

# Justin N Baker

## List of Publications by Year in descending order

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Version: 2024-02-01

207  
papers

6,090  
citations

71102

41  
h-index

110387

64  
g-index

211  
all docs

211  
docs citations

211  
times ranked

4826  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-Wide Analyses Identify Recurrent Amplifications of Receptor Tyrosine Kinases and Cell-Cycle Regulatory Genes in Diffuse Intrinsic Pontine Glioma. <i>Journal of Clinical Oncology</i> , 2011, 29, 3999-4006.	1.6	286
2	“Trying to Be a Good Parent” As Defined By Interviews With Parents Who Made Phase I, Terminal Care, and Resuscitation Decisions for Their Children. <i>Journal of Clinical Oncology</i> , 2009, 27, 5979-5985.	1.6	236
3	Patients’ and Parents’ Needs, Attitudes, and Perceptions About Early Palliative Care Integration in Pediatric Oncology. <i>JAMA Oncology</i> , 2017, 3, 1214.	7.1	146
4	The COVID-19 pandemic: A rapid global response for children with cancer from SIOP, COG, SIOP-E, SIOP-PODC, IPSO, PROS, CCI, and St Jude Global. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28409.	1.5	113
5	Key Factors Affecting Dying Children and Their Families. <i>Journal of Palliative Medicine</i> , 2005, 8, s-70-s-78.	1.1	111
6	Phase I Study of Vandetanib During and After Radiotherapy in Children With Diffuse Intrinsic Pontine Glioma. <i>Journal of Clinical Oncology</i> , 2010, 28, 4762-4768.	1.6	108
7	Early Integration of Palliative Care for Children with High-Risk Cancer and Their Families. <i>Pediatric Blood and Cancer</i> , 2016, 63, 593-597.	1.5	106
8	NCCN Guidelines Insights: Palliative Care, Version 2.2017. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2017, 15, 989-997.	4.9	98
9	Research Priorities in Pediatric Palliative Care. <i>Journal of Pediatrics</i> , 2015, 167, 467-470.e3.	1.8	94
10	Palliative Care Involvement Is Associated with Less Intensive End-of-Life Care in Adolescent and Young Adult Oncology Patients. <i>Journal of Palliative Medicine</i> , 2017, 20, 509-516.	1.1	94
11	Decision Making by Parents of Children With Incurable Cancer Who Opt for Enrollment on a Phase I Trial Compared With Choosing a Do Not Resuscitate/Terminal Care Option. <i>Journal of Clinical Oncology</i> , 2010, 28, 3292-3298.	1.6	93
12	Pediatric palliative care in the community. <i>Ca-A Cancer Journal for Clinicians</i> , 2015, 65, 315-333.	329.8	90
13	National Survey of Pediatric Residency Program Directors and Residents Regarding Education in Palliative Medicine and End-of-Life Care. <i>Journal of Palliative Medicine</i> , 2007, 10, 420-429.	1.1	87
14	Integration of Palliative Care Practices into the Ongoing Care of Children with Cancer: Individualized Care Planning and Coordination. <i>Pediatric Clinics of North America</i> , 2008, 55, 223-250.	1.8	87
15	Adolescents’ preferences for treatment decisional involvement during their cancer. <i>Cancer</i> , 2015, 121, 4416-4424.	4.1	87
16	Phase I Trial, Pharmacokinetics, and Pharmacodynamics of Vandetanib and Dasatinib in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma. <i>Clinical Cancer Research</i> , 2013, 19, 3050-3058.	7.0	82
17	Patterns of End-of-Life Care in Children With Advanced Solid Tumor Malignancies Enrolled on a Palliative Care Service. <i>Journal of Pain and Symptom Management</i> , 2015, 50, 305-312.	1.2	78
18	Providing Research Results to Participants: Attitudes and Needs of Adolescents and Parents of Children With Cancer. <i>Journal of Clinical Oncology</i> , 2009, 27, 878-883.	1.6	76

