Donald L Patrick

List of Publications by Year in descending order

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456 papers 55,304 citations

107 h-index 222 g-index

463 all docs

463 docs citations

times ranked

463

43079 citing authors

#	Article	IF	CITATIONS
1	Measuring Health-Related Quality of Life. Annals of Internal Medicine, 1993, 118, 622.	2.0	2,982
2	The COSMIN study reached international consensus on taxonomy, terminology, and definitions of measurement properties for health-related patient-reported outcomes. Journal of Clinical Epidemiology, 2010, 63, 737-745.	2.4	2,945
3	The COSMIN checklist for assessing the methodological quality of studies on measurement properties of health status measurement instruments: an international Delphi study. Quality of Life Research, 2010, 19, 539-549.	1.5	2,751
4	Assessing health status and quality-of-life instruments: attributes and review criteria. Quality of Life Research, 2002, 11, 193-205.	1.5	1,959
5	COSMIN guideline for systematic reviews of patient-reported outcome measures. Quality of Life Research, 2018, 27, 1147-1157.	1.5	1,600
6	Generic and Disease-Specific Measures in Assessing Health Status and Quality of Life. Medical Care, 1989, 27, S217-S232.	1.1	1,496
7	The validity of self-reported smoking: a review and meta-analysis American Journal of Public Health, 1994, 84, 1086-1093.	1.5	1,468
8	COSMIN Risk of Bias checklist for systematic reviews of Patient-Reported Outcome Measures. Quality of Life Research, 2018, 27, 1171-1179.	1.5	1,264
9	The COSMIN checklist for evaluating the methodological quality of studies on measurement properties: A clarification of its content. BMC Medical Research Methodology, 2010, 10, 22.	1.4	1,178
10	Reproducibility and responsiveness of health status measures statistics and strategies for evaluation. Contemporary Clinical Trials, 1991, 12, S142-S158.	2.0	1,166
11	COSMIN methodology for evaluating the content validity of patient-reported outcome measures: a Delphi study. Quality of Life Research, 2018, 27, 1159-1170.	1.5	1,020
12	Content Validityâ€"Establishing and Reporting the Evidence in Newly Developed Patient-Reported Outcomes (PRO) Instruments for Medical Product Evaluation: ISPOR PRO Good Research Practices Task Force Report: Part 2â€"Assessing Respondent Understanding. Value in Health, 2011, 14, 978-988.	0.1	766
13	Content Validityâ€"Establishing and Reporting the Evidence in Newly Developed Patient-Reported Outcomes (PRO) Instruments for Medical Product Evaluation: ISPOR PRO Good Research Practices Task Force Report: Part 1â€"Eliciting Concepts for a New PRO Instrument. Value in Health, 2011, 14, 967-977.	0.1	747
14	Assessing Health-Related Quality of Life in Patients With Sciatica. Spine, 1995, 20, 1899-1908.	1.0	666
15	Quality of life in persons with irritable bowel syndrome: development and validation of a new measure. Digestive Diseases and Sciences, 1998, 43, 400-411.	1.1	594
16	Patient-Reported Outcomes to Support Medical Product Labeling Claims: FDA Perspective. Value in Health, 2007, 10, S125-S137.	0.1	583
17	Depressive Symptoms and the Cost of Health Services in HMO Patients Aged 65 Years and Older. JAMA - Journal of the American Medical Association, 1997, 277, 1618.	3.8	546
18	GRADE guidelines: 13. Preparing Summary of Findings tables and evidence profiles—continuous outcomes. Journal of Clinical Epidemiology, 2013, 66, 173-183.	2.4	495

