

Crispin Jenkinson

List of Publications by Year in descending order

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Version: 2024-02-01

74
papers

7,556
citations

109321
35
h-index

82547
72
g-index

76
all docs

76
docs citations

76
times ranked

9600
citing authors

#	ARTICLE	IF	CITATIONS
1	Short form 36 (SF36) health survey questionnaire: normative data for adults of working age.. BMJ: British Medical Journal, 1993, 306, 1437-1440.	2.3	1,383
2	The development and validation of a short measure of functioning and well being for individuals with Parkinson's disease. Quality of Life Research, 1995, 4, 241-248.	3.1	1,004
3	A shorter form health survey: can the SF-12 replicate results from the SF-36 in longitudinal studies?. Journal of Public Health, 1997, 19, 179-186.	1.8	741
4	The Picker Patient Experience Questionnaire: development and validation using data from in-patient surveys in five countries. International Journal for Quality in Health Care, 2002, 14, 353-358.	1.8	491
5	Criterion validity and reliability of the SF-36 in a population sample. Quality of Life Research, 1994, 3, 7-12.	3.1	407
6	Development and Testing of the UK SF-12. Journal of Health Services Research and Policy, 1997, 2, 14-18.	1.7	364
7	The PDQ-8: Development and validation of a short-form parkinson's disease questionnaire. Psychology and Health, 1997, 12, 805-814.	2.2	351
8	Self-report quality of life measure for people with schizophrenia: The SQLS. British Journal of Psychiatry, 2000, 177, 42-46.	2.8	241
9	Comparison of three measures of quality of life outcome in the evaluation of continuous positive airways pressure therapy for sleep apnoea. Journal of Sleep Research, 1997, 6, 199-204.	3.2	181
10	Randomised controlled trial comparing hospital at home care with inpatient hospital care. I: three month follow up of health outcomes. BMJ: British Medical Journal, 1998, 316, 1786-1791.	2.3	181
11	Evaluating the responsiveness of the endometriosis health profile questionnaire: The EHP-30. Quality of Life Research, 2004, 13, 705-713.	3.1	108
12	Using the SF-36 measure to compare the health impact of multiple sclerosis and Parkinson's disease with normal population health profiles. Journal of Neurology, Neurosurgery and Psychiatry, 2003, 74, 710-714.	1.9	105
13	Should breast reduction surgery be rationed? A comparison of the health status of patients before and after treatment: postal questionnaire survey. BMJ: British Medical Journal, 1996, 313, 454-457.	2.3	101
14	Cross-cultural evaluation of the short form 8-item Parkinson's Disease Questionnaire (PDQ-8): Results from America, Canada, Japan, Italy and Spain. Parkinsonism and Related Disorders, 2007, 13, 22-28.	2.2	100
15	Development and validation of a short measure of health status for individuals with amyotrophic lateral sclerosis/ motor neurone disease: the ALSAQ-40. Journal of Neurology, 1999, 246, III16-III21.	3.6	94
16	Importance of sensitivity to change as a criterion for selecting health status measures.. Quality and Safety in Health Care, 1992, 1, 89-93.	2.5	86
17	How should we evaluate health status? A comparison of three methods in patients presenting with obstructive sleep apnoea. Quality of Life Research, 1998, 7, 95-100.	3.1	86
18	The ALS Health Profile Study: quality of life of amyotrophic lateral sclerosis patients and carers in Europe. Journal of Neurology, 2000, 247, 835-840.	3.6	73

