

Jennifer W Mack

List of Publications by Year in descending order

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Version: 2024-02-01

155
papers

10,251
citations

81434

41
h-index

39744

98
g-index

156
all docs

156
docs citations

156
times ranked

7695
citing authors

#	ARTICLE	IF	CITATIONS
1	Caregiver Reported Quality of End-of-Life Care of Adolescent and Young Adult Decedents With Cancer. <i>Journal of Palliative Care</i> , 2022, 37, 87-92.	0.4	5
2	Conflicting goals and obligations: Tensions affecting communication in pediatric oncology. <i>Patient Education and Counseling</i> , 2022, 105, 56-61.	1.0	7
3	A Reliability Generalization Meta-Analysis of 17 Patient-Reported Outcome Measures for Positive Psychosocial Constructs in Children, Adolescents, and Young Adults with Cancer. <i>Journal of Adolescent and Young Adult Oncology</i> , 2022, 11, 163-172.	0.7	3
4	Multilevel barriers and facilitators of communication in pediatric oncology: A systematic review. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29405.	0.8	9
5	Acceptable, hopeful, and useful: development and mixed-method evaluation of an educational tool about reproductive options for people with sickle cell disease or trait. <i>Journal of Assisted Reproduction and Genetics</i> , 2022, 39, 183-193.	1.2	8
6	Early parental knowledge of late effect risks in children with cancer. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29473.	0.8	4
7	Effectiveness of Pediatric Concurrent Hospice Care to Improve Continuity of Care. <i>American Journal of Hospice and Palliative Medicine</i> , 2022, 39, 1129-1136.	0.8	8
8	Missing Voices: Lessons Learned from Nonparticipating Caregivers in Palliative Care Research. <i>Journal of Palliative Medicine</i> , 2022, 25, 455-460.	0.6	1
9	Health Literacy and Clinical Outcomes Following Hematopoietic Stem-Cell Transplantation. <i>JCO Oncology Practice</i> , 2022, , OP2100049.	1.4	1
10	Interdependent functions of communication with adolescents and young adults in oncology. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29588.	0.8	13
11	Lack of Concordance in Symptomatic Adverse Event Reporting by Children, Clinicians, and Caregivers: Implications for Cancer Clinical Trials. <i>Journal of Clinical Oncology</i> , 2022, 40, 1623-1634.	0.8	27
12	Racial and ethnic disparities in communication study enrollment for young people with cancer: A descriptive analysis of the literature. <i>Patient Education and Counseling</i> , 2022, 105, 2067-2073.	1.0	4
13	Bereaved Caregivers Perspectives of Negative Communication Experiences Near the End of Life for Adolescents and Young Adults with Cancer. <i>Journal of Adolescent and Young Adult Oncology</i> , 2022, 11, 498-505.	0.7	4
14	Pediatric Concurrent Hospice Care: Cost Implications of a Hybrid Payment Model. <i>American Journal of Hospice and Palliative Medicine</i> , 2022, , 104990912210893.	0.8	2
15	Co-management of communication and care in adolescent and young adult oncology. <i>Pediatric Blood and Cancer</i> , 2022, 69, .	0.8	4
16	Role of Surgery in Rhabdomyosarcoma of the Head and Neck in Children. <i>Laryngoscope</i> , 2021, 131, E984-E992.	1.1	11
17	Characteristics of uncertainty in advanced pediatric cancer conversations. <i>Patient Education and Counseling</i> , 2021, 104, 1066-1074.	1.0	7
18	Validation of the caregiver Pediatric Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events measure. <i>Cancer</i> , 2021, 127, 1483-1494.	2.0	15

