Elizabeth A Hahn

List of Publications by Year in descending order

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96 papers 8,492 citations

126858 33 h-index 49868 87 g-index

98 all docs 98 docs citations

98 times ranked 10730 citing authors

#	Article	IF	CITATIONS
1	The Patient-Reported Outcomes Measurement Information System (PROMIS) developed and tested its first wave of adult self-reported health outcome item banks: 2005–2008. Journal of Clinical Epidemiology, 2010, 63, 1179-1194.	2.4	3,521
2	Reliability and validity of the Functional Assessment of Cancer Therapy-Colorectal (FACT-C) quality of life instrument. Quality of Life Research, 1999, 8, 181-195.	1.5	374
3	Sexual Satisfaction and the Importance of Sexual Health to Quality of Life Throughout the Life Course of U.S. Adults. Journal of Sexual Medicine, 2016, 13, 1642-1650.	0.3	316
4	Long-Term Health-Related Quality of Life, Growth, and Spiritual Well-Being After Hematopoietic Stem-Cell Transplantation. Journal of Clinical Oncology, 2005, 23, 599-608.	0.8	284
5	Measuring social health in the patient-reported outcomes measurement information system (PROMIS): item bank development and testing. Quality of Life Research, 2010, 19, 1035-1044.	1.5	279
6	New English and Spanish social health measures will facilitate evaluating health determinants Health Psychology, 2014, 33, 490-499.	1.3	240
7	Environmental Barriers and Supports to Everyday Participation: A Qualitative Insider Perspective From People With Disabilities. Archives of Physical Medicine and Rehabilitation, 2015, 96, 578-588.	0.5	227
8	Quality of Life in Patients With Newly Diagnosed Chronic Phase Chronic Myeloid Leukemia on Imatinib Versus Interferon Alfa Plus Low-Dose Cytarabine: Results From the IRIS Study. Journal of Clinical Oncology, 2003, 21, 2138-2146.	0.8	191
9	Patient-reported outcomes measurement information system (PROMIS) domain names and definitions revisions: further evaluation of content validity in IRT-derived item banks. Quality of Life Research, 2010, 19, 1311-1321.	1.5	165
10	Validation of the PROMIS physical function measures in a diverse US population-based cohort of cancer patients. Quality of Life Research, 2015, 24, 2333-2344.	1.5	162
11	Effect of a Home-Based Exercise Intervention of Wearable Technology and Telephone Coaching on Walking Performance in Peripheral Artery Disease. JAMA - Journal of the American Medical Association, 2018, 319, 1665.	3 . 8	151
12	Precision of Health-Related Quality-of-Life Data Compared With Other Clinical Measures. Mayo Clinic Proceedings, 2007, 82, 1244-1254.	1.4	131
13	Development and Initial Validation of the PROMIS® Sexual Function and Satisfaction Measures Version 2.0. Journal of Sexual Medicine, 2015, 12, 1961-1974.	0.3	130
14	Health outcomes assessment in vulnerable populations: Measurement challenges and recommendations. Archives of Physical Medicine and Rehabilitation, 2003, 84, S35-S42.	0.5	120
15	Assessment is not enough: a randomized controlled trial of the effects of HRQL assessment on quality of life and satisfaction in oncology clinical practice. Psycho-Oncology, 2007, 16, 1069-1079.	1.0	118
16	The PROMIS satisfaction with social participation measures demonstrated responsiveness in diverse clinical populations. Journal of Clinical Epidemiology, 2016, 73, 135-141.	2.4	97
17	Statistical analysis of quality of life with missing data in cancer clinical trials. , 1998, 17, 653-666.		87
18	The talking touchscreen: A new approach to outcomes assessment in low literacy. Psycho-Oncology, 2004, 13, 86-95.	1.0	82

