

For Deaf-Blind Project Office Use ONLY: ID#:

Kidcode:

2019 NM Deaf-Blind Census Reporting Form
UNM Project For New Mexico Children and Youth Who Are Deaf-Blind
2300 Menaul Blvd., NE
Albuquerque, New Mexico 87107
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505.272.0321 or Toll Free 877.614.4051
FAX: 505.272.3140

STOP!! Complete this form ONLY for individuals who have BOTH a visual and auditory impairment.

Today's Date: / /

Part I: Information about individual with deaf-blindness:

Name: First: _____ Last: _____

Date of Birth: (MM/DD/YYYY) ____/____/____ **Gender:** Male Female

Race/Ethnicity (select the ONE that best describes the individual's race/ethnicity)

- | | | |
|---|--|---|
| <input type="checkbox"/> 1. American Indian or Alaskan Native | <input type="checkbox"/> 4. Hispanic/Latino | <input type="checkbox"/> 7. Two or more races |
| <input type="checkbox"/> 2. Asian | <input type="checkbox"/> 5. White (Not Hispanic) | |
| <input type="checkbox"/> 3. Black or African American | <input type="checkbox"/> 6. Native Hawaiian/Pacific Islander | |

Living Setting: (Select the ONE setting that best describes where the individual resides the majority of the year):

- | | | |
|--|---|--|
| <input type="checkbox"/> 1. Home: Birth/Adoptive Parents | <input type="checkbox"/> 5. Private Residential Facility | <input type="checkbox"/> 9. Pediatric Nursing Home |
| <input type="checkbox"/> 2. Home: Extended Family | <input type="checkbox"/> 6. Group Home (less than 6 residents) | <input type="checkbox"/> 555: Other _____ |
| <input type="checkbox"/> 3. Home: Foster Parents | <input type="checkbox"/> 7. Group Home (6 or more residents) | |
| <input type="checkbox"/> 4. State: Residential Facility | <input type="checkbox"/> 8. Apartment (with non-family persons) | |

Parent/Guardian Name 1 First: _____ Last: _____

Street Address: _____ City: _____ State: ____ Zip: _____

Telephone: (____) _____ County of Residence: _____

Email address: (please print) _____

Parent/Guardian Name 2 First: _____ Last: _____

Street Address: _____ City: _____ State: ____ Zip: _____

Telephone: (____) _____ County of Residence: _____

Email address: (please print) _____

Part II: Individuals Medical Background/Handicapping Conditions:

Primary Classification of Visual Impairment (select the ONE that best describes the primary classification of the individual's visual impairment)

- | | | |
|---|--|---|
| <input type="checkbox"/> 1. Low Vision | <input type="checkbox"/> 4. Totally Blind | <input type="checkbox"/> 9. Documented Functional Vision Loss |
| <input type="checkbox"/> 2. Legally Blind | <input type="checkbox"/> 6. Diagnosed Progressive Loss | |
| <input type="checkbox"/> 3. Light Perception Only | <input type="checkbox"/> 7. Further Testing Needed | |

Cortical Vision Impairment? Yes No Unknown

Primary Classification of Hearing Impairment (select the ONE that best describes the primary classification of the individual's auditory impairment):

- | | | |
|---|--|--|
| <input type="checkbox"/> 1. Mild | <input type="checkbox"/> 4. Severe | <input type="checkbox"/> 7. Further Testing Needed (1 year only) |
| <input type="checkbox"/> 2. Moderate | <input type="checkbox"/> 5. Profound | <input type="checkbox"/> 9. Documented Functional Hearing Loss |
| <input type="checkbox"/> 3. Moderately Severe | <input type="checkbox"/> 6. Diagnosed Progressive Loss | |

Central Auditory Processing Disorder (CAPD)? Yes No Unknown
Auditory Neuropathy? Yes No Unknown
Cochlear Implant? Yes No Unknown

Other Impairments? (indicate Yes OR No for each):
 Orthopedic/Physical Impairments Yes No Cognitive Impairments Yes No
 Behavioral Disorder Yes No Complex Health Care Needs Yes No
 Communication Impairments Yes No Other: _____ Yes No

Etiology (please indicate the ONE etiology from the list below that best describes the primary etiology of the individual's primary disability. Please indicate "Other" if none of the listed etiologies is the primary disability).