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19	Helping parents live with the hole in their heart: The role of health care providers and institutions in the bereaved parents' grief journeys. <i>Cancer</i> , 2016, 122, 2757-2765.	4.1	75
20	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.	6.2	73
21	Communicating and Understanding the Purpose of Pediatric Phase I Cancer Trials. <i>Journal of Clinical Oncology</i> , 2012, 30, 4367-4372.	1.6	72
22	Prospective collection of tissue samples at autopsy in children with diffuse intrinsic pontine glioma. <i>Cancer</i> , 2010, 116, 4632-4637.	4.1	70
23	Child and adolescent self-report symptom measurement in pediatric oncology research: a systematic literature review. <i>Quality of Life Research</i> , 2018, 27, 291-319.	3.1	67
24	Adolescent perspectives on phase I cancer research. <i>Pediatric Blood and Cancer</i> , 2013, 60, 873-878.	1.5	63
25	Suggestions from adolescents, young adults, and parents for improving informed consent in phase 1 pediatric oncology trials. <i>Cancer</i> , 2013, 119, 4154-4161.	4.1	59
26	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.	6.3	59
27	Use of epidural and peripheral nerve blocks at the end of life in children and young adults with cancer: the collaboration between a pain service and a palliative care service. <i>Paediatric Anaesthesia</i> , 2010, 20, 1070-1077.	1.1	58
28	Pediatric Palliative Sedation Therapy with Propofol: Recommendations Based on Experience in Children with Terminal Cancer. <i>Journal of Palliative Medicine</i> , 2012, 15, 1082-1090.	1.1	58
29	Integrating next-generation sequencing into pediatric oncology practice: An assessment of physician confidence and understanding of clinical genomics. <i>Cancer</i> , 2017, 123, 2352-2359.	4.1	58
30	Ethics Consultation in Pediatrics: Long-Term Experience From a Pediatric Oncology Center. <i>American Journal of Bioethics</i> , 2015, 15, 3-17.	0.9	57
31	Palliative Care Version 1.2016. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2016, 14, 82-113.	4.9	57
32	Empowering Bereaved Parents Through the Development of a Comprehensive Bereavement Program. <i>Journal of Pain and Symptom Management</i> , 2017, 53, 767-775.	1.2	57
33	Decision-making by Adolescents and Parents of Children With Cancer Regarding Health Research Participation. <i>Pediatrics</i> , 2009, 124, 959-965.	2.1	56
34	Reported Availability and Gaps of Pediatric Palliative Care in Low- and Middle-Income Countries: A Systematic Review of Published Data. <i>Journal of Palliative Medicine</i> , 2014, 17, 1369-1383.	1.1	55
35	Illness and end-of-life experiences of children with cancer who receive palliative care. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26895.	1.5	53
36	Availability of palliative care services for children with cancer in economically diverse regions of the world. <i>European Journal of Cancer</i> , 2010, 46, 2260-2266.	2.8	52

