

Penny Wright

List of Publications by Year in descending order

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57
papers

1,369
citations

361045
20
h-index

360668
35
g-index

57
all docs

57
docs citations

57
times ranked

2356
citing authors

#	ARTICLE	IF	CITATIONS
1	A data flow process for confidential data and its application in a health research project. PLoS ONE, 2022, 17, e0262609.	1.1	3
2	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. Supportive Care in Cancer, 2022, 30, 3151-3164.	1.0	1
3	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. PLoS ONE, 2022, 17, e0266804.	1.1	2
4	Strategies for living well with hormone-responsive advanced prostate cancer—a qualitative exploration. Supportive Care in Cancer, 2021, 29, 1317-1325.	1.0	2
5	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). Gerontologist, The, 2021, 61, e1-e11.	2.3	8
6	Experiences of Support for Sexual Dysfunction in Men with Prostate Cancer: Findings from a U.K.-Wide Mixed Methods Study. Journal of Sexual Medicine, 2021, 18, 515-525.	0.3	4
7	Quality of Life After Bladder Cancer: A Cross-sectional Survey of Patient-reported Outcomes. European Urology, 2021, 79, 621-632.	0.9	77
8	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. Cancer Epidemiology, 2020, 69, 101830.	0.8	6
9	UK General Population Utility Values for the SIDECAR-D Instrument Measuring the Impact of Caring for People With Dementia. Value in Health, 2020, 23, 1079-1086.	0.1	0
10	A Qualitative Exploration of Prostate Cancer Survivors Experiencing Psychological Distress: Loss of Self, Function, Connection, and Control. Oncology Nursing Forum, 2020, 47, 318-330.	0.5	9
11	Transforming discrete choice experiment latent scale values for EQ-5D-3L using the visual analogue scale. European Journal of Health Economics, 2020, 21, 787-800.	1.4	13
12	Decision regret in men living with and beyond nonmetastatic prostate cancer in the United Kingdom: A population-based patient-reported outcome study. Psycho-Oncology, 2020, 29, 886-893.	1.0	26
13	Cancer-related symptoms, mental well-being, and psychological distress in men diagnosed with prostate cancer treated with androgen deprivation therapy. Quality of Life Research, 2019, 28, 2741-2751.	1.5	21
14	Quality of life among symptomatic compared to PSA-detected prostate cancer survivors - results from a UK wide patient-reported outcomes study. BMC Cancer, 2019, 19, 947.	1.1	4
15	Quality of life in men living with advanced and localised prostate cancer in the UK: a population-based study. Lancet Oncology, The, 2019, 20, 436-447.	5.1	100
16	The psychological impact of being on a monitoring pathway for localised prostate cancer: A UK-wide mixed methods study. Psycho-Oncology, 2019, 28, 1567-1575.	1.0	6
17	Regional Variations in Quality of Survival Among Men with Prostate Cancer Across the United Kingdom. European Urology, 2019, 76, 228-237.	0.9	6
18	Key factors associated with social distress after prostate cancer: Results from the United Kingdom Life after Prostate Cancer diagnosis study. Cancer Epidemiology, 2019, 60, 201-207.	0.8	15

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19	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.	0.4	43
20	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.	0.8	15
21	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.	0.8	6
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.	1.1	12
23	Urinary, bowel and sexual health in older men from Northern Ireland. <i>BJU International</i> , 2018, 122, 845-857.	1.3	18
24	A Needs-led Framework for Understanding the Impact of Caring for a Family Member With Dementia. <i>Gerontologist</i> , The, 2018, 58, e68-e77.	2.3	21
25	Health-related quality of life after treatment for bladder cancer in England. <i>British Journal of Cancer</i> , 2018, 118, 1518-1528.	2.9	55
26	Evaluating patient-reported outcome measures (<sc>PROM</sc>s) for bladder cancer: a systematic review using the <sc>CO</sc>nsensus-based Standards for the selection of health Measurement Instruments (<sc>COSMIN</sc>) checklist. <i>BJU International</i> , 2018, 122, 760-773.	1.3	34
27	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.	1.5	20
28	Changing Current Practice in Urology: Improving Guideline Development and Implementation Through Stakeholder Engagement. <i>European Urology</i> , 2017, 72, 161-163.	0.9	13
29	Health-related quality of life in cancer survivorship: Predictive power of the Social Difficulties Inventory. <i>Psycho-Oncology</i> , 2017, 26, 1994-1997.	1.0	6
30	Maximising Recruitment to Randomised Controlled Trials: The Role of Qualitative Research to Inform Recruitment Challenges. <i>European Urology</i> , 2017, 72, 799-800.	0.9	1
31	The economic burden of cancer in the UK: a study of survivors treated with curative intent. <i>Psycho-Oncology</i> , 2016, 25, 77-83.	1.0	24
32	Ethnicity and the prostate cancer experience: a qualitative metasynthesis. <i>Psycho-Oncology</i> , 2016, 25, 1147-1156.	1.0	46
33	Life after prostate cancer diagnosis: protocol for a UK-wide patient-reported outcomes study. <i>BMJ Open</i> , 2016, 6, e013555.	0.8	27
34	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, 1463-1470.	1.0	38
35	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, 616-624.	0.8	114
36	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, 3423-3430.	0.8	30

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37	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.	0.8	0
38	Traditional and Rasch psychometric analyses of the Quality of Life in Adult Cancer Survivors (QLACS) questionnaire in shorter-term cancer survivors 15months post-diagnosis. <i>Journal of Psychosomatic Research</i> , 2014, 77, 322-329.	1.2	15
39	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, 1435-1440.	1.5	8
40	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, 556-562.	1.2	39
41	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.	2.1	80
42	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, 3365-3372.	1.0	13
43	Managing social difficulties: roles and responsibilities of patients and staff. <i>Psycho-Oncology</i> , 2012, 21, 20-28.	1.0	13
44	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, 747-757.	0.7	21
45	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, 431-438.	1.5	4
46	Measurement invariance of the 16-item social distress scale. <i>Quality of Life Research</i> , 2011, 20, 507-511.	1.5	2
47	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> , 2011, 11, 66.	1.5	28
48	The Social Difficulties Inventory (SDI): development of subscales and scoring guidance for staff. <i>Psycho-Oncology</i> , 2011, 20, 36-43.	1.0	51
49	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, 154-161.	0.8	1
50	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, 2381-2388.	1.3	150
51	Routine assessment of social difficulties in cancer patients: are we opening Pandora's box?. <i>Supportive Care in Cancer</i> , 2009, 17, 1425-1432.	1.0	11
52	Validation of an item bank for detecting and assessing psychological distress in cancer patients. <i>Psycho-Oncology</i> , 2009, 18, 195-199.	1.0	19
53	Measurement and interpretation of social distress using the social difficulties inventory (SDI). <i>European Journal of Cancer</i> , 2008, 44, 1529-1535.	1.3	18
54	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.	1.0	45

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55	Measuring social difficulties in routine patient-centred assessment: a Rasch analysis of the social difficulties inventory. <i>Quality of Life Research</i> , 2007, 16, 823-831.	1.5	23
56	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.	0.7	1
57	Using International Web Sites as a Cancer Resource. <i>Cancer Practice</i> , 2002, 10, 167-169.	0.8	1