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19	Evaluating quality-of-life and health status instruments: development of scientific review criteria. Clinical Therapeutics, 1996, 18, 979-992.	1.1	482
20	Family satisfaction with family conferences about end-of-life care in the intensive care unit: Increased proportion of family speech is associated with increased satisfaction*. Critical Care Medicine, 2004, 32, 1484-1488.	0.4	472
21	A Prospective Study of Advance Directives for Life-Sustaining Care. New England Journal of Medicine, 1991, 324, 882-888.	13.9	464
22	Depressive symptoms and the cost of health services in HMO patients aged 65 years and older. A 4-year prospective study. JAMA - Journal of the American Medical Association, 1997, 277, 1618-1623.	3.8	449
23	Quality of life of persons with urinary incontinence: Development of a new measure. Urology, 1996, 47, 67-71.	0.5	441
24	Quality of life of women with urinary incontinence: further development of the incontinence quality of life instrument (I-QOL). Urology, 1999, 53, 71-76.	0.5	419
25	Estimating clinically significant differences in quality of life outcomes. Quality of Life Research, 2005, 14, 285-295.	1.5	406
26	Health-related quality of life in inflammatory bowel disease. Digestive Diseases and Sciences, 1989, 34, 1379-1386.	1.1	383
27	A Measure of the Quality of Dying and Death. Journal of Pain and Symptom Management, 2002, 24, 17-31.	0.6	379
28	ORIGINAL RESEARCHâ€"EJACULATORY DISORDERS: Premature Ejaculation: An Observational Study of Men and Their Partners. Journal of Sexual Medicine, 2005, 2, 358-367.	0.3	374
29	Validation of the United States' version of the World Health Organization Quality of Life (WHOQOL) instrument. Journal of Clinical Epidemiology, 2000, 53, 1-12.	2.4	370
30	The family conference as a focus to improve communication about end-of-life care in the intensive care unit: Opportunities for improvement. Critical Care Medicine, 2001, 29, N26-N33.	0.4	345
31	Further validation of the IBS-QOL: a disease-specific quality-of-life questionnaire. American Journal of Gastroenterology, 2000, 95, 999-1007.	0.2	340
32	The Qualityity-of-Life (QOL) Research Movement: Past, Present, and Future. Social Indicators Research, 2006, 76, 343-466.	1.4	324
33	Pediatric Patient-Reported Outcome Instruments for Research to Support Medical Product Labeling: Report of the ISPOR PRO Good Research Practices for the Assessment of Children and Adolescents Task Force. Value in Health, 2013, 16, 461-479.	0.1	322
34	Measuring Preferences for Health States Worse than Death. Medical Decision Making, 1994, 14, 9-18.	1.2	315
35	Self-Efficacy and Health Behavior Among Older Adults. Journal of Health and Social Behavior, 1993, 34, 89.	2.7	308
36	The Maine Lumbar Spine Study, Part II. Spine, 1996, 21, 1777-1786.	1.0	305

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37	Use of Existing Patient-Reported Outcome (PRO) Instruments and Their Modification: The ISPOR Good Research Practices for Evaluating and Documenting Content Validity for the Use of Existing Instruments and Their Modification PRO Task Force Report. Value in Health, 2009, 12, 1075-1083.	0.1	305
38	Seven chronic conditions: their impact on US adults' activity levels and use of medical services American Journal of Public Health, 1995, 85, 173-182.	1.5	299
39	The rating form of IBD patient concerns: a new measure of health status Psychosomatic Medicine, 1991, 53, 701-712.	1.3	298
40	The Concept of Quality of Life of Dying Persons in the Context of Health Care. Journal of Pain and Symptom Management, 1999, 17, 93-108.	0.6	295
41	The Maine Lumbar Spine Study, Part III. Spine, 1996, 21, 1787-1794.	1.0	287
42	Sleep problems and their correlates in a working population. Journal of General Internal Medicine, 1995, 10, 25-32.	1.3	284
43	Advancing Symptom Science Through Symptom Cluster Research: Expert Panel Proceedings and Recommendations. Journal of the National Cancer Institute, 2017, 109, djw253.	3.0	275
44	Evaluating the Quality of Dying and Death. Journal of Pain and Symptom Management, 2001, 22, 717-726.	0.6	269
45	The German version of the Oral Health Impact Profile - translation and psychometric properties. European Journal of Oral Sciences, 2002, 110, 425-433.	0.7	258
46	Assessment of Health-Related Quality of Life in Patients With Interstitial Lung Disease. Chest, 1999, 116, 1175-1182.	0.4	252
47	Social functioning as an outcome measure in schizophrenia studies. Acta Psychiatrica Scandinavica, 2007, 116, 403-418.	2.2	251
48	Prevalence of Secondary Conditions Among People With Disabilities. American Journal of Public Health, 2004, 94, 443-445.	1.5	249
49	Barriers and Facilitators to End-of-Life Care Communication for Patients with COPD. Chest, 2005, 127, 2188-2196.	0.4	245
50	Life-sustaining treatments during terminal illness. Journal of General Internal Medicine, 1993, 8, 361-368.	1.3	242
51	Protocol of the COSMIN study: COnsensus-based Standards for the selection of health Measurement INstruments. BMC Medical Research Methodology, 2006, 6, 2.	1.4	238
52	Patient-physician communication about end-of-life care for patients with severe COPD. European Respiratory Journal, 2004, 24, 200-205.	3.1	232
53	Measurement of Health Status in the 1990s. Annual Review of Public Health, 1990, 11, 165-183.	7.6	222
54	Quality of life following intensive care. Journal of General Internal Medicine, 1988, 3, 218-223.	1.3	217

