
36	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> , <b>2019</b> , 9, e030618	3	8
35	Life and bladder cancer: protocol for a longitudinal and cross-sectional patient-reported outcomes study of Yorkshire (UK) patients. <i>BMJ Open</i> , <b>2019</b> , 9, e030850	3	3
34	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> , <b>2019</b> , 12, 125-136	3.7	7
33	Urinary, bowel and sexual health in older men from Northern Ireland. <i>BJU International</i> , <b>2018</b> , 122, 845-858	5.7	10
32	A Needs-led Framework for Understanding the Impact of Caring for a Family Member With Dementia. <i>Gerontologist</i> , <b>2018</b> , 58, e68-e77	5	17
31	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> , <b>2018</b> , 122, 760-773	5.6	21
30	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> , <b>2018</b> , 12, 669-678	5.1	17
29	Health-related quality of life after treatment for bladder cancer in England. <i>British Journal of Cancer</i> , <b>2018</b> , 118, 1518-1528	8.7	28
28	Health-related quality of life in cancer survivorship: Predictive power of the Social Difficulties Inventory. <i>Psycho-Oncology</i> , <b>2017</b> , 26, 1994-1997	3.9	4
27	Ethnicity and the prostate cancer experience: a qualitative metasynthesis. <i>Psycho-Oncology</i> , <b>2016</b> , 25, 1147-1156	3.9	27
26	Life after prostate cancer diagnosis: protocol for a UK-wide patient-reported outcomes study. <i>BMJ Open</i> , <b>2016</b> , 6, e013555	3	25
25	The economic burden of cancer in the UK: a study of survivors treated with curative intent. <i>Psycho-Oncology</i> , <b>2016</b> , 25, 77-83	3.9	22
24	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> , <b>2015</b> , 33, 616-24	2.2	86
23	Identifying Social Distress: A Cross-Sectional Survey of Social Outcomes 12 to 36 Months After Colorectal Cancer Diagnosis. <i>Journal of Clinical Oncology</i> , <b>2015</b> , 33, 3423-30	2.2	23
22	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> , <b>2015</b> , 24, 1463-70	3.9	29
21	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> , <b>2014</b> , 77, 322-9	4.1	13

20	Using the Clinical Research Network for psychosocial cancer research: lessons learned from two observational studies. <i>BMJ Supportive and Palliative Care</i> , <b>2014</b> , 4, 202-211	2.2	
19	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> , <b>2013</b> , 22, 1435-40	3.7	8
18	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> , <b>2013</b> , 75, 556-62	4.1	29
17	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> , <b>2013</b> , 15, e230	7.6	60
16	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> , <b>2012</b> , 21, 747-57	2.4	15
15	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> , <b>2012</b> , 20, 3365-72	3.9	12
14	Managing social difficulties: roles and responsibilities of patients and staff. <i>Psycho-Oncology</i> , <b>2012</b> , 21, 20-8	3.9	12
13	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> , <b>2011</b> , 20, 431-8	3.7	3
12	Measurement invariance of the 16-item social distress scale. <i>Quality of Life Research</i> , <b>2011</b> , 20, 507-11	3.7	2
11	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> , <b>2011</b> , 11, 66	3.6	26
10	The Social Difficulties Inventory (SDI): development of subscales and scoring guidance for staff. <i>Psycho-Oncology</i> , <b>2011</b> , 20, 36-43	3.9	40
9	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> , <b>2011</b> , 1, 154-61	2.2	
8	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> , <b>2010</b> , 46, 2381-8	7.5	125
7	Routine assessment of social difficulties in cancer patients: are we opening Pandora's box?. <i>Supportive Care in Cancer</i> , <b>2009</b> , 17, 1425-32	3.9	11
6	Validation of an item bank for detecting and assessing psychological distress in cancer patients. <i>Psycho-Oncology</i> , <b>2009</b> , 18, 195-9	3.9	18
5	Measurement and interpretation of social distress using the social difficulties inventory (SDI). <i>European Journal of Cancer</i> , <b>2008</b> , 44, 1529-35	7.5	15
4	A Rasch and factor analysis of the Functional Assessment of Cancer Therapy-General (FACT-G). <i>Health and Quality of Life Outcomes</i> , <b>2007</b> , 5, 19	3	41
3	Measuring social difficulties in routine patient-centred assessment: a Rasch analysis of the social difficulties inventory. <i>Quality of Life Research</i> , <b>2007</b> , 16, 823-31	3.7	18

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| 2 | Improvements in measuring the health-related quality of life of cancer patients. <i>Expert Review of Pharmacoeconomics and Outcomes Research</i> , <b>2006</b> , 6, 97-105 | 2.2 | 1 |
| 1 | Using international web sites as a cancer resource. <i>Cancer Practice</i> , <b>2002</b> , 10, 167-9   |     | 1 |