

Sydney M Dy

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/732336/publications.pdf>

Version: 2024-02-01

75
papers

2,085
citations

236925

25
h-index

254184

43
g-index

76
all docs

76
docs citations

76
times ranked

3010
citing authors

#	ARTICLE	IF	CITATIONS
1	Evidence-Based Standards for Cancer Pain Management. <i>Journal of Clinical Oncology</i> , 2008, 26, 3879-3885.	1.6	149
2	Populations and Interventions for Palliative and End-of-Life Care: A Systematic Review. <i>Journal of Palliative Medicine</i> , 2016, 19, 995-1008.	1.1	128
3	Pharmacotherapy for diabetic peripheral neuropathy pain and quality of life. <i>Neurology</i> , 2017, 88, 1958-1967.	1.1	127
4	End-of-Life Care. <i>Archives of Internal Medicine</i> , 2007, 167, 40.	3.8	124
5	Evidence-Based Recommendations for Cancer Fatigue, Anorexia, Depression, and Dyspnea. <i>Journal of Clinical Oncology</i> , 2008, 26, 3886-3895.	1.6	115
6	Critical Pathway Effectiveness: Assessing the Impact of Patient, Hospital Care, and Pathway Characteristics Using Qualitative Comparative Analysis. <i>Health Services Research</i> , 2005, 40, 499-516.	2.0	106
7	Key concepts relevant to quality of complex and shared decision-making in health care: A literature review. <i>Social Science and Medicine</i> , 2012, 74, 582-587.	3.8	92
8	Multimorbidity and Decision-Making Preferences Among Older Adults. <i>Annals of Family Medicine</i> , 2017, 15, 546-551.	1.9	78
9	A Systematic Review of Communication Quality Improvement Interventions for Patients with Advanced and Serious Illness. <i>Journal of General Internal Medicine</i> , 2013, 28, 570-577.	2.6	76
10	Cancer Qualityâ€ASSIST supportive oncology quality indicator set. <i>Cancer</i> , 2010, 116, 3267-3275.	4.1	66
11	Tell Usâ„¢: A Web-Based Tool for Improving Communication Among Patients, Families, and Providers in Hospice and Palliative Care Through Systematic Data Specification, Collection, and Use. <i>Journal of Pain and Symptom Management</i> , 2011, 42, 526-534.	1.2	57
12	Implementation of the World Health Organization Trauma Care Checklist Program in 11 Centers Across Multiple Economic Strata: Effect on Care Process Measures. <i>World Journal of Surgery</i> , 2017, 41, 954-962.	1.6	57
13	A Framework for Assessing Quality Indicators for Cancer Care at the End of Life. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 903-912.	1.2	51
14	Does Caregiving Strain Increase as Patients With and Without Dementia Approach the End of Life?. <i>Journal of Pain and Symptom Management</i> , 2019, 57, 199-208.e2.	1.2	45
15	Non-pharmacologic treatments for symptoms of diabetic peripheral neuropathy: a systematic review. <i>Current Medical Research and Opinion</i> , 2019, 35, 15-25.	1.9	44
16	Is Race/Ethnicity Related to the Presence or Severity of Pain in Colorectal and Lung Cancer?. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 1050-1059.	1.2	42
17	<p>Analysis of Patient Preferences in Lung Cancer â€“ Estimating Acceptable Tradeoffs Between Treatment Benefit and Side Effects<p>. <i>Patient Preference and Adherence</i> , 2020, Volume 14, 927-937.	1.8	39
18	Lower Patient Ratings of Physician Communication Are Associated With Unmet Need for Symptom Management in Patients With Lung and Colorectal Cancer. <i>Journal of Oncology Practice</i> , 2016, 12, e654-e669.	2.5	37

