## Stephen P Mckenna

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Application of Rasch Analysis in the Development and Application of Quality of Life Instruments. Value in Health, 2004, 7, S22-S26.	0.3	350
2	The QLDS: A scale for the measurement of quality of life in depression. Health Policy, 1992, 22, 307-319.	3.0	285
3	Measuring patient-reported outcomes: moving beyond misplaced common sense to hard science. BMC Medicine, 2011, 9, 86.	5.5	238
4	The QoL-AGHDA: an instrument for the assessment of quality of life in adults with growth hormone deficiency. Quality of Life Research, 1999, 8, 373-383.	3.1	225
5	Adapting Quality of Life Instruments. Value in Health, 2004, 7, S27-S30.	0.3	178
6	Defining Patient-Reported Outcomes. Value in Health, 2004, 7, S4-S8.	0.3	146
7	The Translation and Cultural Adaptation of Patient-Reported Outcome Measures. Value in Health, 2005, 8, 89-91.	0.3	104
8	Can We Rely on the Dermatology Life Quality Index as a Measure of the Impact of Psoriasis or Atopic Dermatitis?. Journal of Investigative Dermatology, 2012, 132, 76-84.	0.7	99
9	The Benefit of Pimecrolimus (Elidel, SDZ ASM 981) on Parents' Quality of Life in the Treatment of Pediatric Atopic Dermatitis. Pediatrics, 2002, 110, 1133-1136.	2.1	80
10	A new measure of quality of life in depression: Testing the reliability and construct validity of the QLDS. Health Policy, 1992, 22, 321-330.	3.0	78
11	Effects of Method of Translation of Patient-Reported Health Outcome Questionnaires: A Randomized Study of the Translation of the Rheumatoid Arthritis Quality of Life (RAQoL) Instrument for Sweden. Value in Health, 2010, 13, 424-430.	0.3	77
12	International development of the Parents' Index of Quality of Life in Atopic Dermatitis (PIQoL-AD). Quality of Life Research, 2005, 14, 231-241.	3.1	76
13	Quality of life of children with atopic dermatitis and their families. Current Opinion in Allergy and Clinical Immunology, 2008, 8, 228-231.	2.3	60
14	Development and validation of a needsâ€based quality of life instrument for osteoarthritis. Arthritis and Rheumatism, 2008, 59, 841-848.	6.7	53
15	Development and psychometric assessment of the COPD and Asthma Sleep Impact Scale (CASIS). Health and Quality of Life Outcomes, 2009, 7, 98.	2.4	49
16	Synchrony of change in depressive symptoms, health status, and quality of life in persons with clinical depression. Health and Quality of Life Outcomes, 2006, 4, 27.	2.4	43
17	Development and validation of a preference based measure derived from the Cambridge Pulmonary Hypertension Outcome Review (CAMPHOR) for use in cost utility analyses. Health and Quality of Life Outcomes, 2008, 6, 65.	2.4	41
18	Needs-based quality of life in adults dependent on home parenteral nutrition. Clinical Nutrition, 2019, 38, 1433-1438.	5.0	39