1. Hereditary/Chromosomal Syndromes and Disorders

- | | |
|---|---|
| <input type="checkbox"/> 101. Aicardi Syndrome | <input type="checkbox"/> 130. Marshall Syndrome |
| <input type="checkbox"/> 102. Alport Syndrome | <input type="checkbox"/> 131. Maroteaux-Lary Syndrome (MPS VI) |
| <input type="checkbox"/> 103. Alstrom Syndrome | <input type="checkbox"/> 132. Moebius Syndrome |
| <input type="checkbox"/> 104. Apert Syndrome (Acrocephalosyndactyly, Type!) | <input type="checkbox"/> 133. Monosomy Tenp |
| <input type="checkbox"/> 105. Bardet-Biedl Syndrome (Laurence Moon-Biedl) | <input type="checkbox"/> 134. Morquio Syndrome (MPS IV-B) |
| <input type="checkbox"/> 106. Batten Disease | <input type="checkbox"/> 135. NF One – Neurofibromatosis |
| <input type="checkbox"/> 107. CHARGE Association | <input type="checkbox"/> 136. NF Two – Bilateral Acoustic Neurofibromatosis |
| <input type="checkbox"/> 108. Chromosome Eighteen, Ring Eighteen | <input type="checkbox"/> 137. Norrie Disease |
| <input type="checkbox"/> 109. Cockayne Syndrome | <input type="checkbox"/> 138. Optico-Cochleo Dentate Degeneration |
| <input type="checkbox"/> 110. Cogan Syndrome | <input type="checkbox"/> 139. Pfeiffer Syndrome |
| <input type="checkbox"/> 111. Cornelial de Lange | <input type="checkbox"/> 140. Prader-Willi |
| <input type="checkbox"/> 112. Cri du chat Syndrome (Chromosome 5p-Syndrome) | <input type="checkbox"/> 141. Pierre-Robin Syndrome |
| <input type="checkbox"/> 113. Crigler-Naggar Syndrome | <input type="checkbox"/> 142. Refsum Syndrome |
| <input type="checkbox"/> 114. Crouzon Syndrome (Craniofacial Dysotosis) | <input type="checkbox"/> 143. Scheie Snyderome (MPS I-S) |
| <input type="checkbox"/> 115. Dandy Walker Syndrome | <input type="checkbox"/> 144. Smith-Lemli – Optiz (SLO) Syndrome |
| <input type="checkbox"/> 116. Down Syndrome (Trisomy Twenty-one) | <input type="checkbox"/> 145. Stickler Syndrome |
| <input type="checkbox"/> 117. Goldenhar Syndrome | <input type="checkbox"/> 146. Sturge-Weber Syndrome |
| <input type="checkbox"/> 118. Hand-Schuller-Christian (Histiocytosis X) | <input type="checkbox"/> 147. Treacher Collins Syndrome |
| <input type="checkbox"/> 119. Hallgren Syndrome | <input type="checkbox"/> 148. Trisomy Thirteen (Patau Syndrome) |
| <input type="checkbox"/> 120. Herpes-Zoster (or Hunt) | <input type="checkbox"/> 149. Trisomy Eighteen (Edwards Syndrome) |
| <input type="checkbox"/> 121. Hunter Syndrome (MPSII) | <input type="checkbox"/> 150. Turner Syndrome |
| <input type="checkbox"/> 122. Hurler Syndrome (MPS I-H) | <input type="checkbox"/> 151. Usher I Syndrome |
| <input type="checkbox"/> 123. Kearns-Sayre Syndrome | <input type="checkbox"/> 152. Usher II Syndrome |
| <input type="checkbox"/> 124. Klippel-Feil Sequence | <input type="checkbox"/> 153. Usher III Syndrome |
| <input type="checkbox"/> 125. Klippel-Trenaunay-Weber Syndrome | <input type="checkbox"/> 154. Vgot-Koyanagi-Harada Syndrome |
| <input type="checkbox"/> 126. Kniest Dysplasia | <input type="checkbox"/> 155. Waardenburg Syndrome |
| <input type="checkbox"/> 127. Leber Congenital Anaurosis | <input type="checkbox"/> 156. Wildervanck Syndrome |
| <input type="checkbox"/> 128. Leigh Disease | <input type="checkbox"/> 157. Wolf-Hirschhorn Syndrome (Trisomy 4p) |
| <input type="checkbox"/> 129. Marfan Syndrome | <input type="checkbox"/> 199. Other _____ |

