

Elizabeth A Hahn

List of Publications by Year in descending order

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96
papers

8,492
citations

145106

33
h-index

56606

87
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98
all docs

98
docs citations

98
times ranked

11367
citing authors

#	ARTICLE	IF	CITATIONS
1	Research engagement and experiences of patients pre- and post-implant of a left ventricular assist device from the mechanical circulatory support measures of adjustment and quality of life (MCS) Tj ETQq1 1 0.7843.54 rgBT (Overlock		
2	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
3	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
4	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
5	Assessing meaning & purpose 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
6	Understanding domains that influence perceived stigma in individuals with Huntington disease.. Rehabilitation Psychology, 2020, 65, 113-121.	0.7	4
7	Health literacy, health outcomes, and the caregiver role in traumatic brain injury.. Rehabilitation Psychology, 2020, 65, 401-408.	0.7	11
8	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
9	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
10	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
11	Sociocultural Factors Influencing Caregiver Appraisals Following Traumatic Brain Injury. Archives of Physical Medicine and Rehabilitation, 2019, 100, S58-S64.	0.5	8
12	<p>Survey response burden in caregivers of civilians and service members/veterans with traumatic brain injury</p>. Patient Related Outcome Measures, 2019, Volume 10, 59-66.	0.7	2
13	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
14	End-of-life measures in Huntington disease: HDQLIFE Meaning and Purpose, Concern with Death and Dying, and End-of-Life Planning. Journal of Neurology, 2019, 266, 2406-2422.	1.8	9
15	Investigation of a multimedia, computer-based approach to improve knowledge, attitudes, self-efficacy, and receptivity to cancer clinical trials among newly diagnosed patients with diverse health literacy skills. Cancer, 2019, 125, 2066-2075.	2.0	9
16	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
17	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
18	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

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19	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
20	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
21	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
22	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
23	Factors Associated With End-of-Life Planning in Huntington Disease. American Journal of Hospice and Palliative Medicine, 2018, 35, 440-447.	0.8	16
24	Understanding the need for assistance with survey completion in people with Huntington disease. Quality of Life Research, 2018, 27, 801-810.	1.5	1
25	Implementation of the PROMIS-29 in Routine Care for People With Diabetes. Journal of Ambulatory Care Management, 2018, 41, 274-287.	0.5	24
26	Addressing Lupus Health Disparities: The MONARCAS Community and Academic Collaborative Program. Health Equity, 2018, 2, 145-151.	0.8	7
27	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
28	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
29	Health care barriers and quality of life in central centrifugal cicatricial alopecia patients. Cutis, 2018, 102, 427-432.	0.4	3
30	Patient-reported outcomes in Huntington's disease: Quality of life in neurological disorders (Neuro-QoL) and Huntington's disease health-related quality of life (HDQLIFE) physical function measures. Movement Disorders, 2017, 32, 1096-1102.	2.2	9
31	Responsiveness of 8 Patient-Reported Outcomes Measurement Information System (PROMIS) measures in a large, community-based cancer study cohort. Cancer, 2017, 123, 327-335.	2.0	69
32	The challenge of measuring intra-individual change in fatigue during cancer treatment. Quality of Life Research, 2017, 26, 259-271.	1.5	11
33	Health and Functional Literacy in Physical Rehabilitation Patients. Health Literacy Research and Practice, 2017, 1, e71-e85.	0.5	13
34	Piloting breast cancer survivorship care planning in a safety net hospital.. Journal of Clinical Oncology, 2017, 35, 65-65.	0.8	0
35	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
36	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