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19	Subjective Toxicity Profiles of Children in Treatment for Cancer: A New Guide to Supportive Care?. Journal of Pain and Symptom Management, 2021, 61, 1188-1195.e2.	0.6	15
20	Parental views on communication between children and clinicians in pediatric oncology: a qualitative study. Supportive Care in Cancer, 2021, 29, 4957-4968.	1.0	19
21	Multilevel barriers to communication in pediatric oncology: Clinicians' perspectives. Cancer, 2021, 127, 2130-2138.	2.0	19
22	Patients'™ understanding of communication about palliative care and health condition in Japanese patients with unresectable or recurrent cancer: a cross-sectional survey. Annals of Palliative Medicine, 2021, 10, 2650-2661.	0.5	4
23	“Don't be afraid to speak up” Communication advice from parents and clinicians of children with cancer. Pediatric Blood and Cancer, 2021, 68, e29052.	0.8	5
24	Medical complexity and concurrent hospice care: A national study of Medicaid children from 2011 to 2013. Journal for Specialists in Pediatric Nursing, 2021, 26, e12333.	0.6	12
25	“What Matters to Me” What pediatric stem cell transplant patients want their providers to know. Journal of Psychosocial Oncology, 2021, 39, 586-593.	0.6	1
26	Differences in characteristics of children with cancer who receive standard versus concurrent hospice care. Pediatric Blood and Cancer, 2021, 68, e29106.	0.8	5
27	Evaluation of Patient-Reported Outcome Measures of Positive Psychosocial Constructs in Children and Adolescent/Young Adults with Cancer: A Systematic Review of Measurement Properties. Journal of Adolescent and Young Adult Oncology, 2021, , .	0.7	7
28	The evolution of parents'™ beliefs about childhood cancer during diagnostic communication: a qualitative study in Guatemala. BMJ Global Health, 2021, 6, e004653.	2.0	7
29	Prognostic Communication Between Oncologists and Parents of Children With Advanced Cancer. Pediatrics, 2021, 147, .	1.0	20
30	Assume It Will Break: Parental Perspectives on Negative Communication Experiences in Pediatric Oncology. JCO Oncology Practice, 2021, 17, e859-e871.	1.4	12
31	Factors related to specialized palliative care use and aggressive care at end of life in Japanese patients with advanced solid cancers: a cohort study. Supportive Care in Cancer, 2021, 29, 7805-7813.	1.0	2
32	Validity and reliability of the Japanese version of the Peace, Equanimity, and Acceptance in the Cancer Experience (PEACE) questionnaire. Palliative and Supportive Care, 2021, , 1-7.	0.6	0
33	Identification of Evidence for Key Positive Psychological Constructs in Pediatric and Adolescent/Young Adult Patients with Cancer: A Scoping Review. Journal of Adolescent and Young Adult Oncology, 2021, 10, 247-259.	0.7	7
34	Abstract 10: The Evolution of Parents' Beliefs about Childhood Cancer: A Qualitative Study in Guatemala. , 2021, , .		0
35	Who is coordinating pediatric concurrent hospice care?. Journal of Pain and Symptom Management, 2021, 62, e1-e4.	0.6	4
36	Patients, caregivers, and clinicians differ in performance status ratings: Implications for pediatric cancer clinical trials. Cancer, 2021, 127, 3664-3670.	2.0	15