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55	Barriers to the Use of Health Status Measures in Clinical Investigation, Patient Care, and Policy Research. Medical Care, 1989, 27, S254-S268.	1.1	216
56	Inter-rater agreement and reliability of the COSMIN (COnsensus-based Standards for the selection of) Tj ETQq0	0 0 ₁ .4BT /	Overlock 10 T
57	Stability of Choices about Life-Sustaining Treatments. Annals of Internal Medicine, 1994, 120, 567.	2.0	211
58	Toward an Operational Definition of Health. Journal of Health and Social Behavior, 1973, 14, 6.	2.7	205
59	COSMIN Risk of Bias tool to assess the quality of studies on reliability or measurement error of outcome measurement instruments: a Delphi study. BMC Medical Research Methodology, 2020, 20, 293.	1.4	205
60	Assessment of Community Functioning in People With Schizophrenia and Other Severe Mental Illnesses: A White Paper Based on an NIMH-Sponsored Workshop. Schizophrenia Bulletin, 2007, 33, 805-822.	2.3	201
61	Tranexamic Acid Treatment for Heavy Menstrual Bleeding. Obstetrics and Gynecology, 2010, 116, 865-875.	1.2	195
62	Clinical Outcome Assessments: Conceptual Foundationâ€"Report of the ISPOR Clinical Outcomes Assessment â€" Emerging Good Practices for Outcomes Research Task Force. Value in Health, 2015, 18, 741-752.	0.1	195
63	Studying communication about end-of-life care during the ICU family conference: Development of a framework. Journal of Critical Care, 2002, 17, 147-160.	1.0	186
64	The Psychological Burden of Premature Ejaculation. Journal of Urology, 2007, 177, 1065-1070.	0.2	185
65	Why Don't Patients and Physicians Talk About End-of-Life Care?: Barriers to Communication for Patients With Acquired Immunodeficiency Syndrome and Their Primary Care Clinicians. Archives of Internal Medicine, 2000, 160, 1690-1696.	4.3	184
66	Adolescent quality of life, Part II: initial validation of a new instrument. Journal of Adolescence, 2002, 25, 287-300.	1.2	183
67	The Role of Technical Advances in the Adoption and Integration of Patient-reported Outcomes in Clinical Care. Medical Care, 2015, 53, 153-159.	1.1	183
68	Validation of a new quality of life questionnaire for benign prostatic hyperplasia. Journal of Clinical Epidemiology, 1992, 45, 1431-1445.	2.4	177
69	Untreated Anxiety Among Adult Primary Care Patients in a Health Maintenance Organization. Archives of General Psychiatry, 1994, 51, 740.	13.8	175
70	Assessing Meaningful Change in Quality of Life Over Time: A Users' Guide for Clinicians. Mayo Clinic Proceedings, 2002, 77, 561-571.	1.4	175
71	Quality Adjusted Life Years in Older Adults With Depressive Symptoms and Chronic Medical Disorders. International Psychogeriatrics, 2000, 12, 15-33.	0.6	173
72	Selected Methodological Issues in Evaluating Community-Based Health Promotion and Disease Prevention Programs. Annual Review of Public Health, 1992, 13, 31-57.	7.6	168