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37	Communication in Pediatric Oncology: A Qualitative Study. <i>Pediatrics</i> , 2020, 146, .	2.1	52
38	Eliciting the child's voice in adverse event reporting in oncology trials: Cognitive interview findings from the Pediatric Patientâ€Reported Outcomes version of the Common Terminology Criteria for Adverse Events initiative. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26261.	1.5	50
39	Palliative Care, Version 1.2014. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2014, 12, 1379-1388.	4.9	48
40	Parental Grief Following the Death of a Child from Cancer: The Ongoing Odyssey. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1594-1602.	1.5	48
41	Strange Bedfellows No More: How Integrated Stem-Cell Transplantation and Palliative Care Programs Can Together Improve End-of-Life Care. <i>Journal of Oncology Practice</i> , 2017, 13, 569-577.	2.5	43
42	Provision of Palliative and Hospice Care to Children in the Community: A Population Study of Hospice Nurses. <i>Journal of Pain and Symptom Management</i> , 2019, 57, 241-250.	1.2	43
43	Congruence Gaps Between Adolescents With Cancer and Their Families Regarding Values, Goals, and Beliefs About End-of-Life Care. <i>JAMA Network Open</i> , 2020, 3, e205424.	5.9	41
44	NCCN Guidelines® Insights: Palliative Care, Version 2.2021. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2021, 19, 780-788.	4.9	41
45	A Process to Facilitate Decision Making in Pediatric Stem Cell Transplantation: The Individualized Care Planning and Coordination Model. <i>Biology of Blood and Marrow Transplantation</i> , 2007, 13, 245-254.	2.0	40
46	Parental Perspectives of Communication at the End of Life at a Pediatric Oncology Institution. <i>Journal of Palliative Medicine</i> , 2016, 19, 326-332.	1.1	40
47	Communicating Effectively in Pediatric Cancer Care: Translating Evidence into Practice. <i>Children</i> , 2018, 5, 40.	1.5	39
48	â€Please, I want to go homeâ€: ethical issues raised when considering choice of place of care in palliative care. <i>Postgraduate Medical Journal</i> , 2007, 83, 643-648.	1.8	38
49	Factors That Distinguish Symptoms of Most Concern to Parents from Other Symptoms of Dying Children. <i>Journal of Pain and Symptom Management</i> , 2010, 39, 627-636.	1.2	38
50	Informed consent for pediatric phase 1 cancer trials: Physicians' perspectives. <i>Cancer</i> , 2010, 116, 3244-3250.	4.1	37
51	Pediatric Palliative Oncology: Bridging Silos of Care Through an Embedded Model. <i>Journal of Clinical Oncology</i> , 2017, 35, 2740-2744.	1.6	37
52	Pediatric palliative oncology: the state of the science and art of caring for children with cancer. <i>Current Opinion in Pediatrics</i> , 2018, 30, 40-48.	2.0	37
53	Regret and unfinished business in parents bereaved by cancer: A mixed methods study. <i>Palliative Medicine</i> , 2020, 34, 367-377.	3.1	37
54	Best practices for pediatric palliative cancer care: a primer for clinical providers. <i>The Journal of Supportive Oncology</i> , 2013, 11, 114-125.	2.3	37

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55	The first step to integrating the child's voice in adverse event reporting in oncology trials: A content validation study among pediatric oncology clinicians. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1231-1236.	1.5	36
56	Predictors of Late Palliative Care Referral in Children With Cancer. <i>Journal of Pain and Symptom Management</i> , 2018, 55, 1550-1556.	1.2	36
57	“What if?” Addressing uncertainty with families. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27699.	1.5	36
58	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.	1.5	36
59	Integrating Palliative Care in Pediatric Oncology: Evidence for an Evolving Paradigm for Comprehensive Cancer Care. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2016, 14, 741-748.	4.9	35
60	Cognitive Interview-Based Validation of the Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events in Adolescents with Cancer. <i>Journal of Pain and Symptom Management</i> , 2017, 53, 759-766.	1.2	35
61	Models of Pediatric Palliative Oncology Outpatient Care—Benefits, Challenges, and Opportunities. <i>Journal of Oncology Practice</i> , 2019, 15, 476-487.	2.5	35
62	Practical communication guidance to improve phase 1 informed consent conversations and decision-making in pediatric oncology. <i>Cancer</i> , 2015, 121, 2439-2448.	4.1	34
63	Does phase 1 trial enrollment preclude quality end-of-life care? Phase 1 trial enrollment and end-of-life care characteristics in children with cancer. <i>Cancer</i> , 2015, 121, 1508-1512.	4.1	34
64	Unrealistic parental expectations for cure in poor-prognosis childhood cancer. <i>Cancer</i> , 2020, 126, 416-424.	4.1	34
65	End-of-Life Care for Hospitalized Children. <i>Pediatric Clinics of North America</i> , 2014, 61, 835-854.	1.8	32
66	Concept elicitation phase for the development of the pediatric patient-reported outcome version of the Common Terminology Criteria for Adverse Events. <i>Cancer</i> , 2016, 122, 141-148.	4.1	32
67	Methadone prolongs cardiac conduction in young patients with cancer-related pain. <i>Journal of Opioid Management</i> , 2016, 12, 131-138.	0.5	32
68	Race Does Not Influence Do-Not-Resuscitate Status or the Number or Timing of End-of-Life Care Discussions at a Pediatric Oncology Referral Center. <i>Journal of Palliative Medicine</i> , 2009, 12, 71-76.	1.1	31
69	Recommendations for the Return of Research Results to Study Participants and Guardians: A Report From the Children's Oncology Group. <i>Journal of Clinical Oncology</i> , 2012, 30, 4573-4579.	1.6	31
70	Association of Religious and Spiritual Factors With Patient-Reported Outcomes of Anxiety, Depressive Symptoms, Fatigue, and Pain Interference Among Adolescents and Young Adults With Cancer. <i>JAMA Network Open</i> , 2020, 3, e206696.	5.9	30
71	Patient Involvement in Informed Consent for Pediatric Phase I Cancer Research. <i>Journal of Pediatric Hematology/Oncology</i> , 2014, 36, 635-640.	0.6	29
72	Speaking genomics to parents offered germline testing for cancer predisposition: Use of a 2-visit consent model. <i>Cancer</i> , 2019, 125, 2455-2464.	4.1	29