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19	Content validity in the PROMIS social-health domain: a qualitative analysis of focus-group data. Quality of Life Research, 2008, 17, 737-749.	1.5	76
20	Using Multiple Anchor- and Distribution-Based Estimates to Evaluate Clinically Meaningful Change on the Functional Assessment of Cancer Therapy-Biologic Response Modifiers (FACT-BRM) Instrument. Value in Health, 2005, 8, 117-127.	0.1	75
21	Responsiveness of 8 Patientâ€Reported Outcomes Measurement Information System (PROMIS) measures in a large, communityâ€based cancer study cohort. Cancer, 2017, 123, 327-335.	2.0	69
22	Environmental Factors Item Development for Persons WithÂStroke, Traumatic Brain Injury, and Spinal Cord Injury. Archives of Physical Medicine and Rehabilitation, 2015, 96, 589-595.	0.5	65
23	Health-Related Quality of Life in Caregivers of Individuals With Traumatic Brain Injury: Development of a Conceptual Model. Archives of Physical Medicine and Rehabilitation, 2015, 96, 105-113.	0.5	63
24	Pragmatic randomised clinical trial of proton versus photon therapy for patients with non-metastatic breast cancer: the Radiotherapy Comparative Effectiveness (RadComp) Consortium trial protocol. BMJ Open, 2019, 9, e025556.	0.8	60
25	Health Literacy Assessment Using Talking Touchscreen Technology (Health LiTT): A New Item Response Theory-Based Measure of Health Literacy. Journal of Health Communication, 2011, 16, 150-162.	1.2	58
26	Predictors of health-related quality of life in patients with colorectal cancer. Health and Quality of Life Outcomes, 2008, 6, 66.	1.0	57
27	Development and validation of the functional assessment of chronic illness therapy treatment satisfaction (FACIT TS) measures. Quality of Life Research, 2014, 23, 815-824.	1.5	53
28	Reliability and construct validity of PROMIS® measures for patients with heart failure who undergo heart transplant. Quality of Life Research, 2015, 24, 2591-2599.	1.5	51
29	Caring for our wounded warriors: A qualitative examination of health-related quality of life in caregivers of individuals with military-related traumatic brain injury. Journal of Rehabilitation Research and Development, 2016, 53, 669-680.	1.6	51
30	The TBI-CareQOL Measurement System: Development and Preliminary Validation of Health-Related Quality of Life Measures for Caregivers of Civilians and Service Members/Veterans With Traumatic Brain Injury. Archives of Physical Medicine and Rehabilitation, 2019, 100, S1-S12.	0.5	49
31	Measuring Participation: The Patient-Reported Outcomes Measurement Information System Experience. Archives of Physical Medicine and Rehabilitation, 2010, 91, S60-S65.	0.5	42
32	Bilingual health literacy assessment using the Talking Touchscreen/la Pantalla Parlanchina: Development and pilot testing. Patient Education and Counseling, 2009, 75, 295-301.	1.0	41
33	Measuring Social Well-Being in People with Chronic Illness. Social Indicators Research, 2010, 96, 381-401.	1.4	38
34	Calibration and initial validation of a general self-efficacy item bank and short form for the NIH PROMIS®. Quality of Life Research, 2019, 28, 2513-2523.	1.5	38
35	The impact of literacy on health-related quality of life measurement and outcomes in cancer outpatients. Quality of Life Research, 2007, 16, 495-507.	1.5	37
36	Health Literacy and Patient-Reported Outcomes: A Cross-Sectional Study of Underserved English- and Spanish-Speaking Patients With Type 2 Diabetes. Journal of Health Communication, 2015, 20, 4-15.	1.2	37

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37	Acceptability of the Talking Touchscreen for Health Literacy Assessment. Journal of Health Communication, 2010, 15, 80-92.	1.2	35
38	Addressing low literacy and health literacy in clinical oncology practice. The Journal of Supportive Oncology, 2010, 8, 64-9.	2.3	31
39	Item Banks and Their Potential Applications to Health Status Assessment in Diverse Populations. Medical Care, 2006, 44, S189-S197.	1.1	30
40	Health-related quality of life in mechanical circulatory support: Development of a new conceptual model and items for self-administration. Journal of Heart and Lung Transplantation, 2015, 34, 1292-1304.	0.3	29
41	Comparability of Interview- and Self-Administration of the Functional Assessment of Cancer Therapy-General (FACT-G) in English- and Spanish-Speaking Ambulatory Cancer Patients. Medical Care, 2008, 46, 423-431.	1.1	28
42	Understanding Health-Related Quality of Life of Caregivers of Civilians and Service Members/Veterans With Traumatic Brain Injury: Establishing the Reliability and Validity of PROMIS Social Health Measures. Archives of Physical Medicine and Rehabilitation, 2019, 100, S110-S118.	0.5	28
43	Assessing meaning & Durpose in life: development and validation of an item bank and short forms for the NIH PROMIS®. Quality of Life Research, 2020, 29, 2299-2310.	1.5	27
44	Missing data in quality of life research in Eastern Cooperative Oncology Group (ECOG) clinical trials: problems and solutions., 1998, 17, 547-559.		24
45	Implementation of the PROMIS-29 in Routine Care for People With Diabetes. Journal of Ambulatory Care Management, 2018, 41, 274-287.	0.5	24
46	The Development of a New Computer-Adaptive Test to Evaluate Strain in Caregivers of Individuals With TBI: TBI-CareQOL Caregiver Strain. Archives of Physical Medicine and Rehabilitation, 2019, 100, S13-S21.	0.5	24
47	Cross-Cultural Evaluation of Health Status Using Item Response Theory. Evaluation and the Health Professions, 2005, 28, 233-259.	0.9	23
48	The association between health literacy and indicators of cognitive impairment in a diverse sample of primary care patients. Patient Education and Counseling, 2013, 93, 319-326.	1.0	21
49	Development of Self-Report Measures of Social Attitudes That Act As Environmental Barriers and Facilitators for People With Disabilities. Archives of Physical Medicine and Rehabilitation, 2015, 96, 596-603.	0.5	21
50	Literacy-Fair Measurement of Health-Related Quality of Life Will Facilitate Comparative Effectiveness Research in Spanish-Speaking Cancer Outpatients. Medical Care, 2010, 48, S75-S82.	1.1	20
51	Development and psychometric characteristics of the SCI-QOL Ability to Participate and Satisfaction with Social Roles and Activities item banks and short forms. Journal of Spinal Cord Medicine, 2015, 38, 397-408.	0.7	20
52	Using the ICF's environmental factors framework to develop an item bank measuring built and natural environmental features affecting persons with disabilities. Quality of Life Research, 2016, 25, 2775-2786.	1.5	18
53	Health-related quality of life among women with ductal carcinoma <i>in situ</i> or early invasive breast cancer: validation of the FACT-B (version 4). Expert Review of Quality of Life in Cancer Care, 2016, 1, 99-109.	0.6	17
54	The Development of a New Computer Adaptive Test to Evaluate Feelings of Being Trapped in Caregivers of Individuals With Traumatic Brain Injury: TBI-CareQOL Feeling Trapped Item Bank. Archives of Physical Medicine and Rehabilitation, 2019, 100, S43-S51.	0.5	17

