



## Monitoring Children with Risk Factors

**Policy** Children with risk factors related to late onset or progressive hearing loss will be monitored so that if hearing loss develops, it will be detected as early as possible.

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**Background** The Iowa EHDI High-Risk Monitoring Protocol is based on the Joint Committee on Infant Hearing 2007 position statement. Emphasis is placed on follow-up as deemed appropriate by the primary health care provider and audiologist. The Iowa protocol describes the follow-up process for children with risk factors.

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**Hospital/AEA/  
Private  
Audiologists/  
Health Care  
Provider  
Protocol** Best practice indicates that risk factors for late onset hearing loss be reported through the statewide EHDI data management or by reporting the risk factors on the screening and diagnostic reporting forms. Follow-up with families of children at risk for hearing loss can only be successful if risk factors for late onset hearing loss are reported to the EHDI program.

See the appendix for a list of risk factors for late onset hearing loss.

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**Risk Factors  
Follow-Up  
Procedures** These procedures will apply to babies born in 2008 and beyond.

- Each month, EHDI staff members will search for babies born two months prior who have risk factors listed in the statewide EHDI data management system. (For example, in March, staff members will run a report of babies born in January who have a risk factor for hearing loss.)
- Babies will be sorted based on the risk factor they have. The risk factors will determine when follow-up is recommended.
- EHDI staff members will send the letters in Appendix K to families and primary health care providers of children with risk factors. The letters state:
  - The child should see an audiologist for a hearing evaluation by six months of age if one or more of the following risk factors are present:
    - Bacterial and viral meningitis
    - Congenital Cytomegalovirus (CMV) confirmed in infant
    - Extra-corporeal membrane oxygenation (ECMO)

- Family history of permanent, sensorineural hearing loss during childhood
  - Head injury
  - Neurodegenerative disorder
  - Other postnatal infections
  - Parental concern regarding hearing status
  - Syndromes
- The child should see an audiologist for a hearing evaluation by 24 to 30 months of age if one or more of the following risk factors are present:
  - Cranio-facial anomalies
  - Exchange transfusion for elevated bilirubin
  - Herpes infection confirmed in infant
  - NICU stay longer than five days
  - Other congenital infection
  - Ototoxic medications administered
  - PPHN (persistent pulmonary hypertension) associated with mechanical ventilation
  - Rubella infection confirmed in infant
  - Syphilis infection confirmed in infant
  - Toxoplasmosis infection confirmed in infant
- Babies having a risk factor(s) requiring follow-up at six months and a risk factor(s) requiring follow-up at 24 to 30 months will receive only a six month letter.
- The EHDI staff will periodically search for babies born in earlier months so children whose records are entered into statewide EHDI data management system late will receive risk factor follow-up.

## **Risk Factors Associated with Childhood Hearing Loss**

### **Care Giver Concern of Hearing Loss**

### **Chemotherapy**

### **Congenital Infection - confirmed in infant**

Includes: cytomegalovirus, rubella, syphilis, herpes, toxoplasmosis

### **Craniofacial Anomalies**

Includes: Cleft lip or palate, microtia (abnormally small ear), atresia (blocked or abnormally small ear canal), choanal atresia



### **Family History**

A family history of permanent, sensorineural hearing loss during childhood

### **Head Trauma**

Especially basal skull/temporal bone fracture requiring hospitalization

### **Hyperbilirubinemia**

Requiring exchange transfusion

### **Mechanical Ventilation or Extracorporeal Membrane Oxygenation (ECMO)**

Associated with respiratory complications such as bronchopulmonary dysplasia (BPD), persistent pulmonary hypertension of the newborn (PPHN), and respiratory distress syndrome (RDS)

### **Neonatal Intensive Care Unit (NICU) Admission Greater Than Five Days**

Generally indicates health complications, and possible ototoxic medication treatment

### **Neurodegenerative Disorders**

Hunter syndrome, Friedreich's ataxia, Charcot-Marie-Tooth syndrome

### **Neurological Conditions and Syndromes**

Includes: seizures, hydrocephalus, intra-ventricular hemorrhage, retinopathy of prematurity

### **Ototoxic Medications**

Includes: Gentamycin, Vacomycin, Kanamycin, Streptomycin, Tobramycin

### **Physical Findings Associated with Sensorineural or Permanent Conductive Hearing Loss**

Example: White forelock

**Postnatal Infection**

Includes: Bacterial meningitis, herpes, varicella

**Recurrent or Persistent Otitis Media with Effusion Greater Than Three Months****Syndromes Associated with Sensorineural, Conductive, or Progressive Hearing Loss**

Includes: Trisomy 21-Down syndrome, Goldenhar, Pierre Robin, CHARGE association, Rubinstein-Taybi, Stickler, Usher, osteopetrosis, Neurofibromatosis type II, Treacher Collins

*This list of risk factors above is based on the Joint Committee on Infant Hearing Recommendations (JCIH ) published Fall 2007.*

**Sources:**

Cone-Wesson, B, Vohr, BR, Slinger, YS, Widen, JE, Folsom, RC, Gorga, MP, Norton, SJ. (2000). Identification of Neonatal Hearing Impairment: Infants with Hearing Loss. *Ear and Hearing*. 21(5): 488-507.

Joint Committee on Infant Hearing. (2007). Year 2007 Position Statement: Principles and Guidelines for Early Hearing Detection and Intervention Programs. *American Academy of Pediatrics*. 120: 898-921. <http://pediatrics.aappublications.org/cgi/content/full/120/4/898>.