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73	Are we meeting the informational needs of cancer patients and families? Perception of physician communication in pediatric oncology. <i>Cancer</i> , 2019, 125, 1518-1526.	4.1	28
74	The Impact of Specialty Palliative Care in Pediatric Oncology: A Systematic Review. <i>Journal of Pain and Symptom Management</i> , 2021, 61, 1060-1079.e2.	1.2	28
75	Methylnaltrexone for Opioid-Induced Constipation in Children and Adolescents and Young Adults with Progressive Incurable Cancer at the End of Life. <i>Journal of Palliative Medicine</i> , 2015, 18, 631-633.	1.1	27
76	Predictors of Location of Death for Children with Cancer Enrolled on a Palliative Care Service. <i>Oncologist</i> , 2018, 23, 1525-1532.	3.7	27
77	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.	1.2	27
78	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.	1.6	27
79	Treatment intensity and symptom burden in hospitalized adolescent and young adult hematopoietic cell transplant recipients at the end of life. <i>Bone Marrow Transplantation</i> , 2018, 53, 84-90.	2.4	26
80	Impact of Race and Ethnicity on End-of-Life Experiences for Children With Cancer. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 767-774.	1.4	26
81	Bereaved Parents' Intentions and Suggestions about Research Autopsies in Children with Lethal Brain Tumors. <i>Journal of Pediatrics</i> , 2013, 163, 581-586.	1.8	25
82	Occult post-contrast signal enhancement in pediatric diffuse intrinsic pontine glioma is the MRI marker of angiogenesis?. <i>Neuroradiology</i> , 2014, 56, 405-412.	2.2	25
83	Changes in Medical Care at a Pediatric Oncology Referral Center after Placement of a Do-Not-Resuscitate Order. <i>Journal of Palliative Medicine</i> , 2010, 13, 1349-1352.	1.1	24
84	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.	1.8	24
85	Grief and Bereavement in Fathers After the Death of a Child: A Systematic Review. <i>Pediatrics</i> , 2021, 147, .	2.1	24
86	Keeping Users at the Center: Developing a Multimedia Interface for Informed Consent. <i>Technical Communication Quarterly</i> , 2008, 17, 335-357.	1.6	23
87	Patient-controlled analgesia at the end of life at a pediatric oncology institution. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1237-1244.	1.5	23
88	Communication about the risks and benefits of phase I pediatric oncology trials. <i>Contemporary Clinical Trials</i> , 2015, 41, 139-145.	1.8	23
89	Children with minimal chance for cure: parent proxy of the child's health-related quality of life and the effect on parental physical and mental health during treatment. <i>Journal of Neuro-Oncology</i> , 2016, 129, 373-381.	2.9	23
90	Ethical considerations surrounding germline next-generation sequencing of children with cancer. <i>Expert Review of Molecular Diagnostics</i> , 2017, 17, 523-534.	3.1	23