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55	Quality of life assessment for low literacy Latinos: a new multimedia program for self-administration. The Journal of Oncology Management: the Official Journal of the American College of Oncology Administrators, 2003, 12, 9-12.	0.0	17
56	Conducting routine fatigue assessments for use in clinical oncology practice: patient and provider perspectives. Supportive Care in Cancer, 2008, 16, 379-386.	1.0	16
57	Factors Associated With End-of-Life Planning in Huntington Disease. American Journal of Hospice and Palliative Medicine, 2018, 35, 440-447.	0.8	16
58	High-Risk Human Papillomavirus Detection. American Journal of Clinical Pathology, 2004, 121, 537-545.	0.4	14
59	The Development of Two New Computer Adaptive Tests To Evaluate Feelings of Loss in Caregivers of Individuals With Traumatic Brain Injury: TBI-CareQOL Feelings of Loss-Self and Feelings of Loss-Person With Traumatic Brain Injury. Archives of Physical Medicine and Rehabilitation, 2019, 100, S31-S42.	0.5	14
60	Health and Functional Literacy in Physical Rehabilitation Patients. Health Literacy Research and Practice, 2017, 1, e71-e85.	0.5	13
61	Evaluating linguistic equivalence of patient-reported outcomes in a cancer clinical trial. Clinical Trials, 2006, 3, 280-290.	0.7	12
62	Patient attitudes and preferences regarding literacy screening in ambulatory cancer care clinics. Patient Related Outcome Measures, 2010, 1, 19.	0.7	12
63	The challenge of measuring intra-individual change in fatigue during cancer treatment. Quality of Life Research, 2017, 26, 259-271.	1.5	11
64	Health literacy, health outcomes, and the caregiver role in traumatic brain injury Rehabilitation Psychology, 2020, 65, 401-408.	0.7	11
65	English-Spanish Equivalence of the Health Literacy Assessment Using Talking Touchscreen Technology (Health LiTT). Journal of Health Communication, 2014, 19, 285-301.	1.2	10
66	Validation of the Functional Assessment of Cancer Therapy-Breast (FACT-B) quality of life instrument Journal of Clinical Oncology, 2015, 33, e17753-e17753.	0.8	10
67	Quality of life on imatinib. Seminars in Hematology, 2003, 40, 31-36.	1.8	9
68	Patientâ€reported outcomes in Huntington's disease: Quality of life in neurological disorders (Neuroâ€QoL) and Huntington's disease healthâ€related quality of life (HDQLIFE) physical function measures. Movement Disorders, 2017, 32, 1096-1102.	2.2	9
69	End-of-life measures in Huntington disease: HDQLIFE Meaning and Purpose, Concern with Death and Dying, and EndÂofÂLife Planning. Journal of Neurology, 2019, 266, 2406-2422.	1.8	9
70	Investigation of a multimedia, computerâ€based approach to improve knowledge, attitudes, selfâ€efficacy, and receptivity to cancer clinical trials among newly diagnosed patients with diverse health literacy skills. Cancer, 2019, 125, 2066-2075.	2.0	9
71	Patient-reported outcome measures in Huntington disease: Quality of life in neurological disorders (Neuro-QoL) social functioning measures Psychological Assessment, 2018, 30, 450-458.	1.2	9
72	Refining and supplementing candidate measures of psychological well-being for the NIH PROMIS®: qualitative results from a mixed cancer sample. Quality of Life Research, 2018, 27, 2471-2476.	1.5	8

