

- 2014;63(2):1–21.
- Chawarska K, Shic F, Macari S, et al. 18-month predictors of later outcomes in younger siblings of children with autism spectrum disorder: a baby siblings research consortium study. *J Am Acad Child Adolesc Psychiatry*. 2014;53(12):1317–1327.
- Clifford S, Dissanayake C, Bui QM, et al. Autism spectrum phenotype in males and females with fragile X full mutation and permutation. *J Autism Dev Disord*. 2007;37(4):738–747.
- Colella-Santos MF, Hein TA, de Souza GL, et al. Newborn hearing screening and early diagnostic in the NICU. *Biomed Res Int*. 2014;845308.
- Fabry DA, Davila EP, Arheart KL, et al. Secondhand smoke exposure and the risk of hearing loss. *Tob Control*. 2011;20(1):82–85.
- Fernell E, Eriksson MA, Gillberg C. Early diagnosis of autism and impact on prognosis: a narrative review. *Clin Epidemiol*. 2013;5:33–43.
- Finucane B, Abrams L, Cronister A, et al. Genetic counseling and testing for FMRI gene mutations: practice guidelines of the National Society of Genetic Counselors. *J Genet Couns*. 2012;21(6):752–760.
- Golnik A, Maccabee-Ryaboy N. Autism: clinical pearls for primary care. *Contemp Pediatr*. 2010;42–60.
- Grafodatskaya D, Chung B, Szatmari P, et al. Autism spectrum disorders and epigenetics. *J Am Acad Child Adolesc Psychiatry*. 2010;49(8):794–809.
- Grindle CR. Pediatric hearing loss. *Pediatr Rev*. 2014;35(11):456–463.
- Haddad J. Hearing loss. Kliegman RM, Stanton RF, St. Geme III, et al. *Nelson textbook of pediatrics*. ed 18. Elsevier/Saunders: Philadelphia; 2011.
- Hagerman RJ, Berry-Kravis E, Kaufmann WE, et al. Advances in the treatment of fragile X syndrome. *Pediatrics*. 2009;123(1):378–390.
- Hagerman R, Lauterborn J, Au J, et al. Fragile X syndrome and targeted treatment trials. *Results Probl Cell Differ*. 2012;54:297–335.