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37	Broaching goals of care conversations in advancing pediatric cancer. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29270.	0.8	7
38	Clinicians' Perspectives on the Functions of Communication in Pediatric Oncology. <i>Journal of Palliative Medicine</i> , 2021, 24, 1545-1549.	0.6	5
39	Thinking ahead: Parents' worries about late effects of childhood cancer treatment. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29335.	0.8	6
40	Communication Priorities and Experiences of Caregivers of Children With Cancer in Guatemala. <i>JCO Global Oncology</i> , 2021, 7, 1529-1536.	0.8	5
41	A National Study to Compare Effective Management of Constipation in Children Receiving Concurrent Versus Standard Hospice Care. <i>Journal of Hospice and Palliative Nursing</i> , 2021, Publish Ahead of Print, .	0.5	0
42	Parent and Clinician Perspectives on Challenging Parent-Clinician Relationships in Pediatric Oncology. <i>JAMA Network Open</i> , 2021, 4, e2132138.	2.8	5
43	Mapping child and adolescent self-reported symptom data to clinician-reported adverse event grading to improve pediatric oncology care and research. <i>Cancer</i> , 2020, 126, 140-147.	2.0	14
44	Longitudinal prognostic communication needs of adolescents and young adults with cancer. <i>Cancer</i> , 2020, 126, 400-407.	2.0	26
45	Unrealistic parental expectations for cure in poor-prognosis childhood cancer. <i>Cancer</i> , 2020, 126, 416-424.	2.0	34
46	The evolution of regret: decision-making for parents of children with cancer. <i>Supportive Care in Cancer</i> , 2020, 28, 1215-1222.	1.0	20
47	Racial and Ethnic Differences in Parental Decision-Making Roles in Pediatric Oncology. <i>Journal of Palliative Medicine</i> , 2020, 23, 192-197.	0.6	15
48	End-of-life care among adolescent and young adult patients with cancer living in poverty. <i>Cancer</i> , 2020, 126, 886-893.	2.0	24
49	Expanding construct validity of established and new PROMIS Pediatric measures for children and adolescents receiving cancer treatment. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28160.	0.8	36
50	Emotional Communication in Advanced Pediatric Cancer Conversations. <i>Journal of Pain and Symptom Management</i> , 2020, 59, 808-817.e2.	0.6	21
51	Peace of mind among adolescents and young adults with cancer. <i>Psycho-Oncology</i> , 2020, 29, 572-578.	1.0	7
52	Communication during childhood cancer: Systematic review of patient perspectives. <i>Cancer</i> , 2020, 126, 701-716.	2.0	50
53	Alveolar rhabdomyosarcoma presenting as a pleural effusion: An atypical presentation of a malignancy. <i>Pediatric Pulmonology</i> , 2020, 55, 3231-3233.	1.0	0
54	Pediatric cancer communication in low- and middle-income countries: A scoping review. <i>Cancer</i> , 2020, 126, 5030-5039.	2.0	18

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55	The use of interval-compressed chemotherapy with the addition of vincristine, irinotecan, and temozolomide for pediatric patients with newly diagnosed desmoplastic small round cell tumor. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28559.	0.8	13
56	Impact of cancer on school, work, and financial independence among adolescents and young adults. <i>Cancer</i> , 2020, 126, 4400-4406.	2.0	24
57	Interdisciplinary care of pediatric oncology patients in Central America and the Caribbean. <i>Cancer</i> , 2020, 127, 2579-2586.	2.0	4
58	Qualitative Study of Pediatric Early Warning Systems™ Impact on Interdisciplinary Communication in Two Pediatric Oncology Hospitals With Varying Resources. <i>JCO Global Oncology</i> , 2020, 6, 1079-1086.	0.8	24
59	Care experiences that foster trust between parents and physicians of children with cancer. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28399.	0.8	16
60	Agreement Between Child Self-report and Caregiver-Proxy Report for Symptoms and Functioning of Children Undergoing Cancer Treatment. <i>JAMA Pediatrics</i> , 2020, 174, e202861.	3.3	73
61	Communication in Pediatric Oncology: A Qualitative Study. <i>Pediatrics</i> , 2020, 146, .	1.0	52
62	Early information needs of adolescents and young adults about late effects of cancer treatment. <i>Cancer</i> , 2020, 126, 3281-3288.	2.0	20
63	Parental Considerations Regarding Cure and Late Effects for Children With Cancer. <i>Pediatrics</i> , 2020, 145, e20193552.	1.0	10
64	Psychosocial care for children receiving chimeric antigen receptor (CAR) T-cell therapy. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28249.	0.8	8
65	Racial and Ethnic Differences in Communication and Care for Children With Advanced Cancer. <i>Journal of Pain and Symptom Management</i> , 2020, 60, 782-789.	0.6	27
66	Exercise and Well-being in Adult Survivors of Childhood Cancer—Time for Interventions. <i>JAMA Oncology</i> , 2020, 6, 1170.	3.4	0
67	Pediatric Concurrent Hospice Care. <i>Journal of Hospice and Palliative Nursing</i> , 2020, 22, 238-245.	0.5	28
68	On listening. <i>Cancer</i> , 2020, 126, 1828-1831.	2.0	1
69	Oncologists' Communication and Decision-Making Behaviors Affect Perceptions of Sensitive Information Exchange for Adolescent and Young Adult Patients. <i>Journal of Palliative Medicine</i> , 2020, 23, 310-311.	0.6	0
70	Validity and Reliability of the Pediatric Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events. <i>Journal of the National Cancer Institute</i> , 2020, 112, 1143-1152.	3.0	59
71	Teamwork in prognostic communication: Addressing bottlenecks and barriers. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28192.	0.8	9
72	Humour and laughing in patients with prolonged incurable cancer: an ethnographic study in a comprehensive cancer centre. <i>Quality of Life Research</i> , 2020, 29, 2425-2434.	1.5	14