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73	Sociocultural Factors Influencing Caregiver Appraisals Following Traumatic Brain Injury. Archives of Physical Medicine and Rehabilitation, 2019, 100, S58-S64.	0.5	8
74	Variation and Change Over Time in PROMIS-29 Survey Results Among Primary Care Patients With Type 2 Diabetes. Journal of Patient-centered Research and Reviews, 2019, 6, 135-147.	0.6	8
75	Measuring social function in diverse cancer populations: Evaluation of measurement equivalence of the Patient Reported Outcomes Measurement Information System (PROMIS) Ability to Participate in Social Roles and Activities short form. Psychological Test and Assessment Modeling, 2016, 58, 403-421.	0.6	8
76	Addressing Lupus Health Disparities: The MONARCAS Community and Academic Collaborative Program. Health Equity, 2018, 2, 145-151.	0.8	7
77	Measuring Access to Information and Technology: Environmental Factors Affecting Persons With Neurologic Disorders. Archives of Physical Medicine and Rehabilitation, 2016, 97, 1284-1294.	0.5	5
78	Factors affecting enrollment in literacy studies for English―and Spanishâ€speaking cancer patients. Statistics in Medicine, 2008, 27, 4119-4131.	0.8	4
79	The impact of missing data on estimation of health-related quality of life outcomes: an analysis of a randomized longitudinal clinical trial. Health Services and Outcomes Research Methodology, 2011, 11, 134-144.	0.8	4
80	Development of a new multimedia instrument to measure cancer-specific quality of life in Portuguese-speaking patients with varying literacy skills. SpringerPlus, 2016, 5, 972.	1,2	4
81	Understanding domains that influence perceived stigma in individuals with Huntington disease Rehabilitation Psychology, 2020, 65, 113-121.	0.7	4
82	Measure validation is an ongoing process: the Functional Assessment of Cancer Therapy-Breast Symptom Index as a case example. Annals of Palliative Medicine, 2012, 1, 207-10.	0.5	4
83	Patient preferences regarding medical decision making in the emergency care setting: a pilot-study. American Journal of Emergency Medicine, 2015, 33, 719-721.	0.7	3
84	Rasch Analysis of Social Attitude Barriers and Facilitators to Participation for Individuals with Disabilities. Archives of Physical Medicine and Rehabilitation, 2021, 102, 675-686.	0.5	3
85	Validation of a Short Form for Health Literacy Assessment Using Talking Touchscreen Technology. Health Literacy Research and Practice, 2020, 4, e200-e207.	0.5	3
86	Health care barriers and quality of life in central centrifugal cicatricial alopecia patients. Cutis, 2018, 102, 427-432.	0.4	3
87	Survey response burden in caregivers of civilians and service members/veterans with traumatic brain injury. Patient Related Outcome Measures, 2019, Volume 10, 59-66.	0.7	2
88	A Low Literacy, Multimedia Health Information Technology Intervention to Enhance Patient-Centered Cancer Care in Safety Net Settings Increased Cancer Knowledge in a Randomized Controlled Trial. Cancer Control, 2021, 28, 107327482110367.	0.7	2
89	Missing data in quality of life research in Eastern Cooperative Oncology Group (ECOG) clinical trials: problems and solutions. Statistics in Medicine, 1998, 17, 547-559.	0.8	2
90	Understanding the need for assistance with survey completion in people with Huntington disease. Quality of Life Research, 2018, 27, 801-810.	1.5	1

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91	Responsiveness to Change Over Time: An Examination of the Neuro-QoL Social Function Measures in Persons with Huntington's Disease. Journal of Huntington's Disease, 2020, 9, 83-97.	0.9	1
92	Quality of life on imatinib. Seminars in Hematology, 2003, 40, 31-36.	1.8	1
93	In response to "Scope of shared decision making in patients with psychologic complaints― American Journal of Emergency Medicine, 2015, 33, 841.	0.7	O
94	Piloting breast cancer survivorship care planning in a safety net hospital Journal of Clinical Oncology, 2017, 35, 65-65.	0.8	0
95	Resource and reimbursement barriers to comprehensive cancer care (CCC) delivery: An association of community cancer centers (ACCC) survey research analysis Journal of Clinical Oncology, 2020, 38, 31-31.	0.8	0

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