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37	Using the ICF's environmental factors framework to develop an item bank measuring built and natural environmental features affecting persons with disabilities. <i>Quality of Life Research</i> , 2016, 25, 2775-2786.	1.5	18
38	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). <i>Expert Review of Quality of Life in Cancer Care</i> , 2016, 1, 99-109.	0.6	17
39	The PROMIS satisfaction with social participation measures demonstrated responsiveness in diverse clinical populations. <i>Journal of Clinical Epidemiology</i> , 2016, 73, 135-141.	2.4	97
40	Measuring Access to Information and Technology: Environmental Factors Affecting Persons With Neurologic Disorders. <i>Archives of Physical Medicine and Rehabilitation</i> , 2016, 97, 1284-1294.	0.5	5
41	Caring for our wounded warriors: A qualitative examination of health-related quality of life in caregivers of individuals with military-related traumatic brain injury. <i>Journal of Rehabilitation Research and Development</i> , 2016, 53, 669-680.	1.6	51
42	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. <i>Psychological Test and Assessment Modeling</i> , 2016, 58, 403-421.	0.6	8
43	Health-related quality of life in mechanical circulatory support: Development of a new conceptual model and items for self-administration. <i>Journal of Heart and Lung Transplantation</i> , 2015, 34, 1292-1304.	0.3	29
44	Development and Initial Validation of the PROMIS® Sexual Function and Satisfaction Measures Version 2.0. <i>Journal of Sexual Medicine</i> , 2015, 12, 1961-1974.	0.3	130
45	Health Literacy and Patient-Reported Outcomes: A Cross-Sectional Study of Underserved English- and Spanish-Speaking Patients With Type 2 Diabetes. <i>Journal of Health Communication</i> , 2015, 20, 4-15.	1.2	37
46	Development and psychometric characteristics of the SCI-QOL Ability to Participate and Satisfaction with Social Roles and Activities item banks and short forms. <i>Journal of Spinal Cord Medicine</i> , 2015, 38, 397-408.	0.7	20
47	Patient preferences regarding medical decision making in the emergency care setting: a pilot-study. <i>American Journal of Emergency Medicine</i> , 2015, 33, 719-721.	0.7	3
48	In response to "Scope of shared decision making in patients with psychologic complaints" <i>American Journal of Emergency Medicine</i> , 2015, 33, 841.	0.7	0
49	Environmental Barriers and Supports to Everyday Participation: A Qualitative Insider Perspective From People With Disabilities. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015, 96, 578-588.	0.5	227
50	Validation of the PROMIS physical function measures in a diverse US population-based cohort of cancer patients. <i>Quality of Life Research</i> , 2015, 24, 2333-2344.	1.5	162
51	Reliability and construct validity of PROMIS® measures for patients with heart failure who undergo heart transplant. <i>Quality of Life Research</i> , 2015, 24, 2591-2599.	1.5	51
52	Health-Related Quality of Life in Caregivers of Individuals With Traumatic Brain Injury: Development of a Conceptual Model. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015, 96, 105-113.	0.5	63
53	Environmental Factors Item Development for Persons With Stroke, Traumatic Brain Injury, and Spinal Cord Injury. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015, 96, 589-595.	0.5	65
54	Development of Self-Report Measures of Social Attitudes That Act As Environmental Barriers and Facilitators for People With Disabilities. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015, 96, 596-603.	0.5	21

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55	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
56	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
57	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
58	New English and Spanish social health measures will facilitate evaluating health determinants.. Health Psychology, 2014, 33, 490-499.	1.3	240
59	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
60	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
61	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
62	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
63	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
64	Measuring Social Well-Being in People with Chronic Illness. Social Indicators Research, 2010, 96, 381-401.	1.4	38
65	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
66	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
67	Patient attitudes and preferences regarding literacy screening in ambulatory cancer care clinics. Patient Related Outcome Measures, 2010, 1, 19.	0.7	12
68	Acceptability of the Talking Touchscreen for Health Literacy Assessment. Journal of Health Communication, 2010, 15, 80-92.	1.2	35
69	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
70	Measuring Participation: The Patient-Reported Outcomes Measurement Information System Experience. Archives of Physical Medicine and Rehabilitation, 2010, 91, S60-S65.	0.5	42
71	Addressing low literacy and health literacy in clinical oncology practice. The Journal of Supportive Oncology, 2010, 8, 64-9.	2.3	31
72	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