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73	Variation In State Medicaid Implementation Of The ACA: The Case Of Concurrent Care For Children. Health Affairs, 2020, 39, 1770-1775.	2.5	30
74	Knowing versus doing: The value of behavioral change models for emotional communication in oncology. Patient Education and Counseling, 2019, 102, 2344-2348.	1.0	8
75	Persistently low peace of mind in parents of cancer patients: A five-year follow-up study. Pediatric Blood and Cancer, 2019, 66, e27609.	0.8	7
76	Things that matter: Adolescent and young adult patients'™ priorities during cancer care. Pediatric Blood and Cancer, 2019, 66, e27883.	0.8	22
77	Effective Communication for Newly Diagnosed Pediatric Patients With Cancer: Considerations for the Patients, Family Members, Providers, and Multidisciplinary Team. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2019, 39, 573-581.	1.8	10
78	Adolescent and Young Adult Cancer Patients'™ Experiences With Treatment Decision-making. Pediatrics, 2019, 143, .	1.0	52
79	Inclusion of children in the initial conversation about their cancer diagnosis: impact on parent experiences of the communication process. Supportive Care in Cancer, 2019, 27, 1319-1324.	1.0	7
80	How Do Blood Cancer Doctors Discuss Prognosis? Findings from a National Survey of Hematologic Oncologists. Journal of Palliative Medicine, 2019, 22, 677-684.	0.6	20
81	Mucoepidermoid carcinoma of the head and neck in children. International Journal of Pediatric Otorhinolaryngology, 2019, 120, 93-99.	0.4	16
82	Duality of Purpose: Participant and Parent Understanding of the Purpose of Genomic Tumor Profiling Research Among Children and Young Adults With Solid Tumors. JCO Precision Oncology, 2019, 3, 1-17.	1.5	11
83	Communication interventions in adult and pediatric oncology: A scoping review and analysis of behavioral targets. PLoS ONE, 2019, 14, e0221536.	1.1	23
84	Decisional burden among parents of children with cancer. Cancer, 2019, 125, 1365-1372.	2.0	27
85	Squamous cell carcinoma of the head and neck in children. International Journal of Pediatric Otorhinolaryngology, 2019, 117, 131-137.	0.4	9
86	Adolescent patient involvement in discussions about relapsed or refractory cancer with oncology clinicians. Pediatric Blood and Cancer, 2018, 65, e26918.	0.8	9
87	Parent perspectives on information about late effects of childhood cancer treatment and their role in initial treatment decision making. Pediatric Blood and Cancer, 2018, 65, e26978.	0.8	17
88	Reply to Cross-cultural communication in pediatric oncology: Catch them when they're young. Cancer, 2018, 124, 1517-1517.	2.0	1
89	Parent understanding of the risk of future limitations secondary to pediatric cancer treatment. Pediatric Blood and Cancer, 2018, 65, e27020.	0.8	20
90	Sources of parental hope in pediatric oncology. Pediatric Blood and Cancer, 2018, 65, e26981.	0.8	28