2. Pre-Natal/Congenital Complications

- | | | |
|---|--|--|
| <input type="checkbox"/> 201. Congenital Rubella Syndrome | <input type="checkbox"/> 205. Fetal Alcohol Syndrome | <input type="checkbox"/> 209. Neonatal Herpes Simplex) |
| <input type="checkbox"/> 202. Congenital Syphilis | <input type="checkbox"/> 206. Hydrocephaly | <input type="checkbox"/> 299. Other _____ |
| <input type="checkbox"/> 203. Congenital Toxoplasmosis | <input type="checkbox"/> 207. Maternal Drug Use | |
| <input type="checkbox"/> 204. Cytomegalovirus (CMV) | <input type="checkbox"/> 208. Macrocephaly | |

3. Post-Natal/Non-Congenital Complications

- | | | | |
|---|--|--------------------------------------|--|
| <input type="checkbox"/> 301. Asphyxia | <input type="checkbox"/> 304. Infections | <input type="checkbox"/> 307. Stroke | <input type="checkbox"/> 309. Chemically Induced |
| <input type="checkbox"/> 302. Direct Trauma to the Eye and/or Ear | <input type="checkbox"/> 305. Meningitis | <input type="checkbox"/> 308. Tumors | <input type="checkbox"/> 399. Other |
| <input type="checkbox"/> 303. Encephalitis | <input type="checkbox"/> 306. Severe Head Injury | | |

4. Related to Prematurity

401. Complications of Prematurity

5. Undiagnosed

501. No Determination of Etiology

Part III: IDEA

Part C - Birth through 2 Years

Special Education Status/Part C Exiting (please indicate the ONE code that best describes the individual's Special Education Program status):

- 0. In a Part C early intervention program
- 1. Completion of IFSP prior to reaching max age for Pt C
- 2. Eligible for IDEA Part B
- 3. Not eligible for Part B, referral to other program
- 4. Not eligible for Part B, exit with no referral
- 5. Part B eligibility not determined
- 6. Deceased
- 7. Moved out of state
- 8. Withdrawal by Parent/Guardian
- 9. Attempt(s) to reach parent/child unsuccessful.

Part C Category Code (please indicate the primary category code under which the individual was reported on the Part C, IDEA Child Count. Select only ONE. See attached guideline for additional information if needed):

- 1. At risk for developmental delays
- 2. Developmentally delayed
- 888 Not reported under Part C of IDEA

Part B – 3 through 21 Years

Special Education Status / Part B Exiting (please indicate the ONE code that best describes the individual's Special Education Program status):

- 0. In ECSE or school aged Special Education Program
- 1. Transferred to regular education
- 2. Graduated with regular diploma
- 3. Received a certificate
- 4. Reached maximum age
- 5. Died
- 6. Moved, known to be continuing
- 7. (intentionally not used)
- 8. Dropped out

Part B Category Code (please indicate the primary category code under which the individual; was reported on Part B IDEA Child Count. Select ONE only. See attached Guideline for additional information, if needed):

