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List of Publications by Year in descending order

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218592 118793 5,394 63 26 62 h-index citations g-index papers 65 65 65 7193 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Racial differences in user experiences and perceived value of electronic symptom monitoring in a cohort of black and white bladder and prostate cancer patients. Quality of Life Research, 2021, 30, 3213-3227.	1.5	12
2	Composite grading algorithm for the National Cancer Institute's Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE). Clinical Trials, 2021, 18, 104-114.	0.7	80
3	Feasibility and delivery of patient-reported outcomes in clinical practice among racially diverse bladder and prostate cancer patients. Urologic Oncology: Seminars and Original Investigations, 2021, 39, 77.e1-77.e8.	0.8	6
4	The usefulness of the Electronic Patient Visit Assessment (ePVA) \hat{A} \otimes as a clinical support tool for real-time interventions in head and neck cancer. MHealth, 2021, 7, 7-7.	0.9	4
5	Comparison of weekly and daily recall of pain as an endpoint in a randomized phase 3 trial of cabozantinib for metastatic castration-resistant prostate cancer. Clinical Trials, 2021, 18, 408-416.	0.7	4
6	Mapping the Memorial Anxiety Scale for Prostate Cancer to the SF-6D. Quality of Life Research, 2021, 30, 2919-2928.	1.5	3
7	Association of step count with PROMIS pediatric health-related quality of life measures in children and adolescents with persistent asthma. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2492-2494.	2.0	2
8	Facility-level characteristics associated with family planning and child immunization services integration in urban areas of Nigeria: a longitudinal analysis. BMC Public Health, 2021, 21, 1379.	1.2	3
9	Measuring Goal-Concordant Care in Palliative Care Research. Journal of Pain and Symptom Management, 2021, 62, e305-e314.	0.6	17
10	Pilot randomized trial of an electronic symptom monitoring and reporting intervention for hospitalized adults undergoing hematopoietic stem cell transplantation. Supportive Care in Cancer, 2020, 28, 1223-1231.	1.0	20
11	Assessment of Adverse Events From the Patient Perspective in a Phase 3 Metastatic Castration-Resistant Prostate Cancer Clinical Trial. JAMA Oncology, 2020, 6, e193332.	3.4	39
12	Elements of Palliative Care in the Last 6ÂMonths of Life: Frequency, Predictors, and Timing. Journal of General Internal Medicine, 2020, 35, 753-761.	1.3	27
13	Does Receipt of Recommended Elements of Palliative Care Precede In-Hospital Death or Hospice Referral?. Journal of Pain and Symptom Management, 2020, 59, 778-786.	0.6	4
14	Applying patient-reported outcome methodology to capture patient-reported health data: Report from an NIH Collaboratory roundtable. Healthcare, 2020, 8, 100442.	0.6	3
15	Selecting Outcomes to Ensure Pragmatic Trials Are Relevant to People Living with Dementia. Journal of the American Geriatrics Society, 2020, 68, S55-S61.	1.3	14
16	Clinical Utility and User Perceptions of a Digital System for Electronic Patient-Reported Symptom Monitoring During Routine Cancer Care: Findings From the PRO-TECT Trial. JCO Clinical Cancer Informatics, 2020, 4, 947-957.	1.0	97
17	A review of Kenya's cancer policies to improve access to cancer testing and treatment in the country. Health Research Policy and Systems, 2020, 18, 2.	1.1	13
18	Performance Measures Based on How Adults With Cancer Feel and Function: Stakeholder Recommendations and Feasibility Testing in Six Cancer Centers. JCO Oncology Practice, 2020, 16, e234-e250.	1.4	7

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19	Associations between prostate cancerâ€related anxiety and healthâ€related quality of life. Cancer Medicine, 2020, 9, 4467-4473.	1.3	17
20	Psychometric properties of the FACT-G quality of life scale for family caregivers of cancer patients. Quality of Life Research, 2020, 29, 2241-2252.	1.5	12