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91	Intended and unintended consequences: Ethics, communication, and prognostic disclosure in pediatric oncology. <i>Cancer</i> , 2018, 124, 1232-1241.	2.0	35
92	Communication in pediatric oncology: State of the field and research agenda. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26727.	0.8	83
93	Fatalism and educational disparities in beliefs about the curability of advanced cancer. <i>Patient Education and Counseling</i> , 2018, 101, 113-118.	1.0	27
94	How Parents of Children With Cancer Learn About Their Children's Prognosis. <i>Pediatrics</i> , 2018, 141, .	1.0	29
95	Longitudinal parental preferences for late effects communication during cancer treatment. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26760.	0.8	11
96	Communication About Prognosis With Adolescent and Young Adult Patients With Cancer: Information Needs, Prognostic Awareness, and Outcomes of Disclosure. <i>Journal of Clinical Oncology</i> , 2018, 36, 1861-1867.	0.8	90
97	Parental distress and desire for information regarding long-term implications of pediatric cancer treatment. <i>Cancer</i> , 2018, 124, 4529-4537.	2.0	25
98	The PRISM intervention for adolescents and young adults with cancer: Paying attention to the patient as a whole person. <i>Cancer</i> , 2018, 124, 3802-3805.	2.0	2
99	Closing the Gap on Pediatric Palliative Oncology Disparities. <i>Seminars in Oncology Nursing</i> , 2018, 34, 294-302.	0.7	6
100	Patient perceptions of curability and physician-reported disclosures of incurability in Japanese patients with unresectable/recurrent cancer: a cross-sectional survey. <i>Japanese Journal of Clinical Oncology</i> , 2018, 48, 913-919.	0.6	8
101	Holding hope. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26570.	0.8	2
102	Prognostic disclosures over time: Parental preferences and physician practices. <i>Cancer</i> , 2017, 123, 4031-4038.	2.0	57
103	Pediatric Palliative Care "A Shared Priority. <i>JAMA Oncology</i> , 2017, 3, 1220.	3.4	2
104	Evaluation of Patient and Family Outpatient Complaints as a Strategy to Prioritize Efforts to Improve Cancer Care Delivery. <i>Joint Commission Journal on Quality and Patient Safety</i> , 2017, 43, 498-507.	0.4	32
105	Disparities in prognosis communication among parents of children with cancer: The impact of race and ethnicity. <i>Cancer</i> , 2017, 123, 3995-4003.	2.0	73
106	A Research Agenda for Communication Between Health Care Professionals and Patients Living With Serious Illness. <i>JAMA Internal Medicine</i> , 2017, 177, 1361.	2.6	175
107	Communication preferences of pediatric cancer patients: talking about prognosis and their future life. <i>Supportive Care in Cancer</i> , 2017, 25, 769-774.	1.0	58
108	Difficult relationships between parents and physicians of children with cancer: A qualitative study of parent and physician perspectives. <i>Cancer</i> , 2017, 123, 675-681.	2.0	31

