

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, 2014**):

- 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 aligned with grade-level achievement standards
 4 Alternate assessments based on alternate achievement standards
 5 Modified achievement standards
 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 2, 2015 to:

Susan Bonner, Deafblind Project Coordinator
 Missouri Deafblind Technical Assistance Project
 3815 Magnolia
 St. Louis, Missouri 63110
susan.bonner@msb.dese.mo.gov