



Citation lists are generated by [PedPalASCNET](#) through monthly searches. Visit our [blog](#) to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click [here](#).

Trends in Pediatric Palliative Care Citation List

2018; Issue #4

Arnolds, M., Xu, L., Hughes, P., McCoy, J., & Meadow, W. (2018). [Worth a Try? Describing the Experiences of Families during the Course of Care in the Neonatal Intensive Care Unit When the Prognosis is Poor.](#) *J Pediatr.*

Balkin, E. M., Sleeper, L. A., Kirkpatrick, J. N., Swetz, K. M., Coggins, M. K., Wolfe, J., & Blume, E. D. (2018). [Physician Perspectives on Palliative Care for Children with Advanced Heart Disease: A Comparison between Pediatric Cardiology and Palliative Care Physicians.](#) *J Palliat Med.*

Butler, A. E., Hall, H., & Copnell, B. (2018). [Becoming a Team: The Nature of the Parent-Healthcare Provider Relationship when a Child is Dying in the Pediatric Intensive Care Unit.](#) *J Pediatr Nurs.*

Campbell, J. D., Whittington, M. D., Kim, C. H., VanderVeen, G. R., Knupp, K. G., & Gammaitoni, A. (2018). [Assessing the impact of caring for a child with Dravet syndrome: Results of a caregiver survey.](#) *Epilepsy Behav, 80,* 152–156.

Ciriello, A. G., Dizon, Z. B., & October, T. W. (n.d.). [Speaking a Different Language: A Qualitative Analysis Comparing Language of Palliative Care and Pediatric Intensive Care Unit Physicians.](#) *American Journal of Hospice & Palliative Medicine.* Retrieved from <http://journals.sagepub.com/doi/full/10.1177/1049909117700101>

Cunningham, M., Williamson, S., Baker, J., Pennington, K., Young, A., Bolick, R., & Norton, C. (2018). [Partnering with Payers to Meet the Needs of Pediatric Palliative Care Patients \(FR473\).](#) *Journal of Pain and Symptom Management, 55*(2), 627–628.

Curcio, D. L. (2017). [The Lived Experiences of Nurses Caring For Dying Pediatric Patients.](#) *Pediatr Nurs, 43*(1), 8–14.

Grunauer, M., & Mikesell, C. (2018). [A Review of the Integrated Model of Care: An Opportunity to Respond to Extensive Palliative Care Needs in Pediatric Intensive Care Units in Under-Resourced Settings.](#) *Front Pediatr, 6,* 3.

Hasan, F., Weingarten, K., Rapoport, A., Bouffet, E., & Bartels, U. (2018). [End-of-life care of children with diffuse intrinsic pontine glioma.](#) *J Neurooncol.*



Citation lists are generated by [PedPalASCNET](#) through monthly searches. Visit our [blog](#) to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click [here](#).

Haug, S., Farooqi, S., Wilson, C. G., Hopper, A., Oei, G., & Carter, B. (2018). [Survey on Neonatal End-of-Life Comfort Care Guidelines Across America](#). *Journal of Pain and Symptom Management*.

Henderson, C. M., Wilfond, B. S., & Boss, R. D. (2018). [Bringing social context into the conversation about pediatric long-term ventilation](#). *Hospital Pediatrics*, 8(2), 102–108.

Hill, D., Faerber, J., Carroll, K., Miller, V., Morrison, W., Hinds, P., & Feudtner, C. (2018). [Changes Over Time in Good-Parent Beliefs Among Parents of Children with Serious Illness: Two Year Cohort Study \(TH320B\)](#). *Journal of Pain and Symptom Management*, 55(2), 570.

Hill, D. L., Nathanson, P. G., Fenderson, R. M., Carroll, K. W., & Feudtner, C. (2017). [Parental Concordance Regarding Problems and Hopes for Seriously Ill Children: A Two-Year Cohort Study](#). *J Pain Symptom Manage*, 53(5), 911–918.

Hiscock, A., Kuhn, I., & Barclay, S. (2017). [Advance care discussions with young people affected by life-limiting neuromuscular diseases: A systematic literature review and narrative synthesis](#). *Neuromuscular Disorders*, 27(2), 115–119.

Kang, T., Lotstein, D., Humphrey, L., Klick, J., & Williams, C. (2018). [We Built It...They All Came...Now How to Keep from Drowning? Pediatric Palliative Care Program Development 202: Skills in Your Toolbox for Growth and Sustainability \(P20\)](#). *Journal of Pain and Symptom Management*, 55(2), 556–557.

Kase, S. M., Waldman, E. D., & Weintraub, A. S. (2018). [A cross-sectional pilot study of compassion fatigue, burnout, and compassion satisfaction in pediatric palliative care providers in the United States](#). *Palliat Support Care*, 1–7.

