

**2016 Census Reporting Form**

Please complete and return to: Susan Bonner, Deafblind Project Director, 3815 Magnolia, St. Louis, Missouri 63110.

**STOP!! Complete this form ONLY for individuals who have both a visual and auditory impairment. DO NOT USE for an individual with only a visual impairment.**

Today's Date: \_\_\_\_\_ MOSIS#: \_\_\_\_\_

Status of this Individual's Report (please check one):

- New (individual being added to the Census)  No revisions to last year's information  
 Information revised from last year's reporting

**Part I: Information about individual with deafblindness**

**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):

- 1 American Indian or Alaskan Native  4 Hispanic/Latino  7 Two or more races  
 2 Asian  5 White  
 3 Black or African American  6 Native Hawaiian or other Pacific Islander

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

- 1 Home: Birth/Adoptive Parents  5 Private Residential  9 Pediatric Nursing Home  
 2 Home: Extended Family  6 Group Home (less than six residents)  555 Other:  
 3 Home: Foster Parents  7 Group Home (six or more residents)  
 4 State Residential  8 Apartment (with non-family person(s))

**Parent/Guardian Name 1** First: \_\_\_\_\_ Last: \_\_\_\_\_

Street Address: \_\_\_\_\_

City: \_\_\_\_\_ State: MO ZIP Code: \_\_\_\_\_

Telephone (with Area Code): \_\_\_\_\_ County of Residence: \_\_\_\_\_

**Parent/Guardian Name 2** First: \_\_\_\_\_ Last: \_\_\_\_\_

Street Address: \_\_\_\_\_

City: \_\_\_\_\_ State: MO ZIP Code: \_\_\_\_\_

Telephone (with Area Code): \_\_\_\_\_ County of Residence: \_\_\_\_\_

**Part II: Individual's Medical Background/Handicapping Conditions**

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

- 1 Low Vision  4 Totally Blind  9 Documented Functional Vision Loss  
 2 Legally Blind  6 Diagnosed Progressive Loss  
 3 Light Perception Only  7 Further Testing Needed

Cortical Vision Impairment?  Yes  No

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

- 1 Mild  4 Severe  7 Further Testing Needed  
 2 Moderate  5 Profound  8 XXX  
 3 Moderately Severe  6 Diagnosed Progressive Loss  9 Documented Funtional Hearing Loss

Central Auditory Processing Disorder (CAPD)?  Yes  NoCochlear Implant?  Yes  NoAuditory Neuropathy?  Yes  No**Other Impairments** (indicate YES or NO forPhysical Impairments  Yes  No Complex Health Care Needs  Yes  NoCognitive Impairments  Yes  No Communication Speech/Lang  Yes  NoBehavioral Disorder  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 are the primary disability):

## Hereditary/Chromosomal Syndromes and Disorders

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

## Pre-Natal/Congenital Complications

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

## Post-Natal/Non-Congenital Complications

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

## Related to Prematurity

 401 Complications of Prematurity

## Undiagnosed

 501 No Determination of Etiology

**Part III: IDEA**

**Funding Category** (please indicate the funding category under which the individual was receiving services on **December 1, 2016**):

- 1 IDEA Part B (three through twenty-one)       3 Not reported under Part B or Part C  
 2 IDEA Part C (birth through two)

**-----Part C-----**

**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       6 Deceased  
 1 Completion of IFSP prior to reaching max age for Pt C       7 Moved out of state  
 2 Eligible for IDEA, Part       8 Withdrawal by parent/guardian  
 3 Not eligible for Pt B, referral to other       9 Attempts to reach parent and/or child unsuccessful  
 4 Not eligible for Pt B, exit w/no  
 5 Part B eligibility not determined

**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 as defined by the State's Plan  
 2 Developmentally Delayed  
 888 Not Reported under Part C of IDEA

**-----Part B-----**

**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       5 Died  
 1 Transferred to regular education       6 Moved, Known to be Continuing  
 2 Graduated with regular diploma       7 (intentionally not used)  
 3 Received a certificate       8 Dropped out  
 4 Reached Maximum Age

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

- 1 Mental Retardation       9 Deaf-blindness  
 2 Hearing Impairment (includes deafness)       10 Multiple Disabilities  
 3 Speech or Language Impairment       11 Autism  
 4 Visual Impairment (includes blindness)       12 Traumatic Brain  
 5 Emotional Disturbance       13 Developmentally Delayed-age 3-9  
 6 Orthopedic Impairment       14 Non-Categorical  
 7 Other Health Impairment       15 Not Reported under Part B of IDEA  
 8 Specific Learning

**Deaf-Blind Project Exiting Status:**

- 0 Eligible to receive services from DB  
 1 No longer eligible to receive services from State DB

**Participation in Statewide Assessments**

- 1 Regular grade-level State assessment  
 2 Regular grade-level State assessment w/accommodations  
 3 Alternate assessments  
 6 Not required at age or grade level

**Educational Setting** (indicate the ONE educational setting code from the appropriate age subcategory 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

### ECSE (3-5) Settings

- 1 Attending a regular EC program at least 80% of the time  
 2 Attending a regular EC program 40% to 79% of the time  
 3 Attending a regular EC program less than 40% of the time  
 4 Attending a separate class  
 5 Attending a separate school  
 6 Attending a residential  
 7 Service provider  
 8 Home

### School aged (6-21) Settings

- 9 Inside the regular class 80% or more of the day  
 10 Inside the regular class 40% to 79% of day  
 11 Inside the regular class less than 40% of the day  
 12 Separate School  
 13 Residential facility  
 14 Homebound/Hospital  
 15 Correctional  
 16 Parentally placed in private schools/Homeschooled

### Assistive Technology

- Corrective Lenses  Yes  No  Unknown  
 Assistive Listening Devices  Yes  No  Unknown  
 Additional Assistive Technology  Yes  No  Unknown

### Intervener (One-on-One Paraprofessional)

Does the student receive one-on-one paraprofessional support?  Yes  No  Unknown

### School Information

Agency/School: \_\_\_\_\_

Street Address: \_\_\_\_\_

City: \_\_\_\_\_

State: \_\_\_\_\_

ZIP Code: \_\_\_\_\_

Telephone Number: \_\_\_\_\_

Fax Number: \_\_\_\_\_

Teacher's Name: \_\_\_\_\_

Teacher's Email: \_\_\_\_\_

School District: \_\_\_\_\_

Please return this form and the appropriate Permission for Release Form (if already returned by the individual or parent/guardian, by February 1, 2017 to:

Susan Bonner, Deafblind Project Director  
 Missouri Deafblind Technical Assistance Project  
 3815 Magnolia  
 St. Louis, Missouri 63110  
[susan.bonner@msb.dese.mo.gov](mailto:susan.bonner@msb.dese.mo.gov)