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109	Eliciting the child's voice in adverse event reporting in oncology trials: Cognitive interview findings from the Pediatric Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events initiative. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26261.	0.8	50
110	How a Child With Cancer Moved From Vulnerability to Resilience. <i>Journal of Clinical Oncology</i> , 2017, 35, 3169-3171.	0.8	3
111	Cancer treatment decision-making among young adults with lung and colorectal cancer: a comparison with adults in middle age. <i>Psycho-Oncology</i> , 2016, 25, 1085-1091.	1.0	23
112	Intensity of Medical Interventions between Diagnosis and Death in Patients with Advanced Lung and Colorectal Cancer: A CanCORS Analysis. <i>Journal of Palliative Medicine</i> , 2016, 19, 42-50.	0.6	13
113	Communication Skills Training in Pediatric Oncology: Moving Beyond Role Modeling. <i>Pediatric Blood and Cancer</i> , 2016, 63, 966-972.	0.8	59
114	What Adult Cancer Care Can Learn From Pediatrics. <i>Journal of Oncology Practice</i> , 2016, 12, 765-767.	2.5	5
115	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
116	Prognostic Disclosures to Children: A Historical Perspective. <i>Pediatrics</i> , 2016, 138, .	1.0	92
117	Decisional Regret Among Parents of Children With Cancer. <i>Journal of Clinical Oncology</i> , 2016, 34, 4023-4029.	0.8	83
118	Patient/parent perspectives on genomic tumor profiling of pediatric solid tumors: The Individualized Cancer Therapy (iCat) experience. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1974-1982.	0.8	49
119	Care in the Final Month of Life among Adolescent and Young Adult Cancer Patients in Kaiser Permanente Southern California. <i>Journal of Palliative Medicine</i> , 2016, 19, 1136-1141.	0.6	20
120	Home care for children with multiple complex chronic conditions at the end of life: The choice of hospice versus home health. <i>Home Health Care Services Quarterly</i> , 2016, 35, 101-111.	0.3	20
121	Parental preparedness for late effects and long-term quality of life in survivors of childhood cancer. <i>Cancer</i> , 2016, 122, 2587-2594.	2.0	33
122	“Don't Try to Cover the Sky with Your Hands”: Parents' Experiences with Prognosis Communication About Their Children with Advanced Cancer. <i>Journal of Palliative Medicine</i> , 2016, 19, 626-631.	0.6	94
123	Clusters of Multiple Complex Chronic Conditions: A Latent Class Analysis of Children at End of Life. <i>Journal of Pain and Symptom Management</i> , 2016, 51, 868-874.	0.6	22
124	When to say when: How aggressively to care for children with multiply relapsed cancer?. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1119-1120.	0.8	2
125	High Intensity of End-of-Life Care Among Adolescent and Young Adult Cancer Patients in the New York State Medicaid Program. <i>Medical Care</i> , 2015, 53, 1018-1026.	1.1	83
126	Deeper Conversations Need Not Wait Until the End. <i>Journal of Clinical Oncology</i> , 2015, 33, 3974-3974.	0.8	7

