

Tenbroeck G Smith

List of Publications by Year in descending order

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Version: 2024-02-01

37
papers

2,716
citations

304368

22
h-index

329751

37
g-index

39
all docs

39
docs citations

39
times ranked

3910
citing authors

#	ARTICLE	IF	CITATIONS
1	Adult cancer survivors: How are they faring?. <i>Cancer</i> , 2005, 104, 2565-2576.	2.0	329
2	Residential segregation and disparities in health-related quality of life among Black and White cancer survivors.. <i>Health Psychology</i> , 2011, 30, 137-144.	1.3	277
3	Differences Between Primary Care Physiciansâ€™ and Oncologistsâ€™ Knowledge, Attitudes and Practices Regarding the Care of Cancer Survivors. <i>Journal of General Internal Medicine</i> , 2011, 26, 1403-1410.	1.3	250
4	Symptom burden in cancer survivors 1 year after diagnosis. <i>Cancer</i> , 2011, 117, 2779-2790.	2.0	226
5	Quality of Life of Couples Dealing with Cancer: Dyadic and Individual Adjustment among Breast and Prostate Cancer Survivors and Their Spousal Caregivers. <i>Annals of Behavioral Medicine</i> , 2008, 35, 230-238.	1.7	198
6	The rationale, design, and implementation of the American Cancer Society's studies of cancer survivors. <i>Cancer</i> , 2007, 109, 1-12.	2.0	190
7	A populationâ€based study of prevalence of complementary methods use by cancer survivors. <i>Cancer</i> , 2008, 113, 1048-1057.	2.0	145
8	Equitably improving outcomes for cancer survivors and supporting caregivers: A blueprint for care delivery, research, education, and policy. <i>Ca-A Cancer Journal for Clinicians</i> , 2019, 69, 35-49.	157.7	124
9	Health-related behavior change after cancer: results of the American Cancer Societyâ€™s studies of cancer survivors (SCS). <i>Journal of Cancer Survivorship</i> , 2010, 4, 20-32.	1.5	119
10	Physician roles in the cancer-related follow-up care of cancer survivors. <i>Family Medicine</i> , 2013, 45, 463-74.	0.3	88
11	Health status and life satisfaction among breast cancer survivor peer support volunteers. <i>Psycho-Oncology</i> , 2002, 11, 199-211.	1.0	65
12	Factors Related to the Use of Dietary Supplements by Cancer Survivors. <i>Journal of Alternative and Complementary Medicine</i> , 2009, 15, 673-680.	2.1	65
13	Cancer-related problems, sleep quality, and sleep disturbance among long-term cancer survivors at 9-years post diagnosis. <i>Sleep Medicine</i> , 2020, 65, 177-185.	0.8	57
14	Causal Attribution Among Cancer Survivors of the 10 Most Common Cancers. <i>Journal of Psychosocial Oncology</i> , 2011, 29, 121-140.	0.6	55
15	Perceptions of Patients With Breast and Colon Cancer of the Management of Cancer-Related Pain, Fatigue, and Emotional Distress in Community Oncology. <i>Journal of Clinical Oncology</i> , 2019, 37, 1666-1676.	0.8	50
16	Sociodemographic inequalities in barriers to cancer pain management: a report from the American Cancer Society's Study of Cancer Survivorsâ€™ (SCSâ€™). <i>Psycho-Oncology</i> , 2016, 25, 1212-1221.	1.0	48
17	Populationâ€based cancer registries for qualityâ€ofâ€life research. <i>Cancer</i> , 2013, 119, 2109-2123.	2.0	44
18	A systematic review of large-scale surveys of cancer survivors conducted in North America, 2000â€2011. <i>Journal of Cancer Survivorship</i> , 2012, 6, 115-145.	1.5	42

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19	The relation between cancer patient treatment decision-making roles and quality of life. <i>Cancer</i> , 2013, 119, 2342-2349.	2.0	32
20	The rationale for patient-reported outcomes surveillance in cancer and a reproducible method for achieving it. <i>Cancer</i> , 2016, 122, 344-351.	2.0	32
21	Exploratory Factor Analysis of the Cancer Problems in Living Scale: A Report from the American Cancer Society's Studies of Cancer Survivors. <i>Journal of Pain and Symptom Management</i> , 2009, 37, 676-686.	0.6	31
22	Factors Associated With Health-Related Quality of Life Among Cancer Survivors in the United States. <i>JNCI Cancer Spectrum</i> , 2021, 5, pkaa123.	1.4	26
23	Understanding Long-Term Cancer Survivors' Preferences for Ongoing Medical Care. <i>Journal of General Internal Medicine</i> , 2019, 34, 2091-2097.	1.3	23
24	Sampling in population-based cancer caregivers research. <i>Quality of Life Research</i> , 2009, 18, 981-989.	1.5	18
25	Assessment of the status of A National Action Plan for Cancer Survivorship in the USA. <i>Journal of Cancer Survivorship</i> , 2013, 7, 425-438.	1.5	18
26	Concordance of cancer registry and self-reported race, ethnicity, and cancer type: a report from the American Cancer Society's studies of cancer survivors. <i>Cancer Causes and Control</i> , 2019, 30, 21-29.	0.8	18
27	Cancer survivor perspectives on sharing patient-generated health data with central cancer registries. <i>Quality of Life Research</i> , 2019, 28, 2957-2967.	1.5	14
28	Effectiveness trial of an online self-help intervention for sexual problems after cancer. <i>Journal of Sex and Marital Therapy</i> , 2020, 46, 576-588.	1.0	14
29	Employment experience of cancer survivors 2 years post-diagnosis in the Study of Cancer Survivors-I. <i>Journal of Cancer Survivorship</i> , 2012, 6, 210-218.	1.5	12
30	Investigating relationships among cancer survivors' engagement in an online support community, social support perceptions, well-being, and moderating effects of existing (offline) social support. <i>Supportive Care in Cancer</i> , 2020, 28, 3791-3799.	1.0	10
31	Exploring measurement invariance by gender in the profile of mood states depression subscale among cancer survivors. <i>Quality of Life Research</i> , 2017, 26, 171-175.	1.5	8
32	Has the Quality of Patient-Provider Communication About Survivorship Care Improved?. <i>Journal of Oncology Practice</i> , 2019, 15, e916-e924.	2.5	7
33	Performance Measures Based on How Adults With Cancer Feel and Function: Stakeholder Recommendations and Feasibility Testing in Six Cancer Centers. <i>JCO Oncology Practice</i> , 2020, 16, e234-e250.	1.4	7
34	Oncology team perspectives on distress screening: a multisite study of a well-established use of patient-reported outcomes for clinical assessment. <i>Supportive Care in Cancer</i> , 2022, 30, 1261-1271.	1.0	5
35	Theory-Based Evaluation of an Online Cancer Fatigue Class. <i>Journal of Cancer Education</i> , 2010, 25, 422-430.	0.6	4
36	The development and acceptability of symptom management quality improvement reports based on patient-reported data: an overview of methods used in PROSSES. <i>Quality of Life Research</i> , 2016, 25, 2833-2843.	1.5	4

#	ARTICLE	IF	CITATIONS
37	Research Participantsâ€™ Perspectives on Using an Electronic Portal for Engagement and Data Collection: Focus Group Results From a Large Epidemiologic Cohort. <i>Journal of Medical Internet Research</i> , 2020, 22, e18556.	2.1	0