21	Electronic Health Record Phenotypes for Identifying Patients with Late-Stage Disease: a Method for Research and Clinical Application. Journal of General Internal Medicine, 2019, 34, 2818-2823.	1.3	11
22	The development, usability, and reliability of the Electronic Patient Visit Assessment (ePVA) for head and neck cancer. MHealth, 2019, 5, 21-21.	0.9	6
23	Usability of PCforMe in Patients With Advanced Cancer Referred to Outpatient Palliative Care: Results of a Randomized, Active-Controlled Pilot Trial. Journal of Pain and Symptom Management, 2019, 58, 382-389.	0.6	12
24	Senior Sway: Using a Mobile Application to Measure Fall Risk. Journal of Geriatric Physical Therapy, 2019, 42, E101-E107.	0.6	2
25	Cabozantinib Versus Mitoxantrone-prednisone in Symptomatic Metastatic Castration-resistant Prostate Cancer: A Randomized Phase 3 Trial with a Primary Pain Endpoint. European Urology, 2019, 75, 929-937.	0.9	41
26	Relationship between sleep and exercise as colorectal cancer survivors transition off treatment. Supportive Care in Cancer, 2018, 26, 2663-2673.	1.0	8
27	Prevalence of patient-reported gastrointestinal symptoms and agreement with clinician toxicity assessments in radiation therapy for anal cancer. Quality of Life Research, 2018, 27, 97-103.	1.5	17
28	Experiences of Inpatient Bone Marrow Transplantation Nurses and Providers Using Electronic Symptom Reporting. Journal of Oncology Practice, 2018, 14, e496-e504.	2.5	3
29	A systematic review of patient reported outcomes in phase II or III clinical trials of myelodysplastic syndromes and acute myeloid leukemia. Leukemia Research, 2018, 70, 106-116.	0.4	9
30	Exploring differences in adverse symptom event grading thresholds between clinicians and patients in the clinical trial setting. Journal of Cancer Research and Clinical Oncology, 2017, 143, 735-743.	1.2	36
31	Feasibility of Patient Reporting of Symptomatic Adverse Events via the Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE) in a Chemoradiotherapy Cooperative Group Multicenter Clinical Trial. International Journal of Radiation Oncology Biology Physics. 2017, 98, 409-418.	0.4	87
32	Evaluation of different recall periods for the US National Cancer Institute's PRO-CTCAE. Clinical Trials, 2017, 14, 255-263.	0.7	58
33	Symptom burden and life challenges reported by adult chordoma patients and their caregivers. Quality of Life Research, 2017, 26, 2237-2244.	1.5	11
34	A Library of Instruments Endorsed by Published Systematic Reviews for Assessing Patients and Their Care Developed by the Palliative Care Research Cooperative. Journal of Palliative Medicine, 2017, 20, 910-911.	0.6	0
35	The prevalence and pattern of chemotherapy-induced peripheral neuropathy among women with breast cancer receiving care in a large community oncology practice. Quality of Life Research, 2017, 26, 2763-2772.	1.5	51
36	Evaluation of pedometry as a patient-centered outcome in patients undergoing hematopoietic cell transplant (HCT): a comparison of pedometry and patient reports of symptoms, health, and quality of life. Quality of Life Research, 2016, 25, 535-546.	1.5	54

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37	The impact of surgical complications on health-related quality of life inÂwomen undergoing gynecologic and gynecologic oncology procedures: a prospective longitudinal cohort study. American Journal of Obstetrics and Gynecology, 2016, 215, 457.e1-457.e13.	0.7	19
38	Mode equivalence and acceptability of tablet computer-, interactive voice response system-, and paper-based administration of the U.S. National Cancer Institute's Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE). Health and Quality of Life Outcomes, 2016, 14, 24.	1.0	91
39	The association between clinician-based common terminology criteria for adverse events (CTCAE) and patient-reported outcomes (PRO): a systematic review. Supportive Care in Cancer, 2016, 24, 3669-3676.	1.0	249