- 1. Intellectual Disability
- 2. Hearing Impairment (includes deafness)
- 3. Speech or Language Impairment
- 4. Visual Impairment (includes blindness)
- 5. Emotional Disturbance
- 6. Orthopedic Impairment
- 7. Other Health Impairment
- 8. Specific Learning Disability
- 9. Deaf-Blindness
- 10. Multiple disabilities
- 11. Autism
- 12. Traumatic Brain Injury
- 13. Developmentally Delayed – age 3 through 9
- 14. Non-Categorical
- 888. Not Reported under Part B of IDEA

Deaf-Blind Project Exiting Status:

- 0. Eligible to receive services from DB Project

Project staff to complete this section

- 1. No longer eligible to receive services from DB Project

Participation in Statewide Assessments

- 1. Regular grade level State assessment
- 2. Regular grade level State assessment w/accommodations
- 3. Alternate assessments
- 4. (intentionally not used)
- 5. (intentionally not used)
- 6. Not yet required at age or grade level
- 7. Parent Opt Out

Educational Setting (indicate the ONE educational setting code from the appropriate age sub category that best describes the individual's educational setting. Please specify "OTHER" if none of the provided codes apply):

Early Intervention Setting

Birth through 2 years of age (if the individual is in this category, please check the ONE box indicating the service(s) setting).

- 1. Home
- 2. Community based setting
- 3. Other setting(s)

Early Childhood Special Education (3-5) Setting

- | | |
|---|--|
| <input type="checkbox"/> 1. In a regular EC program at least 80% of the time | <input type="checkbox"/> 5. Attending a separate school |
| <input type="checkbox"/> 2. In a regular EC program 40% to 79% of the time | <input type="checkbox"/> 6. Attending a residential facility |
| <input type="checkbox"/> 3. In a regular EC program less than 40% of the time | <input type="checkbox"/> 7. Service provider location |
| <input type="checkbox"/> 4. Attending a separate class | <input type="checkbox"/> 8. Home |

School aged (6 – 21) Setting

- | | |
|--|--|
| <input type="checkbox"/> 9. Inside the regular class 80% or more of the day | <input type="checkbox"/> 13. Residential facility |
| <input type="checkbox"/> 10. Inside the regular class 40% to 79% of the day | <input type="checkbox"/> 14. Homebound/Hospital |
| <input type="checkbox"/> 11. Inside the regular classroom less than 40% of the day | <input type="checkbox"/> 15. Correctional facility |
| <input type="checkbox"/> 12. Separate School | <input type="checkbox"/> 16. Parentally placed in private school |

Assistive Technology

- | | | | |
|---------------------------------|------------------------------|-----------------------------|----------------------------------|
| Corrective Lenses | <input type="checkbox"/> Yes | <input type="checkbox"/> No | <input type="checkbox"/> Unknown |
| Assisted Listening Devices | <input type="checkbox"/> Yes | <input type="checkbox"/> No | <input type="checkbox"/> Unknown |
| Additional Assistive Technology | <input type="checkbox"/> Yes | <input type="checkbox"/> No | <input type="checkbox"/> Unknown |

Intervener Services

An Intervener provides consistent one-to-one support to a student who is deaf-blind throughout the instructional day and has completed specialized training in deaf-blindness.

- Is your child/student receiving Intervener Services? Yes No Unknown

School Information

Agency / School: _____

Street Address _____

City: _____ State: _____ Zip: _____

Telephone Number: (____) _____ Fax Number: (____) _____

Teacher/Coordinator Name: *(please print)* _____

Teacher/Coordinator Email Address: *(please print)* _____

School District: _____

Please return this form by April 8th, 2020

If you have any questions, please call

Project for New Mexico Children and Youth Who Are Deaf-Blind

505.272.0321 or Toll Free 877.614.4051

RETURN the Form by FAX: 505.272.3140 or mail:

UNM Project For New Mexico Children and Youth Who Are Deaf-Blind

2300 Menaul Blvd., NE

Albuquerque, New Mexico 87107

Thank you for completing this form, which will assist in program development, and funding!