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127	End-of-Life Care Intensity Among Adolescent and Young Adult Patients With Cancer in Kaiser Permanente Southern California. <i>JAMA Oncology</i> , 2015, 1, 592.	3.4	86
128	Patient beliefs that chemotherapy may be curative and care received at the end of life among patients with metastatic lung and colorectal cancer. <i>Cancer</i> , 2015, 121, 1891-1897.	2.0	111
129	Parental Hope for Children With Advanced Cancer. <i>Pediatrics</i> , 2015, 135, 868-874.	1.0	108
130	Symptom Prevalence in Lung and Colorectal Cancer Patients. <i>Journal of Pain and Symptom Management</i> , 2015, 49, 192-202.	0.6	82
131	Young Adult Perspectives on a Successful Transition from Pediatric to Adult Care in Sickle Cell Disease. <i>Journal of Hematology Research</i> , 2015, 2, 17-24.	0.5	37
132	Communicating About Prognosis: Ethical Responsibilities of Pediatricians and Parents. <i>Pediatrics</i> , 2014, 133, S24-S30.	1.0	103
133	Underuse of Hospice Care by Medicaid-Insured Patients With Stage IV Lung Cancer in New York and California. <i>Journal of Clinical Oncology</i> , 2013, 31, 2569-2579.	0.8	60
134	Parent perceptions of the quality of information received about a child's cancer. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1896-1901.	0.8	49
135	Patients' Expectations about Effects of Chemotherapy for Advanced Cancer. <i>New England Journal of Medicine</i> , 2012, 367, 1616-1625.	13.9	990
136	Reasons Why Physicians Do Not Have Discussions About Poor Prognosis, Why It Matters, and What Can Be Improved. <i>Journal of Clinical Oncology</i> , 2012, 30, 2715-2717.	0.8	256
137	Associations Between End-of-Life Discussion Characteristics and Care Received Near Death: A Prospective Cohort Study. <i>Journal of Clinical Oncology</i> , 2012, 30, 4387-4395.	0.8	457
138	End-of-Life Care Discussions Among Patients With Advanced Cancer. <i>Annals of Internal Medicine</i> , 2012, 156, 204.	2.0	341
139	Communication and Decision Support for Children with Advanced Cancer and Their Families. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2012, , 637-643.	1.8	10
140	Parents' Roles in Decision Making for Children With Cancer in the First Year of Cancer Treatment. <i>Journal of Clinical Oncology</i> , 2011, 29, 2085-2090.	0.8	67
141	End-of-Life Discussions, Goal Attainment, and Distress at the End of Life: Predictors and Outcomes of Receipt of Care Consistent With Preferences. <i>Journal of Clinical Oncology</i> , 2010, 28, 1203-1208.	0.8	685
142	Racial Disparities in the Outcomes of Communication on Medical Care Received Near Death. <i>Archives of Internal Medicine</i> , 2010, 170, 1533-40.	4.3	235
143	Peace of Mind and Sense of Purpose as Core Existential Issues Among Parents of Children With Cancer. <i>JAMA Pediatrics</i> , 2009, 163, 519.	3.6	88
144	Measuring therapeutic alliance between oncologists and patients with advanced cancer. <i>Cancer</i> , 2009, 115, 3302-3311.	2.0	156

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145	Peace, Equanimity, and Acceptance in the cancer experience (PEACE). <i>Cancer</i> , 2008, 112, 2509-2517.	2.0	119
146	Associations Between End-of-Life Discussions, Patient Mental Health, Medical Care Near Death, and Caregiver Bereavement Adjustment. <i>JAMA - Journal of the American Medical Association</i> , 2008, 300, 1665.	3.8	2,213
147	Parents' Views of Cancer-Directed Therapy for Children With No Realistic Chance for Cure. <i>Journal of Clinical Oncology</i> , 2008, 26, 4759-4764.	0.8	95
148	Hope and Prognostic Disclosure. <i>Journal of Clinical Oncology</i> , 2007, 25, 5636-5642.	0.8	320
149	Understanding of Prognosis Among Parents of Children With Cancer: Parental Optimism and the Parent-Physician Interaction. <i>Journal of Clinical Oncology</i> , 2007, 25, 1357-1362.	0.8	163
150	Quality of Health Care for Children. <i>JAMA Pediatrics</i> , 2007, 161, 828.	3.6	13
151	Communication About Prognosis Between Parents and Physicians of Children With Cancer: Parent Preferences and the Impact of Prognostic Information. <i>Journal of Clinical Oncology</i> , 2006, 24, 5265-5270.	0.8	289
152	Parent and Physician Perspectives on Quality of Care at the End of Life in Children With Cancer. <i>Journal of Clinical Oncology</i> , 2005, 23, 9155-9161.	0.8	296
153	The Day One Talk. <i>Journal of Clinical Oncology</i> , 2004, 22, 563-566.	0.8	95
154	The Art of Saying Nothing. <i>Pediatrics</i> , 0, , .	1.0	1
155	Factors Affecting Hospice Use Among Adolescents and Young Adult Cancer Patients. <i>Journal of Adolescent and Young Adult Oncology</i> , 0, , .	0.7	2