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73	Longitudinal investigation of depression outcomes in primary care in six countries: the LIDO Study. Functional status, health service use and treatment of people with depressive symptoms. Psychological Medicine, 2002, 32, 889-902.	2.7	168
74	Health status and health care use in persons with inflammatory bowel disease. Digestive Diseases and Sciences, 1991, 36, 1746-1755.	1.1	167
75	Premature Ejaculation: Results from a Five-Country European Observational Study. European Urology, 2008, 53, 1048-1057.	0.9	164
76	Inclusion of patient-reported outcome measures in registered clinical trials: Evidence from ClinicalTrials.gov (2007–2013). Contemporary Clinical Trials, 2015, 43, 1-9.	0.8	162
77	Validation of the Multiple Sclerosis International Quality of Life questionnaire. Multiple Sclerosis Journal, 2008, 14, 219-230.	1.4	159
78	Reproducibility and Responsiveness of the VF-14. JAMA Ophthalmology, 1995, 113, 1508.	2.6	157
79	Adolescent quality of life, Part I: conceptual and measurement model. Journal of Adolescence, 2002, 25, 275-286.	1.2	156
80	Measurement of Health Outcomes in Treatment Effectiveness Evaluations. Medical Care, 2000, 38, II-14-II-25.	1.1	154
81	Seattle 5 a Day Worksite Program to Increase Fruit and Vegetable Consumption. Preventive Medicine, 2001, 32, 230-238.	1.6	150
82	The mathematical relationship among different forms of responsiveness coefficients. Quality of Life Research, 2007, 16, 815-822.	1.5	149
83	Evaluation of the methodological quality of systematic reviews of health status measurement instruments. Quality of Life Research, 2009, 18, 313-333.	1.5	146
84	The impact of onychomycosis on quality of life: Development of an international onychomycosis-specific questionnaire to measure patient quality of life. Journal of the American Academy of Dermatology, 1999, 41, 189-196.	0.6	143
85	The Quebec Task Force Classification for Spinal Disorders and the Severity, Treatment, and Outcomes of Sciatica and Lumbar Spinal Stenosis. Spine, 1996, 21, 2885-2892.	1.0	142
86	Issues in the measurement of satisfaction with treatment. American Journal of Managed Care, 1997, 3, 579-94.	0.8	140
87	Measuring and Improving the Quality of Dying and Death. Annals of Internal Medicine, 2003, 139, 410.	2.0	138
88	The Premature Ejaculation Profile: validation of selfâ€reported outcome measures for research and practice. BJU International, 2009, 103, 358-364.	1.3	138
89	Clinical Sequencing Exploratory Research Consortium: Accelerating Evidence-Based Practice of Genomic Medicine. American Journal of Human Genetics, 2016, 98, 1051-1066.	2.6	137
90	Methods for measuring levels of well-being for a health status index. Health Services Research, 1973, 8, 228-45.	1.0	137

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91	The Caregiver Activity Survey (CAS): development and validation of a new measure for caregivers of persons with Alzheimer's disease. International Journal of Geriatric Psychiatry, 1997, 12, 978-988.	1.3	135
92	Quality of Dying and Death in Two Medical ICUs. Chest, 2005, 127, 1775-1783.	0.4	135
93	The Cost-effectiveness of Lung Transplantation. Chest, 1995, 108, 1594-1601.	0.4	133
94	Issues in quality-of-life measurement in clinical trials. Contemporary Clinical Trials, 1991, 12, S81-S90.	2.0	129
95	Pain outcomes in patients with advanced breast cancer and bone metastases. Cancer, 2013, 119, 832-838.	2.0	126
96	The use of patient-reported outcomes instruments in registered clinical trials: Evidence from ClinicalTrials.gov. Contemporary Clinical Trials, 2009, 30, 289-292.	0.8	122
97	The quality of patient-doctor communication about end-of-life care: a study of patients with advanced AIDS and their primary care clinicians. Aids, 1999, 13, 1123-1131.	1.0	119
98	Validation of Preferences for Life-Sustaining Treatment: Implications for Advance Care Planning. Annals of Internal Medicine, 1997, 127, 509.	2.0	118
99	Functional status and perceived quality of life in adults with and without chronic conditions. Journal of Clinical Epidemiology, 2000, 53, 779-785.	2.4	117
100	Patient-Reported Outcome and Observer-Reported Outcome Assessment in Rare Disease Clinical Trials: An ISPOR COA Emerging Good Practices Task Force Report. Value in Health, 2017, 20, 838-855.	0.1	117
101	Disease-specific Versus Generic Measurement of Health-related Quality of Life in Insulin-dependent Diabetic Patients. Medical Care, 1993, 31, 629-639.	1.1	116
102	Association of Depression and Life-Sustaining Treatment Preferences in Patients With COPD. Chest, 2005, 127, 328-334.	0.4	116
103	Validation of the EQ-5D in a general population sample in urban China. Quality of Life Research, 2012, 21, 155-160.	1.5	116
104	The assessment of health status among patients with COPD. European Respiratory Journal, 2003, 21, 36S-45s.	3.1	115
105	A structured review of randomized controlled trials of weight loss showed little improvement in health-related quality of life. Journal of Clinical Epidemiology, 2005, 58, 568-578.	2.4	113
106	Quality of life of persons with onychomycosis. Quality of Life Research, 1993, 2, 341-348.	1.5	112
107	Relative Importance of Sexuality and Quality of Life in Patients with Prostatic Symptoms. European Urology, 1997, 31, 272-280.	0.9	112
108	Care for Depression in HMO Patients Aged 65 and Older. Journal of the American Geriatrics Society, 2000, 48, 871-878.	1.3	112

