

- on physician communication in pediatric palliative care. *Palliat Support Care*. 2007;5(4):355–365.
- Huang IC, Kenzik KM, Sanjeev TY, et al. Quality of life information and trust in physicians among families of children with life-limiting conditions. *Patient Relat Outcome Meas*. 2010;2010(1):141–148.
- Hungerbuehler I, Vollrath ME, Landolt MA. Posttraumatic growth in mothers and fathers of children with severe illnesses. *J Health Psychol*. 2011;16(8):1259–1267.
- Immelt S. Psychological adjustment in young children with chronic medical conditions. *J Pediatr Nurs*. 2006;21(5):362–377.
- Jacobs HH. Ethics in pediatric end-of-life care: a nursing perspective. *J Pediatr Nurs*. 2005;20(5):360–369.
- James L, Johnson B. The needs of parents of pediatric oncology patients during the palliative care phase. *J Pediatr Oncol Nurs*. 1997;14(2):83–95.
- Jokinen P. The family life-path theory: a tool for nurses working in partnership with families. *J Child Health Care*. 2004;8(2):124–133.
- Jones TL, Prinz RJ. Potential roles of parental self-efficacy in parent and child adjustment: a review. *Clin Psychol Rev*. 2005;25(3):341–363.
- Kassam A, Wolfe J. The ambiguities of free-standing pediatric hospices. *J Palliat Med*. 2013;16(7):716–717.
- Kavanaugh K, Moro TT, Savage TA. How nurses assist parents regarding life support decisions for extremely premature infants. *J Obstet Gynecol Neonatal Nurs*. 2010;39(2):147–158.
- Keim-Malpass J, Hart TG, Miller JR. Coverage of palliative and hospice care for pediatric patients with a life-limiting illness: a policy brief. *J Pediatr Health Care*. 2013;27(6):511–516.
- Kirk S, Glendinning C, Callery PJ. Parent or nurse? The experience of being the parent of a technology-dependent child. *Adv Nurs*. 2005;51(5):456–464.
- Klass D. The inner representation of the dead child in the psychic and social narratives of bereaved parents. Neimeyer RA. *Meaning reconstruction and the experience of*