Stephen P Mckenna

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19	The Development and Psychometric Properties of the MSQOL. Clinical Drug Investigation, 1998, 15, 413-423.	2.2	38
20	Translation and validation of non-English versions of the Ankylosing Spondylitis Quality of Life (ASQOL) questionnaire. Health and Quality of Life Outcomes, 2007, 5, 7.	2.4	37
21	The Needs-Based Approach to Quality of Life Assessment. Value in Health, 2004, 7, S1-S3.	0.3	35
22	Development and validation of the Parenteral Nutrition Impact Questionnaire (PNIQ), a patient-centric outcome measure for Home Parenteral Nutrition. Clinical Nutrition, 2018, 37, 978-983.	5.0	35
23	Dimensionality of the Dermatology Life Quality Index (DLQI): A Commentary. Acta Dermato-Venereologica, 2006, 86, 284-285.	1.3	34
24	The German adaptation of the Cambridge pulmonary hypertension outcome review (CAMPHOR). Health and Quality of Life Outcomes, 2012, 10, 110.	2.4	34
25	Assessing quality of life in Crohn's disease: development and validation of the Crohn's Life Impact Questionnaire (CLIQ). Quality of Life Research, 2015, 24, 2279-2288.	3.1	34
26	Quality of life in infants and children with atopic dermatitis: Addressing issues of differential item functioning across countries in multinational clinical trials. Health and Quality of Life Outcomes, 2007, 5, 45.	2.4	33
27	Measurement of patient-reported outcomes. 2: Are current measures failing us?. Journal of Medical Economics, 2019, 22, 523-530.	2.1	33
28	Measurement of patient-reported outcomes. 1: The search for the Holy Grail. Journal of Medical Economics, 2019, 22, 516-522.	2.1	31
29	Development and validation of the living with chronic obstructive pulmonary disease questionnaire. Quality of Life Research, 2011, 20, 1043-1052.	3.1	29
30	A qualitative study of the impact of Crohn's disease from a patient's perspective: TableÂ1. Frontline Gastroenterology, 2017, 8, 68-73.	1.8	29
31	Composite outcome measurement in clinical research: the triumph of illusion over reality?. Journal of Medical Economics, 2020, 23, 1196-1204.	2.1	27
32	International Development of the Patient-Reported Outcome Indices for Multiple Sclerosis (PRIMUS). Value in Health, 2010, 13, 946-951.	0.3	25
33	The impact of Home Parenteral Nutrition on the lives of adults with Type 3 Intestinal Failure. Clinical Nutrition ESPEN, 2018, 24, 35-40.	1.2	24
34	Patient value: its nature, measurement, and role in real world evidence studies and outcomes-based reimbursement. Journal of Medical Economics, 2018, 21, 474-480.	2.1	23
35	Integrating Patient-Reported Outcomes. Value in Health, 2004, 7, S9-S12.	0.3	21

 $_{36}$  Interpreting scores on multiple sclerosis-specific patient reported outcome measures (the PRIMUS and) Tj ETQq0 0.0 gBT /Overlock 10

STEPHEN P MCKENNA

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37	COPD and Asthma Fatigue Scale (CAFS): Development and Psychometric Assessment. Health Outcomes Research in Medicine, 2010, 1, e5-e16.	0.6	20
38	Setting and maintaining standards for patient-reported outcome measures: can we rely on the COSMIN checklists?. Journal of Medical Economics, 2021, 24, 502-511.	2.1	18
39	Assessing the quality of life of patients in phases I and II anti-cancer drug trials: Interviews versus questionnaires by Cox K. Social Science and Medicine, 2004, 58, 659-660.	3.8	17
40	Development of the US PSORIQoL: a psoriasis-specific measure of quality of life. International Journal of Dermatology, 2005, 44, 462-469.	1.0	16
41	Adaptation of the rheumatoid arthritis quality of life scale for Estonia. Rheumatology International, 2006, 26, 655-662.	3.0	16
42	International Development of the Unidimensional Fatigue Impact Scale (U-FIS). Value in Health, 2010, 13, 463-468.	0.3	16
43	Development and validation of the Asthma Life Impact Scale (ALIS). Respiratory Medicine, 2010, 104, 633-643.	2.9	16
44	Measurement, modeling and QALYs. F1000Research, 2020, 9, 1048.	1.6	16
45	Adaptation of Chinese and English versions of the Ankylosing Spondylitis quality of life (ASQoL) scale for use in Singapore. BMC Musculoskeletal Disorders, 2017, 18, 353.	1.9	14
46	Adaptation of Chinese and English versions of the Psoriatic Arthritis Quality of Life (PsAQoL) scale for use in Singapore. BMC Musculoskeletal Disorders, 2016, 17, 432.	1.9	13
47	Fundamental Measurement: The Need Fulfilment Quality of Life (N-QOL) Measure. Innovations in Pharmacy, 2021, 12, 6.	0.6	11
48	Summary of Needs-Based Quality of Life Instruments. Value in Health, 2004, 7, S39-S40.	0.3	9
49	Further international adaptation and validation of the Rheumatoid Arthritis Quality of Life (RAQoL) questionnaire. Rheumatology International, 2015, 35, 669-675.	3.0	8
50	Portuguese validation of the Cambridge pulmonary hypertension outcome review (CAMPHOR) questionnaire. Health and Quality of Life Outcomes, 2016, 14, 110.	2.4	8
51	Adaptation and validation of the quality of life assessment of the Cambridge pulmonary hypertension outcome review (CAMPHOR) for Brazil. Journal of Patient-Reported Outcomes, 2020, 4, 43.	1.9	7
52	A Review of the Psychometric Properties and Use of the Rheumatoid Arthritis Quality of Life Questionnaire (RAQoL) in Clinical Research. Current Rheumatology Reviews, 2017, 13, 197-205.	0.8	7
53	Adaptation of the osteoarthritis-specific quality of life scale (the OAQoL) for use in Germany, Hungary, Italy, Spain and Turkey. Rheumatology International, 2017, 37, 727-734.	3.0	6
54	Fundamental Outcome Measurement: Selecting Patient Reported Outcome Instruments and Interpreting the Data they Produce. Innovations in Pharmacy, 2021, 12, 17.	0.6	6