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109	Restricted activity days among older adults American Journal of Public Health, 1992, 82, 1263-1267.	1.5	111
110	Evaluating the credibility of anchor based estimates of minimal important differences for patient reported outcomes: instrument development and reliability study. BMJ, The, 2020, 369, m1714.	3.0	110
111	Clinician-Reported Outcome Assessments of Treatment Benefit: Report of the ISPOR Clinical Outcome Assessment Emerging Good Practices Task Force. Value in Health, 2017, 20, 2-14.	0.1	109
112	A cross-cultural comparison of health status values American Journal of Public Health, 1985, 75, 1402-1407.	1.5	108
113	Reliability, validity and ability to detect change of the clinician-rated Personal and Social Performance scale in patients with acute symptoms of schizophrenia. Current Medical Research and Opinion, 2009, 25, 325-338.	0.9	107
114	Longitudinal Assessment of Quality of Life in Acute Psychiatric Inpatients: Reliability and Validity. Journal of Nervous and Mental Disease, 1997, 185, 166-175.	0.5	107
115	Economic Evaluation of Aquatic Exercise for Persons With Osteoarthritis. Medical Care, 2001, 39, 413-424.	1.1	106
116	Minimally important difference estimates and methods: a protocol. BMJ Open, 2015, 5, e007953.	0.8	103
117	Managed care, physician job satisfaction, and the quality of primary care. Journal of General Internal Medicine, 2005, 20, 271-277.	1.3	100
118	Universal Health Outcome Measures for Older Persons with Multiple Chronic Conditions. Journal of the American Geriatrics Society, 2012, 60, 2333-2341.	1.3	99
119	Characterization of Health Related Quality of Life (HRQOL) for Patients With Functional Bowel Disorder (FBD) and Its Response to Treatment. American Journal of Gastroenterology, 2007, 102, 1442-1453.	0.2	98
120	A comparison of patient, family, and physician assessments of the value of medical intensive care. Critical Care Medicine, 1988, 16, 594-600.	0.4	97
121	Does Adherence Make a Difference?. Nursing Research, 2002, 51, 285-291.	0.8	97
122	Patient-reported respiratory symptoms in cystic fibrosis. Journal of Cystic Fibrosis, 2009, 8, 245-252.	0.3	96
123	Access to Primary Health Care Among Persons With Disabilities in Rural Areas: A Summary of the Literature. Journal of Rural Health, 1996, 12, 45-53.	1.6	93
124	Quality of life and health-risk behaviors among adolescents. Journal of Adolescent Health, 2001, 29, 426-435.	1,2	93
125	"Molecule Corrals" for Studies of Monolayer Organic Films. Science, 1994, 265, 231-234.	6.0	92
126	Quality of Life of Women with Urinary Incontinence: Cross-cultural Performance of 15 Language Versions of the I-QOL. Quality of Life Research, 2005, 14, 1901-1913.	1.5	92

