

Deafblind Student Registry Form



New
 Update
 No Changes
 Deceased
 Delete
 Region

| | | | |
|----------------------------|-------------------------------|--|------------------------------|
| Student Information | Date of Birth / / | Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female | |
| | Last Name: | First Name: | |
| Address | | City | State Zip Code |

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

- 1 American Indian or Alaskan Native
 2 Asian or Pacific Islander
 3 Black (not Hispanic)
 4 Hispanic
 5 White (not Hispanic)

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

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

Parent/Guardian Name 1 First: Last:

Street Address:

City: State: ZIP Code:

Telephone (with Area Code): County of Residence:

Cell Phone: Email:

Parent/Guardian Name 2 First: Last:

Street Address:

City: State: ZIP Code:

Telephone (with Area Code): County of Residence:

Cell Phone: Email:

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

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):

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

Central Auditory Processing Disorder (CAPD)? Yes No UnknownAuditory Neuropathy? Yes No UnknownCochlear Implant? Yes No Unknown**Other Impairments** (indicate YES or NO for each):Physical Impairments Yes No Complex Health Care Needs Yes NoCognitive Impairments Yes No Communication Speech/Lang Yes NoBehavioral Disorder Yes No Other: _____ Yes No**Assistive Technology**Corrective Lenses Yes No Unknown Assistive Listening Devices Yes No UnknownAdditional Assistive Technology Yes No Unknown**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 I) | <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 Neurofibromatosis |
| <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 X) | <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**-----Part C-----**

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

- | | |
|---|--|
| <input type="radio"/> 0 In a Part C early intervention program | <input type="radio"/> 6 Deceased |
| <input type="radio"/> 1 Completion of IFSP prior to reaching max age for Pt C | <input type="radio"/> 7 Moved out of state |
| <input type="radio"/> 2 Eligible for IDEA, Part B | <input type="radio"/> 8 Withdrawal by parent/guardian |
| <input type="radio"/> 3 Not eligible for Pt B, referral to other program | <input type="radio"/> 9 Attempts to reach parent and/or child unsuccessful |
| <input type="radio"/> 4 Not eligible for Pt B, exit w/no referrals | |
| <input type="radio"/> 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
- 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):

- | | |
|--|---|
| <input type="radio"/> 0 In ECSE or school-aged Special Education Program | <input type="radio"/> 5 Died |
| <input type="radio"/> 1 Transferred to regular education | <input type="radio"/> 6 Moved, Known to be Continuing |
| <input type="radio"/> 2 Graduated with regular diploma | <input type="radio"/> 7 (intentionally not used) |
| <input type="radio"/> 3 Received a certificate | <input type="radio"/> 8 Dropped out |
| <input type="radio"/> 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.):

- | | |
|--|--|
| <input type="radio"/> 1 Mental Retardation | <input type="radio"/> 9 Deaf-blindness |
| <input type="radio"/> 2 Hearing Impairment (includes deafness) | <input type="radio"/> 10 Multiple Disabilities |
| <input type="radio"/> 3 Speech or Language Impairment | <input type="radio"/> 11 Autism |
| <input type="radio"/> 4 Visual Impairment (includes blindness) | <input type="radio"/> 12 Traumatic Brain Injury |
| <input type="radio"/> 5 Emotional Disturbance | <input type="radio"/> 13 Developmentally Delayed - age 3 through 9 |
| <input type="radio"/> 6 Orthopedic Impairment | <input type="radio"/> 14 Non-Categorical |
| <input type="radio"/> 7 Other Health Impairment | <input type="radio"/> 888 Not Reported under Part B of IDEA |
| <input type="radio"/> 8 Specific Learning Disability | |

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 settings
- 3 Other Settings

Please see next page for additional Educational Settings.

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 facility
- 7 Service provider location
- 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 facilities
- 16 Parentally placed in private schools

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 yet required

Deaf-Blind Project Exiting Status:

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

School Information

Agency/School: _____

Street Address: _____

City: _____ State: _____ ZIP Code: _____

Telephone Number: _____ Fax Number: _____

Teacher's Name: _____

Teacher's Email: _____

School District: _____

Lyn Ayer, Project Director
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If you have any questions, please call Lyn Ayer at 503-838-8328, or email: ayerl@wou.edu

Thank you for completing this from which will assist in program development and funding.