#	ARTICLE	IF	CITATIONS
19	Comorbid illnesses and health care utilization among Medicare beneficiaries with lung cancer. <i>Critical Reviews in Oncology/Hematology</i> , 2006, 59, 218-225.	4.4	36
20	The Quality Imperative for Palliative Care. <i>Journal of Pain and Symptom Management</i> , 2015, 49, 243-253.	1.2	33
21	Association Between Palliative Case Management and Utilization of Inpatient, Intensive Care Unit, Emergency Department, and Hospice in Medicaid Beneficiaries. <i>American Journal of Hospice and Palliative Medicine</i> , 2015, 32, 216-220.	1.4	33
22	Patient- and Caregiver-Reported Assessment Tools for Palliative Care: Summary of the 2017 Agency for Healthcare Research and Quality Technical Brief. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 961-972.e16.	1.2	33
23	Fatigability and endurance performance in cancer survivors: Analyses from the Baltimore Longitudinal Study of Aging. <i>Cancer</i> , 2018, 124, 1279-1287.	4.1	33
24	Continuity, Coordination, and Transitions of Care for Patients with Serious and Advanced Illness: A Systematic Review of Interventions. <i>Journal of Palliative Medicine</i> , 2013, 16, 436-445.	1.1	31
25	Quality of care and short- and long-term outcomes of laryngeal cancer care in the elderly. <i>Laryngoscope</i> , 2015, 125, 2323-2329.	2.0	27
26	Pharmacologic Interventions for Breathlessness in Patients With Advanced Cancer. <i>JAMA Network Open</i> , 2021, 4, e2037632.	5.9	26
27	A randomized intervention involving family to improve communication in breast cancer care. <i>Npj Breast Cancer</i> , 2021, 7, 14.	5.2	23
28	Financial Burden of Drugs Prescribed for Cancer-Associated Symptoms. <i>JCO Oncology Practice</i> , 2022, 18, 140-147.	2.9	23
29	Health-Related Quality of Life and Functional Status Quality Indicators for Older Persons with Multiple Chronic Conditions. <i>Journal of the American Geriatrics Society</i> , 2013, 61, 2120-2127.	2.6	22
30	Contrasting characteristics of daily physical activity in older adults by cancer history. <i>Cancer</i> , 2018, 124, 4692-4699.	4.1	22
31	Towards a More Patient-Centered Approach to Medication Safety. <i>Journal of Patient Experience</i> , 2018, 5, 83-87.	0.9	22
32	To Stent or Not to Stent: An Evidence-Based Approach to Palliative Procedures at the End of Life. <i>Journal of Pain and Symptom Management</i> , 2012, 43, 795-801.	1.2	18
33	Patient Characteristics and End-of-Life Health Care Utilization Among Medicare Beneficiaries in 1989 and 1999. <i>Medical Care</i> , 2007, 45, 926-930.	2.4	17
34	Palliative Care for Cancer Survivors. <i>Medical Clinics of North America</i> , 2017, 101, 1181-1196.	2.5	17
35	Association of Magnet Status and Nurse Staffing With Improvements in Patient Experience With Hospital Care, 2008-2015. <i>Medical Care</i> , 2018, 56, 111-120.	2.4	17
36	Nonpharmacological Interventions for Managing Breathlessness in Patients With Advanced Cancer. <i>JAMA Oncology</i> , 2021, 7, 290.	7.1	16

#	ARTICLE	IF	CITATIONS
37	Impact of Medicare Annual Wellness Visits on Uptake of Depression Screening. <i>Psychiatric Services</i> , 2015, 66, 1207-1212.	2.0	15
38	Increasing Advance Care Planning Completion at an Academic Internal Medicine Outpatient Clinic. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 383-386.	1.2	15
39	Evaluating the Quality of Supportive Oncology Using Patient-Reported Data. <i>Journal of Oncology Practice</i> , 2014, 10, e223-e230.	2.5	13
40	Project Transform: engaging patient advocates to share their perspectives on improving research, treatment and policy. <i>Current Medical Research and Opinion</i> , 2018, 34, 1755-1762.	1.9	11
41	Patient-reported quality of care and pain severity in cancer. <i>Palliative and Supportive Care</i> , 2015, 13, 875-884.	1.0	10
42	Preparing Older Patients With Serious Illness for Advance Care Planning Discussions in Primary Care. <i>Journal of Pain and Symptom Management</i> , 2019, 58, 244-251.e1.	1.2	8
43	Quality of Preventive and Chronic Illness Care for Insured Adults With Opioid Use Disorder. <i>JAMA Network Open</i> , 2021, 4, e214925.	5.9	8
44	Prescribing of low- versus high-cost Part B drugs in Medicare Advantage and traditional Medicare. <i>Health Services Research</i> , 2022, 57, 537-547.	2.0	8
45	Live, Learn, Pass It on. <i>Journal of Patient Experience</i> , 2017, 4, 162-168.	0.9	7
46	Older Patients' Perspectives on Quality of Serious Illness Care in Primary Care. <i>American Journal of Hospice and Palliative Medicine</i> , 2018, 35, 1330-1336.	1.4	7
47	The Johns Hopkins Primary Care for Cancer Survivor Clinic: lessons learned in our first 4 years. <i>Journal of Cancer Survivorship</i> , 2020, 14, 19-25.	2.9	7
48	Selecting Patient-Reported Outcome Measures to Contribute to Primary Care Performance Measurement: a Mixed Methods Approach. <i>Journal of General Internal Medicine</i> , 2020, 35, 2687-2697.	2.6	7
49	The Palliative Care "Promoting Access and Improvement of the Cancer Experience (PC-PAICE) Project in India: A Multisite International Quality Improvement Collaborative. <i>Journal of Pain and Symptom Management</i> , 2021, 61, 190-197.	1.2	7
50	Measuring the quality of palliative care and supportive oncology: principles and practice. <i>The Journal of Supportive Oncology</i> , 2013, 11, 160-164.	2.3	7
51	Care pathways and patient safety: key concepts, patient outcomes and related interventions. <i>International Journal of Care Pathways</i> , 2010, 14, 124-128.	0.5	6
52	A Quality Improvement Initiative for Improving Appropriateness of Referrals From a Cancer Center to Subacute Rehabilitation. <i>Journal of Pain and Symptom Management</i> , 2014, 48, 127-131.	1.2	6
53	Advance Care Planning Shared Decision-Making Tools for Non-Cancer Chronic Serious Illness: A Mixed Method Systematic Review. <i>American Journal of Hospice and Palliative Medicine</i> , 2021, 38, 1526-1535.	1.4	6
54	A qualitative exploration of favorite patients in primary care. <i>Patient Education and Counseling</i> , 2016, 99, 1888-1893.	2.2	5