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127	Stigma experiences in youth with facial differences: a multi-site study of adolescents and their mothers. Orthodontics and Craniofacial Research, 2007, 10, 96-103.	1.2	90
128	Cross-cultural development of a quality of life measure for men with erection difficulties. Quality of Life Research, 1996, 5, 443-449.	1.5	89
129	Disability and Patient Satisfaction With Medical Care. Medical Care, 1983, 21, 1062-1075.	1.1	88
130	Correlates of exercise maintenance among people with mobility impairments. Disability and Rehabilitation, 1999, 21, 15-22.	0.9	88
131	Use of alternative therapies by older adults with osteoarthritis. Arthritis and Rheumatism, 2001, 45, 222-227.	6.7	87
132	ORIGINAL RESEARCHâ€"EJACULATORY DISORDERS: Interrelationships Among Measures of Premature Ejaculation: The Central Role of Perceived Control. Journal of Sexual Medicine, 2007, 4, 780-788.	0.3	85
133	Depressive Symptoms and Mortality in a Prospective Study of 2,558 Older Adults. American Journal of Geriatric Psychiatry, 2002, 10, 521-530.	0.6	84
134	A New Instrument to Assess the Long-term Quality of Life Effects From Migraine: Development and Psychometric Testing of the MSQOL. Headache, 1996, 36, 484-492.	1.8	83
135	Barriers to communication about end-of-life care in AIDS patients. Journal of General Internal Medicine, 1997, 12, 736-741.	1.3	83
136	Assessing the clinical significance of health-related quality of life (HrQOL) improvements in anaemic cancer patients receiving epoetin alfa. European Journal of Cancer, 2003, 39, 335-345.	1.3	83
137	Depression status, medical comorbidity and resource costs. British Journal of Psychiatry, 2003, 183, 121-131.	1.7	83
138	A good death: A qualitative study of patients with advanced AIDS. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2002, 14, 587-598.	0.6	82
139	Longitudinal Analysis of Sexual Function Reported by Men in the Prostate Cancer Prevention Trial. Journal of the National Cancer Institute, 2007, 99, 1025-1035.	3.0	80
140	Patient-reported outcomes in meta-analyses $\hat{a} \in \text{Part 1: assessing risk of bias and combining outcomes.}$ Health and Quality of Life Outcomes, 2013, 11, 109.	1.0	80
141	Pain and health-related quality of life in patients with advanced solid tumours and bone metastases: integrated results from three randomized, double-blind studies of denosumab and zoledronic acid. Supportive Care in Cancer, 2013, 21, 3497-3507.	1.0	80
142	Rapid translation of quality of life measures for international clinical trials: avoiding errors in the minimalist approach. Quality of Life Research, 1994, 3, 403-412.	1.5	79
143	Transforming Self-Rated Health and the SF-36 Scales to Include Death and Improve Interpretability. Medical Care, 2001, 39, 670-680.	1.1	79
144	Assessing health-related quality of life for clinical decision-making. , 1993, , 11-63.		79

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145	Use of the Health Utilities Index With Stroke Patients and Their Caregivers. Stroke, 1997, 28, 1888-1894.	1.0	79
146	Necessary but not sufficient. Journal of General Internal Medicine, 1994, 9, 606-615.	1.3	77
147	Patterns of Self-Rated Health in Older Adults Before and After Sentinel Health Events. Journal of the American Geriatrics Society, 2001, 49, 36-44.	1.3	76
148	Mode of Communication, Perceived Level of Understanding, and Perceived Quality of Life in Youth Who Are Deaf or Hard of Hearing. Journal of Deaf Studies and Deaf Education, 2011, 16, 512-523.	0.7	76
149	Assessing health-related quality-of-life and health state preference in persons with obesity: a validation study. Quality of Life Research, 1997, 6, 311-322.	1.5	75
150	Managed care, access to mental health specialists, and outcomes among primary care patients with depressive symptoms. Journal of General Internal Medicine, 2002, 17, 258-269.	1.3	75
151	Performance of Two Selfâ€Report Measures for Evaluating Obesity and Weight Loss*. Obesity, 2004, 12, 48-57.	4.0	75
152	Assessing health and quality of life outcomes in dialysis: A report on an institute of medicine workshop. American Journal of Kidney Diseases, 1997, 30, 140-155.	2.1	74
153	Insights pertaining to patient assessments of states worse than death. Journal of Clinical Ethics, 1993, 4, 33-41.	0.1	74
154	The Maine Lumbar Spine Study, Part I. Spine, 1996, 21, 1769-1776.	1.0	73
155	Quality of Life: How Do Adolescents with Facial Differences Compare with Other Adolescents?. Cleft Palate-Craniofacial Journal, 2005, 42, 25-32.	0.5	73
156	Development and Validation of the Diabetes Quality of Life Clinical Trial Questionnaire. Medical Care, 1999, 37, AS45-AS66.	1.1	73
157	Measuring the Quality of Life of Youth with Facial Differences. Cleft Palate-Craniofacial Journal, 2007, 44, 538-547.	0.5	72
158	Evaluation of the Irritable Bowel Syndrome Quality of Life (IBS-QOL) questionnaire in diarrheal-predominant irritable bowel syndrome patients. Health and Quality of Life Outcomes, 2013, 11, 208.	1.0	72
159	Return of incidental findings in genomic medicine: measuring what patients valueâ€"development of an instrument to measure preferences for information from next-generation testing (IMPRINT). Genetics in Medicine, 2013, 15, 873-881.	1.1	72
160	Cultural Adaptation of a Quality-of-Life Measure for Urinary Incontinence. European Urology, 1999, 36, 427-435.	0.9	70
161	Quality of Life of Adolescents With Perceived Disabilities. Journal of Pediatric Psychology, 2003, 28, 233-241.	1.1	69
162	Effects of Self-Perceived Hair Loss in a Community Sample of Men. Dermatology, 1998, 197, 223-229.	0.9	67

