- Child. 2014;99(3):216-220.
- Sullivan-Bolyai S, Sadler L, Knafl KA, et al. Great expectations: a position description for parents as caregivers, part I. *Pediatr Nurs*. 2003;29(6):52–56.
- Swallow V, Macfadyen A, Santacroce SJ, et al. Fathers' contributions to the management of their child's long-term medical condition: a narrative review of the literature. *Health Expect*. 2012;15(2):157–175.
- Thibodeaux AG, Deatrick JA. Cultural influence on family management of children with cancer. *J Pediatr Oncol Nurs*. 2007;24(4):227–233.
- Thomlinson EH. The lived experience of families of children who are failing to thrive. *J Adv Nurs*. 2002;39(6):537–545.
- Toomey SL, Chien AT, Elliott MN, et al. Disparities in unmet need for care coordination: the national survey of children's health. *Pediatrics*. 2013;131(2):217–224.
- Treyvaud K. Parent and family outcomes following very preterm or very low birth weight birth: a review. *Semin Fetal Neonatal Med.* 2014;19(2):131–135.
- Vance JC, Najman JM, Thearle MJ, et al. Psychological changes in parents eight months after the loss of an infant from stillbirth, neonatal death, or sudden infant death syndrome—a longitudinal study. *Pediatrics*. 1995;96(5):933–938.
- von Lützau P, Otto M, Hechler T, et al. Children dying from cancer: parents' perspectives on symptoms, quality of life, characteristics of death, and end-of-life decisions. *J Palliat Care*. 2012;28(4):274–281.
- Wiener L, McConnell DG, Latella L, et al. Cultural and religious considerations in pediatric palliative care. *Palliat Support Care*. 2013;11(1):47–67.
- Whitehead LC, Gosling V. Parent's perceptions of interactions with health professionals in the pathway to gaining a diagnosis of tuberous sclerosis (TS) and beyond. *Res Dev Disabil*. 2003;24(2):109–119.
- Wolfe J, Friebert S, Hilden J. Caring for children with advanced cancer integrating palliative care. *Pediatr Clin North Am.* 2002;49(5):1043–1062.
- Wolfe J, Grier HE, Klar N, et al. Symptoms and suffering at