STEPHEN P MCKENNA

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55	Measuring the Impact of Caring for a Spouse with Alzheimer's Disease: Validation of the Alzheimer's Patient Partners Life Impact Questionnaire (APPLIQue). Journal of Applied Measurement, 2018, 19, 271-282.	0.3	6
56	Functional disability and quality of life in patients with ankylosing spondylitis. Rheumatology International, 2004, 24, 57-58.	3.0	5
57	International Development of the Alzheimer's Patient Partners Life Impact Questionnaire (APPLIQue). American Journal of Alzheimer's Disease and Other Dementias, 2020, 35, 153331752095169.	1.9	5
58	COSMIN reviews: the need to consider measurement theory, modern measurement and a prospective rather than retrospective approach to evaluating patient-based measures. Journal of Medical Economics, 2021, 24, 860-861.	2.1	5
59	The living with chronic obstructive pulmonary disease scale was successfully adapted for use in Southern European (Italian and Spanish) and Eastern European (Russian) cultures. Journal of Clinical Epidemiology, 2012, 65, 906-914.	5.0	4
60	Improving Scoring Precision and Internal Construct Validity of the Bath Ankylosing Spondylitis Disease Activity Index Using Rasch Measurement Theory. Journal of Rheumatology, 2020, 47, 354-361.	2.0	4
61	Psoriatic Arthritis Quality of Life Questionnaire: translation, cultural adaptation and validation into Brazilian Portuguese language. Advances in Rheumatology, 2021, 61, 13.	1.7	4
62	Development and validation of the Bowel Cleansing Impact Review (BOCLIR). Frontline Gastroenterology, 2013, 4, 112-119.	1.8	3
63	Adaptation and validation of the Rheumatoid Arthritis Quality of Life (RAQoL) questionnaire for use in Serbia. Rheumatology International, 2017, 37, 641-646.	3.0	3
64	A qualitative study of the impact of plexiform neurofibromas on need fulfilment in adults with neurofibromatosis type 1. SAGE Open Medicine, 2019, 7, 205031211982968.	1.8	3
65	Translation, cross-cultural adaptation and validation of the Osteoarthritis Quality of Life (OAQoL) questionnaire for use in Portugal. Rheumatology International, 2019, 39, 715-722.	3.0	3
66	The development of the PlexiQoL: A patientâ€reported outcome measure for adults with neurofibromatosis type 1â€associated plexiform neurofibromas. Molecular Genetics & Genomic Medicine, 2020, 8, e1530.	1.2	3
67	Adaptation of the QoL-AGHDA scale for adults with growth hormone deficiency in four Slavic languages. Health and Quality of Life Outcomes, 2011, 9, 60.	2.4	2
68	Adaptation and validation of the Rheumatoid Arthritis Quality of Life (RAQoL) questionnaire for use in Bulgaria. Rheumatology International, 2020, 40, 2077-2083.	3.0	1
69	Evaluation of the Unidimensional Fatigue Impact Scale (U-FIS) in Crohn's Disease: The Importance of Local Item Dependency. Journal of Nursing Measurement, 2021, , JNM-D-20-00116.	0.3	1
70	The Polish adaptation of the CAMbridge Pulmonary Hypertension Outcome Review (CAMPHOR). Cardiology Journal, 2020, 27, 608-615.	1.2	1
71	It is time for an organised, scientific approach to the application of patient-reported outcome measures in clinical studies and trials. Clinical Rheumatology, 2019, 38, 611-615.	2.2	0
72	Assessment of the performance of the Brazilian Portuguese Nottingham Health Profile in adult growth hormone deficiency and pulmonary hypertension. F1000Research, 2020, 9, 1399.	1.6	0

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73	Assessment of the performance of the Brazilian Portuguese Nottingham Health Profile in adult growth hormone deficiency and pulmonary hypertension. F1000Research, 2020, 9, 1399.	1.6	0