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91	Pediatric Oncology: Managing Pain at the End of Life. Paediatric Drugs, 2016, 18, 161-180.	3.1	22
92	Ethical issues in the care of adolescent and young adult oncology patients. Pediatric Blood and Cancer, 2019, 66, e27608.	1.5	22
93	Reconsidering early parental grief following the death of a child from cancer: a new framework for future research and bereavement support. Supportive Care in Cancer, 2020, 28, 4131-4139.	2.2	22
94	Barriers to the early integration of palliative care in pediatric oncology in 11 Eurasian countries. Cancer, 2020, 126, 4984-4993.	4.1	22
95	Parent-Clinician Communication Intervention during End-of-Life Decision Making for Children with Incurable Cancer. Journal of Palliative Medicine, 2012, 15, 916-922.	1.1	21
96	Management of diffuse intrinsic pontine glioma in children: current and future strategies for improving prognosis. CNS Oncology, 2014, 3, 421-431.	3.0	21
97	Going straight to the source: A pilot study of bereaved parentâ€facilitated communication training for pediatric subspecialty fellows. Pediatric Blood and Cancer, 2017, 64, 156-162.	1.5	21
98	Bereaved Parents, Hope, and Realism. Pediatrics, 2020, 145, e20192771.	2.1	21
99	Evaluation of Physician and Nurse Dyad Training Procedures to Deliver a Palliative and End-of-Life Communication Intervention to Parents of Children with a Brain Tumor. Journal of Pediatric Oncology Nursing, 2015, 32, 337-347.	1.5	20
100	Palliative Sedation With Propofol for an Adolescent With a DNR Order. Pediatrics, 2017, 140, .	2.1	20
101	Prognostic Communication Between Oncologists and Parents of Children With Advanced Cancer. Pediatrics, 2021, 147, .	2.1	20
102	Parental views on communication between children and clinicians in pediatric oncology: a qualitative study. Supportive Care in Cancer, 2021, 29, 4957-4968.	2.2	19
103	Multilevel barriers to communication in pediatric oncology: Clinicians' perspectives. Cancer, 2021, 127, 2130-2138.	4.1	19
104	A Model of Interpersonal Trust, Credibility, and Relationship Maintenance. Pediatrics, 2019, 144, e20191319.	2.1	17
105	Facilitators Associated With Building and Sustaining Therapeutic Alliance in Advanced Pediatric Cancer. JAMA Network Open, 2021, 4, e2120925.	5.9	17
106	Creating a Palliative and End-of-Life Program in a Cure-Oriented Pediatric Setting: The Zig-Zag Method. Journal of Pediatric Oncology Nursing, 2007, 24, 246-254.	1.5	16
107	â€œBeing a good patientâ€•during times of illness as defined by adolescent patients with cancer. Cancer, 2016, 122, 2224-2233.	4.1	16
108	Reducing Compassion Fatigue in Inpatient Pediatric Oncology Nurses. , 2019, 46, 338-347.		16



