## NEW MEXICO DEPARTMENT OF

Public Health Division, Family Health Bureau, Children's Medical Services Early Hearing Detection and Intervention (EHDI) Program

## Report of Audiological/Screening/Diagnostic Results (ROAR)

Securely email or fax completed ROAR to NMDOH Early Hearing Detection & Intervention (EHDI) Program: newborn.hearing@doh.nm.gov

Phone: 1-877-890-4692 or Fax: 505-827-5995

Use the ROAR to report screening and diagnostic results for children with a DOB in the current or previous calendar years.

Provide Audiological Report and Cover Sheet (instead of a ROAR) for children up to age 4 when diagnosed with permanent (not fluctuating conductive) hearing loss.

Reporting Date:	Birth Hospital/Midwife Practice:
Audiologist:	Practice:
Physician:	Practice:
Child's First and Last Name:	Child's DOB:
Mother's First and Last Name:	
Parent/Guardian First and Last 1	Name:
Address:	Phone #:
Please complete all appropria	te sections:
Missed Appointment Date:	Next Appointment Date:
Middle Ear Dysfunction Referral to ENT	on (Screening/Diagnostic Postponed)
ENT Appointment Date:	ENT Practice:
Right Ear: Pass Refe	Type of Screening:OAEAABR  rAtresia/MicrotiaUnsuccessful Screening rAtresia/MicrotiaUnsuccessful Screening
	N:  AE/TOAESedated ABRNon-Sedated ABRASSR Play Audiometry (CPA)Visual Reinforcement Audiometry (VRA)
HEARING STATUS:	
Unable to Complete Test (future diagnostics needed)	Fluctuating Hearing Loss (middle ear dysfunction)
Confirmed Normal Hearin	g
Comments:	

## Risk Factor Assessment Checklist for Reporting to the EHDI Program:

Joint Committee on Infant Hearing (JCIH) Risk Factors:

Perinatal:
☐ Family history* of early, progressive, or delayed onset permanent childhood hearing loss
□ Neonatal intensive care of more than 5 days
☐ Hyperbilirubinemia with exchange transfusion regardless of length of stay
☐ Aminoglycoside administration for more than 5 days**
☐ Asphyxia or Hypoxic Ischemic Encephalopathy
☐ Extracorporeal membrane oxygenation (ECMO)*
☐ In utero infections, such as herpes, rubella, syphilis, toxoplasmosis, cytomegalovirus (CMV), Zika
☐ Birth conditions or findings such as:
<ul> <li>Craniofacial malformations including microtia/atresia, ear dysplasia, oral facial cleft lip/cleft palate, white forelock, and microphthalmia</li> </ul>
☐ Congenital microcephaly, congenital or acquired hydrocephalus
☐ Temporal bone abnormalities
☐ Syndromes*** associated with hearing loss or progressive or late-onset hearing loss (see HereditaryHearingLoss.org) that include:
☐ Alport
☐ Goldenhar
☐ Treacher-Collins,
$\Box$ CHARGE
□ Waardenburg
☐ Trisomy 21
$\Box$ BOR
Comments:
Perinatal/Postnatal:
<ul> <li>Culture-positive postnatal infections associated with sensorineural hearing loss***, including confirmed bacterial and viral (especially herpes viruses and varicella) meningitis, or encephalitis.</li> <li>Events associated with hearing loss:</li> </ul>
<ul> <li>☐ Significant head trauma, especially basal skull/temporal bone fractures</li> <li>☐ Chemotherapy*</li> </ul>
☐ Caregiver concern**** regarding hearing, speech, language, or developmental delay
Comments:
* Infants at increased risk of delayed onset or progressive hearing loss
**Infants with toxic levels or with a known genetic susceptibility remain at risk
***Syndromes (Van Camp & Smith, 2016)
****Parental/caregiver concern should always prompt further evaluation.