

Citation lists are generated by <u>PedPalASCNET</u> through monthly searches of PubMed, Medline, and Embase. Visit our <u>blog</u> to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click <u>here</u>.

Trends in Pediatric Palliative Care Citation List 2016; Issue #11

Artis, G., et al. (2016). "Emergency care planning-are we discussing it, are we recording it, and are we acting on it? An audit of ECP implementation in children with life limiting conditions." Archives of Disease in Childhood 101: A288.

Azim, H. (2016). <u>"The spectrum of children's palliative care needs-a useful tool?"</u> Archives of Disease in Childhood 101: A306.

Banaka, M., et al. (2016). <u>"Are we looking after our neonatal nursing colleagues? Experiences from an end of life care support study."</u> Archives of Disease in Childhood 101: A243-A244.

Barnato, A. E., et al. (2016). "Storytelling in the Early Bereavement Period to Reduce Emotional Distress Among Surrogates Involved in a Decision to Limit Life Support in the ICU: A Pilot Feasibility Trial." Crit Care Med.

Beecham, E., et al. (2016). "<u>Keeping all options open: Parents' approaches to advance care planning</u>." Health Expect.

Crooks, H. F., et al. (2016). "An evaluation of non-medical prescribing in a children's hospice service." Archives of Disease in Childhood 101: A307-A308.

Currie, E. R., et al. (2016). <u>"Parent Perspectives of Neonatal Intensive Care at the End-of-Life."</u> J Pediatr Nurs 31(5): 478-489.

Downie, J., et al. (2016). "Advanced planning in perinatal palliative care-are we making clear end of life plans?" Archives of Disease in Childhood 101: A304.

Downie, J., et al. (2016). <u>"Evaluation of the need for a paediatric palliative care service in the west of Scotland."</u> Archives of Disease in Childhood 101: A287-A288.

Gall, C., et al. (2016). <u>"Pediatric Triage in a Severe Pandemic: Maximizing Survival by Establishing Triage Thresholds."</u> Crit Care Med 44(9): 1762-1768.

Garros, D. (2016). "Moral Distress in the Everyday Life of an intensivist." Frontiers in Pediatrics 4: 4.

Goodwin, J., et al. (2016). ""You Don't Know Until You Get There": The Positive and Negative "Lived" Experience of Parenting an Adult Child With 22q11.2 Deletion Syndrome." Health Psychol.

Holland, C., et al. (2016). "Paediatric palliative care: Is it all about communication?-quality improvement project to assess and address background knowledge and training needs of a department." Archives of Disease in Childhood 101: A344-A345.



Citation lists are generated by <u>PedPalASCNET</u> through monthly searches of PubMed, Medline, and Embase. Visit our <u>blog</u> to see monthly commentaries on selected articles. To view a citation in Zotero, or to join our Zotero group, click <u>here</u>.

Hutchinson, A. and S. Bertaud (2016). <u>"Trainee experience and understanding in paediatric palliative care."</u> Archives of Disease in Childhood 101: A304.

Jarvis, S., et al. (2016). "How many children and young people with life-limiting conditions are clinically unstable? A national data linkage study." Arch Dis Child.

Mandrell, B. N., et al. (2016). "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." J Neurooncol 129(2): 373-381.

Mherekumombe, M. F., et al. (2016). <u>"Pop Up: A new model of paediatric palliative care."</u> Journal of Paediatrics & Child Health.

Mitchell, S., et al. (2016). "Do specialist paediatric palliative care services benefit children and young people patients with life-limiting or life-threatening conditions and their families?" Archives of Disease in Childhood 101: A306.

Pace, J. C. and T. S. Mobley (2016). "Rituals at End-of-Life." Nurs Clin North Am 51(3): 471-487.

Rafferty, K. A. and S. L. Sullivan (2016). ""You Know the Medicine, I Know My Kid": How Parents Advocate for Their Children Living With Complex Chronic Conditions." Health Commun: 1-10.

Ray, K. N., et al. (2016). "Use of Adult-Trained Medical Subspecialists by Children Seeking Medical Subspecialty Care." J Pediatr 176: 173-181.e171.

Schonfeld, D. J. and T. Demaria (2016). "Supporting the Grieving Child and Family." Pediatrics 138(3).

Shaw, C., et al. (2016). <u>"Parental involvement in neonatal critical care decision-making."</u> Sociology of Health & Illness.

Warlow, T. A., et al. (2016). "Neonatal palliative care: A quality improvement project." Archives of Disease in Childhood 101: A286-A287.

Williams, M., et al. (2016). "Palliative care for children in the community with static neurological conditions-are we getting it right?" Archives of Disease in Childhood 101: A305.

Wool, C., et al. (2016). "Quality Indicators and Parental Satisfaction With Perinatal Palliative Care in the Intrapartum Setting After Diagnosis of a Life-Limiting Fetal Condition." ANS Adv Nurs Sci.

Wylde, C. (2016). "A study to investigate parental satisfaction with the allocation of respite care for their child by a children's hospice." Archives of Disease in Childhood 101: A306-A307.

Zaal-Schuller, I. H., et al. (2016). "How parents and physicians experience end-of-life decision-making for children with profound intellectual and multiple disabilities." Res Dev Disabil 59: 283-293.