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73	Content validity in the PROMIS social-health domain: a qualitative analysis of focus-group data. <i>Quality of Life Research</i> , 2008, 17, 737-749.	1.5	76
74	Conducting routine fatigue assessments for use in clinical oncology practice: patient and provider perspectives. <i>Supportive Care in Cancer</i> , 2008, 16, 379-386.	1.0	16
75	Factors affecting enrollment in literacy studies for English- and Spanish-speaking cancer patients. <i>Statistics in Medicine</i> , 2008, 27, 4119-4131.	0.8	4
76	Predictors of health-related quality of life in patients with colorectal cancer. <i>Health and Quality of Life Outcomes</i> , 2008, 6, 66.	1.0	57
77	Comparability of Interview- and Self-Administration of the Functional Assessment of Cancer Therapy-General (FACT-G) in English- and Spanish-Speaking Ambulatory Cancer Patients. <i>Medical Care</i> , 2008, 46, 423-431.	1.1	28
78	Precision of Health-Related Quality-of-Life Data Compared With Other Clinical Measures. <i>Mayo Clinic Proceedings</i> , 2007, 82, 1244-1254.	1.4	131
79	Assessment is not enough: a randomized controlled trial of the effects of HRQL assessment on quality of life and satisfaction in oncology clinical practice. <i>Psycho-Oncology</i> , 2007, 16, 1069-1079.	1.0	118
80	The impact of literacy on health-related quality of life measurement and outcomes in cancer outpatients. <i>Quality of Life Research</i> , 2007, 16, 495-507.	1.5	37
81	Item Banks and Their Potential Applications to Health Status Assessment in Diverse Populations. <i>Medical Care</i> , 2006, 44, S189-S197.	1.1	30
82	Evaluating linguistic equivalence of patient-reported outcomes in a cancer clinical trial. <i>Clinical Trials</i> , 2006, 3, 280-290.	0.7	12
83	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. <i>Value in Health</i> , 2005, 8, 117-127.	0.1	75
84	Cross-Cultural Evaluation of Health Status Using Item Response Theory. <i>Evaluation and the Health Professions</i> , 2005, 28, 233-259.	0.9	23
85	Long-Term Health-Related Quality of Life, Growth, and Spiritual Well-Being After Hematopoietic Stem-Cell Transplantation. <i>Journal of Clinical Oncology</i> , 2005, 23, 599-608.	0.8	284
86	High-Risk Human Papillomavirus Detection. <i>American Journal of Clinical Pathology</i> , 2004, 121, 537-545.	0.4	14
87	The talking touchscreen: A new approach to outcomes assessment in low literacy. <i>Psycho-Oncology</i> , 2004, 13, 86-95.	1.0	82
88	Quality of life on imatinib. <i>Seminars in Hematology</i> , 2003, 40, 31-36.	1.8	9
89	Health outcomes assessment in vulnerable populations: Measurement challenges and recommendations. <i>Archives of Physical Medicine and Rehabilitation</i> , 2003, 84, S35-S42.	0.5	120
90	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. <i>Journal of Clinical Oncology</i> , 2003, 21, 2138-2146.	0.8	191

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91	Quality of life on imatinib. <i>Seminars in Hematology</i> , 2003, 40, 31-36.	1.8	1
92	Quality of life assessment for low literacy Latinos: a new multimedia program for self-administration. <i>The Journal of Oncology Management: the Official Journal of the American College of Oncology Administrators</i> , 2003, 12, 9-12.	0.0	17
93	Reliability and validity of the Functional Assessment of Cancer Therapy-Colorectal (FACT-C) quality of life instrument. <i>Quality of Life Research</i> , 1999, 8, 181-195.	1.5	374
94	Missing data in quality of life research in Eastern Cooperative Oncology Group (ECOG) clinical trials: problems and solutions. , 1998, 17, 547-559.		24
95	Statistical analysis of quality of life with missing data in cancer clinical trials. , 1998, 17, 653-666.		87
96	Missing data in quality of life research in Eastern Cooperative Oncology Group (ECOG) clinical trials: problems and solutions. <i>Statistics in Medicine</i> , 1998, 17, 547-559.	0.8	2