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19	Evidence for the validity and reliability of the ALS assessment questionnaire: The ALSAQ-40. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases, 2000, 1, 33-40.	1.2	73
20	Comparison of UK and US methods for weighting and scoring the SF-36 summary measures. Journal of Public Health, 1999, 21, 372-376.	1.8	72
21	The routine collection of patient-reported outcome measures (PROMs) for long-term conditions in primary care: a cohort survey. BMJ Open, 2014, 4, e003968.	1.9	72
22	Properties of the Picker Patient Experience questionnaire in a randomized controlled trial of long versus short form survey instruments. Journal of Public Health, 2003, 25, 197-201.	1.8	65
23	Measuring change over time: a comparison of results from a global single item of health status and the multi-dimensional SF-36 health status survey questionnaire in patients presenting with menorrhagia. Quality of Life Research, 1994, 3, 317-321.	3.1	58
24	Sensitivity to change of health status measures in a randomized controlled trial: comparison of the COOP charts and the SF-36. Quality of Life Research, 1995, 4, 47-52.	3.1	55
25	The development and validation of a quality of life measure for the carers of people with Parkinson's disease (the PDQ-Carer). Parkinsonism and Related Disorders, 2012, 18, 483-487.	2.2	55
26	Measuring the effects of online health information: Scale validation for the e-Health Impact Questionnaire. Patient Education and Counseling, 2015, 98, 1418-1424.	2.2	55
27	The London handicap scale: a re-evaluation of its validity using standard scoring and simple summation. Journal of Neurology, Neurosurgery and Psychiatry, 2000, 68, 365-367.	1.9	54
28	The current practice of handling and reporting missing outcome data in eight widely used PROMs in RCT publications: a review of the current literature. Quality of Life Research, 2016, 25, 1613-1623.	3.1	53
29	Health-Related Quality-of-Life Measurement in Patients with Parkinson's Disease. Pharmacoeconomics, 1999, 15, 157-165.	3.3	52
30	Use of the short form health survey (SF-36) in patients with amyotrophic lateral sclerosis: tests of data quality, score reliability, response rate and scaling assumptions. Journal of Neurology, 2002, 249, 178-183.	3.6	52
31	Making sense of ambiguity: evaluation in internal reliability and face validity of the SF 36 questionnaire in women presenting with menorrhagia.. Quality and Safety in Health Care, 1996, 5, 9-12.	2.5	51
32	Cross-cultural evaluation of the Parkinson's Disease Questionnaire: tests of data quality, score reliability, response rate, and scaling assumptions in the United States, Canada, Japan, Italy, and Spain. Journal of Clinical Epidemiology, 2003, 56, 843-847.	5.0	45
33	Can item response theory reduce patient burden when measuring health status in neurological disorders? Results from Rasch analysis of the SF-36 physical functioning scale (PF-10). Journal of Neurology, Neurosurgery and Psychiatry, 2001, 71, 220-224.	1.9	41
34	Evaluation of the American version of the 30-item Endometriosis Health Profile (EHP-30). Quality of Life Research, 2008, 17, 1147-1152.	3.1	41
35	Informing the NHS Outcomes Framework: evaluating meaningful health outcomes for children with neurodisability using multiple methods including systematic review, qualitative research, Delphi survey and consensus meeting. Health Services and Delivery Research, 2014, 2, 1-224.	1.4	40
36	The Parkinson's Disease Questionnaire (PDQ-39): evidence for a method of imputing missing data. Age and Ageing, 2006, 35, 497-502.	1.6	36

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37	Change in health status in long-term conditions over a one year period: a cohort survey using patient-reported outcome measures. Health and Quality of Life Outcomes, 2014, 12, 123.	2.4	29
38	Comparison of the sensitivity to change of long and short form pain measures. Quality of Life Research, 1995, 4, 353-357.	3.1	27
39	Measuring the experiences of health care for patients with musculoskeletal disorders (MSD): development of the Picker MSD questionnaire. Scandinavian Journal of Caring Sciences, 2002, 16, 329-333.	2.1	27
40	Patient attitudes to clinical trials: development of a questionnaire and results from asthma and cancer patients. Health Expectations, 2005, 8, 244-252.	2.6	26
41	Long-term benefits in self-reported health status of nasal continuous positive airway pressure therapy for obstructive sleep apnoea. QJM - Monthly Journal of the Association of Physicians, 2001, 94, 95-99.	0.5	25
42	Quality of life and burden of morbidity in primary care users with multimorbidity. Patient Related Outcome Measures, 2018, Volume 9, 103-113.	1.2	24
43	Deriving summary indices of health status from the Amyotrophic Lateral Sclerosis Assessment Questionnaires (ALSAQ-40 and ALSAQ-5). Journal of Neurology, Neurosurgery and Psychiatry, 2003, 74, 242-245.	1.9	23
44	Validation of the Oxford Participation and Activities Questionnaire. Patient Related Outcome Measures, 2016, 7, 73.	1.2	21
45	Quality of life in coeliac disease: item reduction, scale development and psychometric evaluation of the Coeliac Disease Assessment Questionnaire (<scp>CDAQ</scp>). Alimentary Pharmacology and Therapeutics, 2018, 48, 852-862.	3.7	21
46	Evaluation of the Dartmouth COOP Charts in a large-scale community survey in the United Kingdom. Journal of Public Health, 2002, 24, 106-111.	1.8	20
47	Data quality, floor and ceiling effects, and test–retest reliability of the Mild Cognitive Impairment Questionnaire. Patient Related Outcome Measures, 2018, Volume 9, 43-47.	1.2	19
48	Experiences of Using Web-Based and Mobile Technologies to Support Self-Management of Type 2 Diabetes: Qualitative Study. JMIR Diabetes, 2018, 3, e9.	1.9	18
49	Evaluating the psychometric properties of an e-based version of the 39-item Parkinson&TM;s Disease Questionnaire. Health and Quality of Life Outcomes, 2015, 13, 5.	2.4	16
50	Patient reported outcomes. European Journal of Cardiovascular Nursing, 2016, 15, 112-113.	0.9	16
51	Long-Term Conditions Questionnaire (LTCQ): initial validation survey among primary care patients and social care recipients in England. BMJ Open, 2017, 7, e019235.	1.9	15
52	Comparison of the 40-item Amyotrophic Lateral Sclerosis Assessment Questionnaire (ALSAQ-40) with a short-form five-item version (ALSAQ-5) in a longitudinal survey. Clinical Rehabilitation, 2007, 21, 266-272.	2.2	14
53	The Long-Term Conditions Questionnaire: conceptual framework and item development. Patient Related Outcome Measures, 2016, Volume 7, 109-125.	1.2	14
54	The development and validation of a patient-reported quality of life measure for people with mild cognitive impairment. International Psychogeriatrics, 2014, 26, 487-497.	1.0	13