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109	Bad News Deserves Better Communication: A Customizable Curriculum for Teaching Learners to Share Life-Altering Information in Pediatrics. MedEdPORTAL: the Journal of Teaching and Learning Resources, 2016, 12, 10438.	1.2	16
110	Natural Language Processing and Machine Learning Methods to Characterize Unstructured Patient-Reported Outcomes: Validation Study. Journal of Medical Internet Research, 2021, 23, e26777.	4.3	16
111	Productivity in Pediatric Palliative Care: Measuring and Monitoring an Elusive Metric. Journal of Pain and Symptom Management, 2017, 53, 952-961.	1.2	15
112	Outcome Measurement for Children and Young People. Journal of Palliative Medicine, 2017, 20, 313-313.	1.1	15
113	Importance ratings on patient-reported outcome items for survivorship care: comparison between pediatric cancer survivors, parents, and clinicians. Quality of Life Research, 2018, 27, 1877-1884.	3.1	15
114	Parent-child communication surrounding genetic testing for Li-Fraumeni syndrome: Living under the cloud of cancer. Pediatric Blood and Cancer, 2018, 65, e27350.	1.5	15
115	Validation of the caregiver Pediatric Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events measure. Cancer, 2021, 127, 1483-1494.	4.1	15
116	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.	1.2	15
117	Patients, caregivers, and clinicians differ in performance status ratings: Implications for pediatric cancer clinical trials. Cancer, 2021, 127, 3664-3670.	4.1	15
118	Development of depression in survivors of childhood and adolescent cancer: a multi-level life course conceptual framework. Supportive Care in Cancer, 2017, 25, 2009-2017.	2.2	14
119	Mapping child and adolescent self-reported symptom data to clinician-reported adverse event grading to improve pediatric oncology care and research. Cancer, 2020, 126, 140-147.	4.1	14
120	Hospice nurses request paediatric-specific educational resources and training programs to improve care for children and families in the community: Qualitative data analysis from a population-level survey. Palliative Medicine, 2020, 34, 403-412.	3.1	14
121	The state of the science for communication training in pediatric oncology: A systematic review. Pediatric Blood and Cancer, 2020, 67, e28607.	1.5	14
122	Clinician Emotions Surrounding Pediatric Oncology Patient Deterioration. Frontiers in Oncology, 2021, 11, 626457.	2.8	14
123	Impact of PEWS on Perceived Quality of Care During Deterioration in Children With Cancer Hospitalized in Different Resource-Settings. Frontiers in Oncology, 2021, 11, 660051.	2.8	14
124	Grief and Bereavement in Parents After the Death of a Child in Low- and Middle-Income Countries. Children, 2020, 7, 39.	1.5	14
125	Pediatric Palliative Oncology: A New Training Model for an Emerging Field. Journal of Clinical Oncology, 2016, 34, 288-289.	1.6	13
126	Communication with Children with Cancer and Their Families Throughout the Illness Journey and at the End of Life. Pediatric Oncology, 2018, , 55-93.	0.5	13