40	Symptom Monitoring With Patient-Reported Outcomes During Routine Cancer Treatment: A Randomized Controlled Trial. Journal of Clinical Oncology, 2016, 34, 557-565.	0.8	1,746
41	Evaluation of mode equivalence of the MSKCC Bowel Function Instrument, LASA Quality of Life, and Subjective Significance Questionnaire items administered by Web, interactive voice response system (IVRS), and paper. Quality of Life Research, 2016, 25, 1123-1130.	1.5	13
42	Emerging uses of patient generated health data in clinical research. Molecular Oncology, 2015, 9, 1018-1024.	2.1	77
43	Comparison of Seven-Day and Repeated 24-Hour Recall of Symptoms in the First 100ÂDays After Hematopoietic Cell Transplantation. Journal of Pain and Symptom Management, 2015, 49, 513-520.	0.6	10
44	Validity and Reliability of the US National Cancer Institute's Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE). JAMA Oncology, 2015, 1, 1051.	3.4	581
45	The Project Data Sphere Initiative: Accelerating Cancer Research by Sharing Data. Oncologist, 2015, 20, 464-e20.	1.9	79
46	Effects of Cabozantinib on Pain and Narcotic Use in Patients with Castration-resistant Prostate Cancer: Results from a Phase 2 Nonrandomized Expansion Cohort. European Urology, 2015, 67, 310-318.	0.9	35
47	Development of the National Cancer Institute's Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE). Journal of the National Cancer Institute, 2014, 106, dju244-dju244.	3.0	689
48	Patient Expectations of Functional Outcomes After Rectal Cancer Surgery. Diseases of the Colon and Rectum, 2014, 57, 151-157.	0.7	28
49	Cognitive interviewing of the US National Cancer Institute's Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE). Quality of Life Research, 2014, 23, 257-269.	1.5	117
50	PRO Data Collection in Clinical Trials Using Mixed Modes: Report of the ISPOR PRO Mixed Modes Good Research Practices Task Force. Value in Health, 2014, 17, 501-516.	0.1	56
51	Prevalence of Pain and Analgesic Use in Men With Metastatic Prostate Cancer Using a Patient-Reported Outcome Measure. Journal of Oncology Practice, 2013, 9, 223-229.	2.5	29
52	Feasibility of Long-Term Patient Self-Reporting of Toxicities From Home via the Internet During Routine Chemotherapy. Journal of Clinical Oncology, 2013, 31, 2580-2585.	0.8	134
53	Reliability of adverse symptom event reporting by clinicians. Quality of Life Research, 2012, 21, 1159-1164.	1.5	186
54	Measurement of Affective and Activity Pain Interference Using the Brief Pain Inventory (BPI): Cancer and Leukemia Group B 70903. Pain Medicine, 2012, 13, 1417-1424.	0.9	17

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55	Comparison of 7-Day Recall and Daily Diary Reports of COPD Symptoms and Impacts. Value in Health, 2012, 15, 466-474.	0.1	22
56	Electronic patientâ€reported outcome systems in oncology clinical practice. Ca-A Cancer Journal for Clinicians, 2012, 62, 336-347.	157.7	279
57	Patient versus clinician symptom reporting during chemoradiation for rectal cancer Journal of Clinical Oncology, 2012, 30, 646-646.	0.8	3
58	Patient-Reported Outcomes vs. Clinician Symptom Reporting During Chemoradiation for Rectal Cancer. Gastrointestinal Cancer Research: GCR, 2012, 5, 119-24.	0.8	28
59	Content validation of two SF-36 subscales for use in type 2 diabetes and non-dialysis chronic kidney disease-related anemia. Quality of Life Research, 2011, 20, 889-901.	1.5	18
60	Impact of Erythropoiesis-Stimulating Agents on Energy and Physical Function in Nondialysis CKD Patients With Anemia: A Systematic Review. American Journal of Kidney Diseases, 2010, 55, 519-534.	2.1	60
61	Evaluation of Inhaler Device Technique in Caregivers of Young Children with Asthma. Pediatric, Allergy, Immunology, and Pulmonology, 2010, 23, 113-120.	0.3	9
62	Comparison of 7-day and repeated 24-hour recall of symptoms of cystic fibrosis. Journal of Cystic Fibrosis, 2010, 9, 419-424.	0.3	23
63	Identifying high-risk asthma with utilization data: a revised HEDIS definition. American Journal of Managed Care, 2008, 14, 450-6.	0.8	6