Lamba, S., Sudore, R., Buckholz, G. T., Radwany, S. M., & Wolfe, J. (2018). [It Takes a Village: Building a Culture of Mentorship and Sponsorship to Thrive Throughout Our Careers \(FR409\)](#). *Journal of Pain and Symptom Management*, 55(2), 598.

Meaney, S., Corcoran, P., & O'Donoghue, K. (2017). [Death of One Twin during the Perinatal Period: An Interpretative Phenomenological Analysis](#). *J Palliat Med*, 20(3), 290–293.

Mooney-Doyle, K., Deatrick, J. A., Ulrich, C. M., Meghani, S. H., & Feudtner, C. (2018). [Parenting in Childhood Life-Threatening Illness: A Mixed-Methods Study](#). *Journal of Palliative Medicine*, 21(2), 208–215.



Citation lists are generated by [PedPalASCNET](#) through monthly searches. Visit our [blog](#) to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click [here](#).

Nelson, K., Rosella, L., Mahant, S., Guttmann, A., & Cohen, E. (2018). [Effect of Permanent Feeding Tube Placement on Healthcare Utilization Among Children with Neurologic Impairment \(TH340A\)](#). *Journal of Pain and Symptom Management*, 55(2), 578–579.

Noritz, G., & Feudtner, C. (2018). [Walking the Line Between Hope And Hype: Palliative Care in the Era of New and Expensive Neuromuscular Therapies for Children and Adults \(TH317\)](#). *Journal of Pain and Symptom Management*, 55(2), 567–568.

Parravicini, E., Daho, M., Foe, G., Steinwurtzel, R., & Byrne, M. (2018). [Parental assessment of comfort in newborns affected by lifelimiting conditions treated by a standardized neonatal comfort care program](#). *Journal of Perinatology*, 38(2), 142–147.

Pentaris, P., Papadatou, D., Jones, A., & Hosang, G. M. (2018). [Palliative care professional's perceptions of barriers and challenges to accessing children's hospice and palliative care services in south east London: A preliminary study](#). *Death Stud.*

Perluxo, D., & Francisco, R. (2018). [Use of Facebook in the maternal grief process: An exploratory qualitative study](#). *Death Studies*, 42(2), 79–88.

Rossfeld, Z. M., Tumin, D., & Humphrey, L. M. (2018). [Self-Assessment of Skills and Competencies among Residents Participating in a Pediatric Hospice and Palliative Medicine Elective Rotation](#). *Journal of Palliative Medicine*, 21, 2.

Snaman, J., Kaye, E., Levine, D., Clark, L., Wilcox, R., Cunningham, M., & Baker, J. (2018). [Incorporating Bereaved Parents as Facilitators and Educators in Teaching Principles of Palliative and End-of-Life Care \(FR452\)](#). *Journal of Pain and Symptom Management*, 55(2), 619.

Snaman, J. M., Kaye, E., Cunningham, M. J., Sykes, A., Levine, D. R., Mahoney, D., & Baker, J. N. (2018). [Going Straight to the Source: A Pilot Study of Bereaved Parent-facilitated Communication Training for Pediatric Subspecialty Fellows](#). *Pediatrics*, 141(1), 381–381.

Stutz, M., Kao, R. L., Huard, L., Grotts, J., Sanz, J., & Ross, M. K. (2018). [Associations Between Pediatric Palliative Care Consultation and End-of-Life Preparation at an Academic Medical Center: A Retrospective EHR Analysis](#). *Hosp Pediatr.*

Ullrich, C., Levine, D., Li, A., Ma, C., Wolfe, J., & Holder, R. (2018). [Family Satisfaction with Symptom Management Practices and Child Comfort During Pediatric Hematopoietic Stem Cell Transplant \(TH320C\)](#). *Journal of Pain and Symptom Management*, 55(2), 570–571.



Citation lists are generated by [PedPalASCNET](#) through monthly searches. Visit our [blog](#) to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click [here](#).

Verberne, L. M., Kars, M. C., Schepers, S. A., Schouten-Van Meeteren, A. Y. N., Grootenhuis, M. A., & Van Delden, J. J. M. (2018). [Barriers and facilitators to the implementation of a paediatric palliative care team.](#) *BMC Palliative Care*, 17(1).

Weaver, M., Wichman, C., Darnall, C., Bace, S., Vail, C., & Macfadyen, A. (2018). [Proxy-Reported Quality of Life and Family Impact for Children Followed Longitudinally by a Pediatric Palliative Care Team.](#) *Journal of Palliative Medicine*, 21(2), 241–244.

Zaal-Schuller, I. H., Willems, D. I., Ewals, F., van Goudoever, J. B., & de Vos, M. A. (2018). [Involvement of nurses in end-of-life discussions for severely disabled children.](#) *Journal of Intellectual Disability Research*.