**16 Newborn Screening**

**Selected References**

Advisory Committee on Heritable Disorders in Newborns and Children. Recommended uniform screening panel. https://www.hrsa.gov/advisory-committees/heritable-disorders/rusp/index.html. Accessed August 28, 2018.

American College of Medical Genetics. Newborn screening: towards a uniform screening panel and system. *Genetic Med*. 2006;8(suppl):S12-S252.

American College of Medical Genetics Newborn Screening Expert Group. Genetics Home Reference. Genetic conditions. Bethesda, MD: Lister Hill National Center for Biomedical Communications, U.S. National Library of Medicine National Institutes of Health Department of Health and Human Services; 2007. http://ghr.nlm.nih.gov. Accessed October 24, 2012.

Bonham JR, Carling RS, Lindner M. Raising awareness of false positive newborn screening results arising from Pivalate containing creams and antibiotics in Europe when screening for isovaleric acidemia. *J Neonatal Screen*. 2018;4(1):8.

Friedman JM, Cornel MC, Goldenberg AJ, et al. Genomic newborn screening: public health policy considerations and recommendations. *BMC Med Genom*. 2017;10(1):9.

Joint Committee on Infant Hearing. Year 2007 position statement: principles and guidelines for early hearing detection and intervention programs. *Pediatrics*. 2007;120(4):898-921.

Kaye CI, the Committee on Genetics. Newborn screening fact sheets. *Pediatrics*. 2006;118(3):e934-e963.

King JS, Smith ME. Whole-genome screening of newborns? The constitutional boundaries of state newborn screening programs. *Pediatrics*. 2016;137(suppl 1):S8-S15.

Levy PA. An overview of newborn screening. *J Dev Behav Pediatr*. 2010;31:622-631.

Lockwood C, Lemons J, eds. *Guidelines for Perinatal Care*. 6th ed. Elk Grove, IL: American Academy of Pediatrics & The American College of Obstetricians and Gynecologists; 2007:223-224.

Lu KD, Engmann C, Moya F, Muhlebach M. Cystic fibrosis in premature infants. *J Perinatol*. 2011;31:504-508.

Newborn Screening Authoring Committee. Newborn screening expands: recommendations for pediatricians and medical homes—implications for the system. *Pediatrics*. 2008;121:192-217.

Sweetman L, Millington DS, Therrell BL, et al. Naming and counting disorders (conditions) included in newborn screening panels. *Pediatrics*. 2006;117(5 Pt 2):S308-S314.

Watson MS, Mann MY, Lloyd-Puryear MY, et al. Newborn screening: toward a uniform screening panel and system—executive summary. *Pediatrics*. 2006;117:S296-S307.