

Table I. Risk Indicators for Progressive or Delayed-Onset Hearing Loss
(For Use with Neonates and Infants Through 2 Years of Age)

Family history of permanent childhood hearing loss		
<ul style="list-style-type: none"> • Mother of child • Father of child 	<ul style="list-style-type: none"> • Grandmother of child • Grandfather of child 	<ul style="list-style-type: none"> • 1st cousin of child • More than one relative of the same parent
<ul style="list-style-type: none"> • Sister of child • Brother of child 	<ul style="list-style-type: none"> • Aunt of child • Uncle of child 	
Stigmata or other findings associated with a syndrome known to include a sensorineural or conductive hearing loss or Eustachian tube dysfunction		
<ul style="list-style-type: none"> • Branchio-oto-renal (BOR) • Noonan • CHARGE association • Pierre Robin 	<ul style="list-style-type: none"> • Stickler • Williams • Zellweger • Goldenhar (oculo-auriculo-vertebral or OAV) • <u>Trisomy 8</u> (Warkany Syndrome) or Trisomy 9 (Mosaic Syndrome) 	<ul style="list-style-type: none"> • Trisomy 21 – Down syndrome • Trisomy 18 – Edwards syndrome • Trisomy 13 – Patau syndrome
Postnatal infections associated with sensorineural hearing loss		
<ul style="list-style-type: none"> • Confirmed bacterial meningitis 	<ul style="list-style-type: none"> • Confirmed viral meningitis 	
In utero infections		
<ul style="list-style-type: none"> • Cytomegalovirus • Herpes 	<ul style="list-style-type: none"> • Rubella • Syphilis 	<ul style="list-style-type: none"> • Toxoplasmosis
Neonatal indicators		
<ul style="list-style-type: none"> • Intensive care greater than (>) 5 days • Extracorporeal membrane oxygenation (ECMO) 	<ul style="list-style-type: none"> • Exposure to ototoxic medications: at risk aminoglycoside exposure • Mechanical ventilation 	<ul style="list-style-type: none"> • Hyperbilirubinemia requiring exchange transfusion
Syndromes associated with progressive hearing loss		
<ul style="list-style-type: none"> • Neurofibromatosis • Osteopetrosis • Alport 	<ul style="list-style-type: none"> • Jervell & Lange-Nielson • Waardenburg • Pendred 	<ul style="list-style-type: none"> • Usher
Neurodegenerative disorders, such as		
<ul style="list-style-type: none"> • Hunter syndrome 	<ul style="list-style-type: none"> • Charcot-Marie-Tooth syndrome 	<ul style="list-style-type: none"> • Friedreich's ataxia
Head trauma requiring hospitalization		
<ul style="list-style-type: none"> • Basal skull/temporal bone fracture 	Other – specify if chosen	
Parental or caregiver concern regarding hearing, speech, language, and or developmental delay		
Craniofacial Anomalies		
<ul style="list-style-type: none"> • Pinna • Cleft palate 	<ul style="list-style-type: none"> • Atresia • Microtia 	<ul style="list-style-type: none"> • Choanal atresia • Temporal bone anomalies
Chemotherapy		

Based on Year 2007 Position Statement: Principles and Guidelines for Early Hearing Detection and Intervention Programs, Joint Committee on Infant Hearing.