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127	A pediatric brain tumor consortium phase II trial of capecitabine rapidly disintegrating tablets with concomitant radiation therapy in children with newly diagnosed diffuse intrinsic pontine gliomas. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26832.	1.5	13
128	Incorporating Bereaved Parents as Faculty Facilitators and Educators in Teaching Principles of Palliative and End-of-Life Care. <i>American Journal of Hospice and Palliative Medicine</i> , 2018, 35, 1518-1525.	1.4	13
129	“Living life as if I never had cancer”: A study of the meaning of living well in adolescents and young adults who have experienced cancer. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28599.	1.5	13
130	Thoughts from the threshold: patient and family hopes, fears, values, and goals at the onset of pediatric hematopoietic cell transplantation. <i>Bone Marrow Transplantation</i> , 2020, 55, 1103-1113.	2.4	13
131	Pediatric Advance Care Planning and Families’ Positive Caregiving Appraisals: An RCT. <i>Pediatrics</i> , 2021, 147, .	2.1	13
132	Interdependent functions of communication with adolescents and young adults in oncology. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29588.	1.5	13
133	Physician Perceptions of Palliative Care for Children With Cancer in Latin America. <i>JAMA Network Open</i> , 2022, 5, e221245.	5.9	13
134	A multicountry assessment in Eurasia: Alignment of physician perspectives on palliative care integration in pediatric oncology with World Health Organization guidelines. <i>Cancer</i> , 2020, 126, 3777-3787.	4.1	12
135	Assume It Will Break: Parental Perspectives on Negative Communication Experiences in Pediatric Oncology. <i>JCO Oncology Practice</i> , 2021, 17, e859-e871.	2.9	12
136	Racial and Ethnic Disparities in Health Outcomes Among Long-Term Survivors of Childhood Cancer: A Scoping Review. <i>Frontiers in Public Health</i> , 2021, 9, 741334.	2.7	12
137	The role of the pediatric anesthesiologist in relieving suffering at the end of life: when is palliative sedation appropriate in pediatrics?. <i>Paediatric Anaesthesia</i> , 2017, 27, 443-444.	1.1	11
138	The bereavement experience of adolescents and early young adults with cancer: Peer and parental loss due to death is associated with increased risk of adverse psychological outcomes. <i>PLoS ONE</i> , 2017, 12, e0181024.	2.5	11
139	Role of Amputation in Improving Mobility, Pain Outcomes, and Emotional and Psychological Well-Being in Children With Metastatic Osteosarcoma. <i>American Journal of Hospice and Palliative Medicine</i> , 2019, 36, 105-110.	1.4	11
140	The association of age, literacy, and race on completing patient-reported outcome measures in pediatric oncology. <i>Quality of Life Research</i> , 2019, 28, 1793-1801.	3.1	11
141	Engaging Parents of Children Who Died From Cancer in Research on the Early Grief Experience. <i>Journal of Pain and Symptom Management</i> , 2021, 61, 781-788.	1.2	11
142	Hope-Colored Glasses: Perceptions of Prognosis Among Pediatric Oncology Patients and Their Parents. <i>JCO Oncology Practice</i> , 2021, 17, e730-e739.	2.9	11
143	The intersectionality of gender and poverty on symptom suffering among adolescents with cancer. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29144.	1.5	11
144	Profile Comparison of Patient-Reported and Proxy-Reported Symptoms in Pediatric Patients With Cancer Receiving Chemotherapy. <i>JAMA Network Open</i> , 2022, 5, e221855.	5.9	11

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145	Bereaved Parents' Perceptions About When Their Child's Cancer-Related Death Would Occur. <i>Journal of Pain and Symptom Management</i> , 2009, 38, 561-567.	1.2	10
146	The Impact of Pediatric Palliative Care Involvement in the Care of Critically Ill Patients without Complex Chronic Conditions. <i>Journal of Palliative Medicine</i> , 2019, 22, 553-556.	1.1	10
147	MyPref: pilot study of a novel communication and decision-making tool for adolescents and young adults with advanced cancer. <i>Supportive Care in Cancer</i> , 2021, 29, 2983-2992.	2.2	10
148	Multidisciplinary Clinician Perspectives on Embedded Palliative Care Models in Pediatric Cancer. <i>Journal of Pain and Symptom Management</i> , 2022, 64, 222-233.	1.2	10
149	Methylnaltrexone Use in a Seventeen-Month-Old Female with Progressive Cancer and Rectal Prolapse. <i>Journal of Palliative Medicine</i> , 2013, 16, 1486-1488.	1.1	9
150	Microethics of Communication—Hidden Roles of Bias and Heuristics in the Words We Choose. <i>JAMA Pediatrics</i> , 2018, 172, 1115.	6.2	9
151	Status of Palliative Oncology Care for Children and Young People in Sub-Saharan Africa: A Perspective Paper on Priorities for New Frontiers. <i>JCO Global Oncology</i> , 2021, 7, 1395-1405.	1.8	9
152	Global Experiences of Pediatric Palliative Care Teams During the First 6 Months of the SARS-CoV-2 Pandemic. <i>Journal of Pain and Symptom Management</i> , 2021, 62, e91-e99.	1.2	9
153	Community Hospice Nurses' Perspectives on Needs, Preferences, and Challenges Related to Caring for Children With Serious Illness. <i>JAMA Network Open</i> , 2021, 4, e2127457.	5.9	9
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