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55	Development of the Oxford Participation and Activities Questionnaire: constructing an item pool. Patient Related Outcome Measures, 2015, 6, 145.	1.2	12
56	Interpreting change scores on the Amyotrophic Lateral Sclerosis Assessment Questionnaire (ALSAQ-40). Clinical Rehabilitation, 2003, 17, 380-385.	2.2	11
57	Comparison of statistical approaches for analyzing incomplete longitudinal patient-reported outcome data in randomized controlled trials. Patient Related Outcome Measures, 2018, Volume 9, 197-209.	1.2	10
58	An evaluation of a near real-time survey for improving patients'™ experiences of the relational aspects of care: a mixed-methods evaluation. Health Services and Delivery Research, 2018, 6, 1-174.	1.4	10
59	Quality of life in coeliac disease: qualitative interviews to develop candidate items for the Coeliac Disease Assessment Questionnaire. Patient Related Outcome Measures, 2018, Volume 9, 211-220.	1.2	9
60	The Amyotrophic Lateral Sclerosis Assessment Questionnaire (ALSAQ-40): Evidence for a method of imputing missing data. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2007, 8, 90-95.	2.1	8
61	The Oxford Participation and Activities Questionnaire: study protocol. Patient Related Outcome Measures, 2013, 5, 1.	1.2	7
62	<p>The Oxford Participation and Activities Questionnaire (Ox-PAQ): development of a short form and index measure</p>. Patient Related Outcome Measures, 2019, Volume 10, 227-232.	1.2	7
63	Withdrawing gluten-free food from prescriptions in England: a mixed-methods study to examine the impact of policy changes on quality of life. Journal of Human Nutrition and Dietetics, 2020, 33, 453-464.	2.5	7
64	Social Class Differences in Morbidity Using the New U.K. National Statistics Socio-Economic Classification: Do Class Differences in Employment Relations Explain Class Differences in Health?. Annals of the New York Academy of Sciences, 1999, 896, 313-315.	3.8	6
65	Measuring mental health in amyotrophic lateral sclerosis (ALS): A comparison of the SF-36 Mental Health Index with the Psychological General Well-Being Index. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases, 2001, 2, 197-201.	1.2	6
66	<p>Measuring improvement in health-status with the Oxford Participation and Activities Questionnaire (Ox-PAQ)</p>. Patient Related Outcome Measures, 2019, Volume 10, 153-156.	1.2	6
67	Burden of cancer trial participation: A qualitative sub-study of the INTERIM feasibility RCT. Chronic Illness, 2023, 19, 81-94.	1.5	6
68	The Relational Aspects of Care Questionnaire: item reduction and scoring using inpatient and accident and emergency data in England. Patient Related Outcome Measures, 2018, Volume 9, 173-181.	1.2	5
69	Healthcare experiences and quality of life of adults with coeliac disease: a cross-sectional study. Journal of Human Nutrition and Dietetics, 2020, 33, 741-751.	2.5	5
70	Administering the Routine Activities domain of the Oxford Participation and Activities Questionnaire as a stand-alone scale: the Oxford Routine Activities Measure. Patient Related Outcome Measures, 2018, Volume 9, 239-243.	1.2	2
71	Development and validation of a short measure of health status for individuals with amyotrophic lateral sclerosis/ motor neurone disease: the ALSAQ-40. Journal of Neurology, 1999, 246, s016-s021.	3.6	2
72	Use of the Oxford Participation and Activities Questionnaire in chronic obstructive pulmonary disease. Chronic Respiratory Disease, 2019, 16, 147997311986221.	2.4	1

#	ARTICLE	IF	CITATIONS
73	Web-Based and mHealth Technologies to Support Self-Management in People Living With Type 2 Diabetes: Validation of the Diabetes Self-Management and Technology Questionnaire (DSMT-Q). JMIR Diabetes, 2020, 5, e18208.	1.9	1
74	Measures of health status-the nottingham health profile and the SF-36. Critical Public Health, 1993, 4, 15-20.	2.4	0