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163	Measuring health and health state preferences among critically ill patients. Intensive Care Medicine, 1996, 22, 1379-1384.	3.9	66
164	A Review of Selected Patient-Generated Outcome Measures and Their Application in Clinical Trials. Value in Health, 2003, 6, 595-603.	0.1	66
165	Validation of Electronic Data Capture of the Irritable Bowel Syndromeâ€"Quality of Life Measure, the Work Productivity and Activity Impairment Questionnaire for Irritable Bowel Syndrome and the EuroQol. Value in Health, 2006, 9, 98-105.	0.1	65
166	SPIRIT-PRO Extension explanation and elaboration: guidelines for inclusion of patient-reported outcomes in protocols of clinical trials. BMJ Open, 2021, 11, e045105.	0.8	65
167	Evaluating CAM treatment at the end of life: A review of clinical trials for massage and meditation. Complementary Therapies in Medicine, 2006, 14, 100-112.	1.3	64
168	The international development of the RGHQoL: a quality of life measure for recurrent genital herpes. Quality of Life Research, 1998, 7, 143-153.	1.5	61
169	Pain and analgesic use associated with skeletal-related events in patients with advanced cancer and bone metastases. Supportive Care in Cancer, 2016, 24, 1327-1337.	1.0	61
170	Measurement of Health-Related Quality of Life in Patients With Amyotrophic Lateral Sclerosis in Clinical Trials of New Therapies. Medical Care, 1999, 37, 15-26.	1.1	61
171	Integrating Mental Health Services into Primary Medical Care. Medical Care, 1976, 14, 654-661.	1.1	60
172	Environmental Indicators: A Tool for Evaluating Community-Based Health-Promotion Programs. American Journal of Preventive Medicine, 1992, 8, 345-350.	1.6	60
173	Validation of a symptoms questionnaire for benign prostatic hyperplasia. Prostate, 1992, 21, 247-254.	1.2	60
174	Psychosocial support and change in the health status of physically disabled people. Social Science and Medicine, 1986, 22, 1347-1354.	1.8	59
175	Rethinking Prevention for People with Disabilities Part I: A Conceptual Model for Promoting Health. American Journal of Health Promotion, 1997, 11, 257-260.	0.9	59
176	A new approach to multicultural item generation in the development of two obesity-specific measures: The obesity and weight loss quality of life (OWLQOL) questionnaire and the weight-related symptom measure (WRSM). Clinical Therapeutics, 2002, 24, 690-700.	1.1	59
177	Use of Quality-Adjusted Life Year Weights with Dose-Response Models for Public Health Decisions: A Case Study of the Risks and Benefits of Fish Consumption. Risk Analysis, 2000, 20, 529-542.	1.5	58
178	Profile of Men Randomized to the Prostate Cancer Prevention Trial: Baseline Health-Related Quality of Life, Urinary and Sexual Functioning, and Health Behaviors. Journal of Clinical Oncology, 2000, 18, 1942-1953.	0.8	58
179	Managed Care and Primary Physician Satisfaction. Journal of the American Board of Family Medicine, 2003, 16, 383-393.	0.8	58
180	Dissatisfaction with medical services among medicare beneficiaries with disabilities. Archives of Physical Medicine and Rehabilitation, 2002, 83, 1335-1341.	0.5	57

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181	Quality of life of adolescent males with Attention-Deficit Hyperactivity Disorder. Journal of Attention Disorders, 2004, 7, 163-173.	1.5	56
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