#	ARTICLE	IF	CITATIONS
55	Quality of Buprenorphine Care for Insured Adults With Opioid Use Disorder. <i>Medical Care</i> , 2021, 59, 393-401.	2.4	5
56	Use of Peripheral μ -Opioid Receptor Antagonists for Treating Opioid-Induced Constipation among US Medicare Beneficiaries from 2014 to 2018. <i>Journal of Palliative Medicine</i> , 2021, 24, 1236-1239.	1.1	5
57	Implementation and Effectiveness of Integrating Palliative Care Into Ambulatory Care of Noncancer Serious Chronic Illness: Mixed Methods Review and Meta-Analysis. <i>Annals of Family Medicine</i> , 2022, 20, 77-83.	1.9	5
58	Evaluation of the Measuring and Improving Quality in Palliative Care Survey. <i>Journal of Oncology Practice</i> , 2018, 14, e834-e843.	2.5	4
59	Pain Experiences of Adults With Sickle Cell Disease and Hematopoietic Stem Cell Transplantation: A Qualitative Study. <i>Pain Medicine</i> , 2021, 22, 1753-1759.	1.9	4
60	The Key Role of Nonpharmacologic Management of Cachexia in Persons With Advanced Illness. <i>JAMA Internal Medicine</i> , 2021, 181, 978-979.	5.1	4
61	End-of-Life Health Care Utilization in Hospitals with Compared to Those without Palliative Care Programs. <i>Journal of Palliative Medicine</i> , 2014, 17, 877-878.	1.1	3
62	Asking questions of a palliative care nurse practitioner on a pancreatic cancer website. <i>Palliative and Supportive Care</i> , 2015, 13, 787-793.	1.0	3
63	A Survey to Evaluate Facilitators and Barriers to Quality Measurement and Improvement: Adapting Tools for Implementation Research in Palliative Care Programs. <i>Journal of Pain and Symptom Management</i> , 2017, 54, 806-814.	1.2	3
64	What matters when doctors die: A qualitative study of family perspectives. <i>PLoS ONE</i> , 2020, 15, e0235138.	2.5	3
65	Perceptions of Facilitators and Barriers to Measuring and Improving Quality in Palliative Care Programs. <i>American Journal of Hospice and Palliative Medicine</i> , 2020, 37, 1022-1028.	1.4	2
66	Patient perceptions of safety in primary care: a qualitative study to inform care. <i>Current Medical Research and Opinion</i> , 2021, 37, 1-9.	1.9	2
67	Conformance to Depression Process Measures of Medicare Part B Beneficiaries in Primary Care Settings. <i>Journal of the American Geriatrics Society</i> , 2015, 63, 1338-1345.	2.6	1
68	Outcomes After Referral for Acute Inpatient Rehabilitation in Hospitalized Patients With Cancer. <i>Journal of Pain and Symptom Management</i> , 2015, 50, e1-e3.	1.2	1
69	Patient Safety in Primary Care: Conceptual Meanings to the Health Care Team and Patients. <i>Journal of the American Board of Family Medicine</i> , 2020, 33, 754-764.	1.5	1
70	Challenges and Strategies for Patient Safety in Primary Care: A Qualitative Study. <i>American Journal of Medical Quality</i> , 2022, Publish Ahead of Print, .	0.5	1
71	INVOLVING FAMILY TO IMPROVE COMMUNICATION IN PRIMARY CARE FOR PRIMARY CARE PATIENTS WITH DEMENTIA. <i>Innovation in Aging</i> , 2019, 3, S364-S364.	0.1	0
72	What matters when doctors die: A qualitative study of family perspectives. , 2020, 15, e0235138.		0

#	ARTICLE	IF	CITATIONS
73	What matters when doctors die: A qualitative study of family perspectives. , 2020, 15, e0235138.		0
74	What matters when doctors die: A qualitative study of family perspectives. , 2020, 15, e0235138.		0
75	What matters when doctors die: A qualitative study of family perspectives. , 2020, 15, e0235138.		0