Risk Factors Associated with Childhood Hearing Loss

Regardless of previous hearing-screening outcomes, all infants with or without risk factors should receive ongoing surveillance of communicative development beginning at 2 months of age during well-child visits in the medical home (AAP Committee, 2017). This recommendation provides an alternative, more inclusive strategy of surveillance of all children within the medical home based on the pediatric periodicity schedule (AAP Committee, 2017; AAP, 2014a). Should a parent have concern regarding speech, language, developmental delay or developmental regression, an immediate referral should be made to an audiologist and speech-language pathologist for further evaluation.


A child should be seen by an audiologist for a hearing evaluation no later than three months after the occurrence if one or more of the following risk factors are present:

- Bacterial and viral meningitis (especially herpes viruses and varicella) or encephalitis
- Congenital Cytomegalovirus (CMV) confirmed in infant
- Extra-corporeal membrane oxygenation (ECMO)
- Head injury (especially basal skull/temporal bone fracture)
- Chemotherapy

A child should see an audiologist for a hearing evaluation by nine months of age if one or more of the following risk factors are present in the period immediately before or right after birth.

- Family history of hearing loss (permanent, sensorineural hearing loss since childhood)
- Cranio-facial anomalies (includes cleft lip or palate, microtia (abnormally small ear), atresia (blocked or abnormally small ear canal), ear dysplasia, microphthalmia, white forelock, congenital microphthalmia, congenital or acquired hydrocephalus, or temporal bone abnormalities
- Exchange transfusion for elevated bilirubin regardless of length of stay
- NICU stay longer than five days
- Aminoglycoside (includes: Gentamycin, Vacomycin, Kanamycin, Streptomycin, Tobramycin) administered for more than five days
- In utero infections such as herpes, rubella, syphilis, and toxoplasmosis
- Asphyxia or Hypoxic Ischemic Encephalopathy
- Syndromes (includes: Trisomy 21-Down syndrome, Goldenhar, Pierre Robin, CHARGE association, Rubinstein-Taybi, Stickler, Usher, osteopetrosis, Neurofibromatosis type II, Treacher Collins, Hunter syndrome, Friedreich’s ataxia, Charcot-Marie-Tooth syndrome or visit the Hereditary Hearing Loss website)