National Deaf-Blind Census: Ind	○ Update ○ No change ○ Female	
	No longer in district. Moved to:	
Student's Last Name Student's First		
Address	O SSID#(10 digit student ID#)	
	Date of Birth:	
City State	Zip Code / / Month Day Year	
Parent(s)/Guardian	Phone	
PRIMARY IDENTIFIED ETIOLOGY: Sel	ect ONE from the list below that best describes	
	etiology of the individual's primary disability.	
	hromosomal Syndromes and Disorders	
101 Aicardi syndrome	130 Marshall syndrome	
102 Alport syndrome103 Alstrom syndrome	131 Maroteaux-Lamy syndrome (MPS VI)132 Moebius syndrome	
103 Alstrom syndrome (Acrocephalosyndactyly, Type 1)	132 Moreoras Syndrome 133 Monosomy 10p	
105 Bardet-Biedl syndrome (Laurence Moon-Biedl)	134 Morquio syndrome (MPS IV-B)	
106 Batten disease	135 NF1 - Neurofibromatosis (von Recklinghausen disease)	
107 CHARGE association	136 NF2 - Bilateral Acoustic Neurofibromatosis	
108 Chromosome 18, Ring 18	137 Norrie disease	
109 Cockayne syndrome	138 Optico-Cochleo-Dentate Degeneration	
110 Cogan Syndrome	139 Pfeiffer syndrome	
111 Cornelia de Lange	140 Prader-Willi	
112 Cri du chat syndrome (Chromosome 5p-syndrome)113 Crigler-Najjar syndrome	141 Pierre-Robin syndrome 142 Refsum syndrome	
114 Crouzon syndrome (Craniofacial Dysotosis)	143 Scheie syndrome (MPS I-S)	
115 Dandy Walker syndrome	144 Smith-Lemli-Opitz (SLO) syndrome	
116 Down syndrome (Trisomy 21 syndrome)	145 Stickler syndrome	
117 Goldenhar syndrome	146 Sturge-Weber syndrome	
118 Hand-Schuller-Christian (Histiocytosis X)	147 Treacher Collins syndrome	
119 Hallgren syndrome	148 Trisomy 13 (Patau syndrome, Trisomy 13-15)	
120 Herpes-Zoster (or Hunt)	149 Trisomy 18 (Edwards syndrome)	
121 Hunter Syndrome (MPS II)	150 Turner syndrome 151 Usher I syndrome	
122 Hurler syndrome (MPS I-H)123 Kearns-Sayre syndrome	151 Usher II syndrome	
124 Klippel-Feil sequence	153 Usher III syndrome	
125 Klippel-Trenaunay-Weber syndrome	154 Vogt-Koyanagi-Harada syndrome	
126 Kniest Dysplasia	155 Waardenburg syndrome	
127 Leber congenital amaurosis	156 Wildervanck syndrome	
128 Leigh disease	157 Wolf-Hirschhorn syndrome (Trisomy 4p)	
129 Marfan syndrome	199 Other	
Pre-Natal/Congenital Complications	Post-Natal/Non-Congenital Complications	
201 Congenital Rubella Syndrome	301 Asphyxia	
202 Congenital Syphilis203 Congenital Toxoplasmosis	302 Direct Trauma (to the eye and/or ear)303 Encephalitis	
203 Congenital Toxoplasmosis 204 Cytomegalovirus (CMV)	304 Infections	
205 Fetal Alcohol Syndrome	305 Meningitis	
206 Hydrocephaly	306 Severe Head Injury	
207 Maternal Drug Use	307 Stroke	
208 Microcephaly	308 Tumors	
209 Neonatal Herpes Simplex (HSV)	309 Chemically Induced	
299 Other 399 Other		
Related to Prematurity	Undiagnosed	
401 Complications of Prematurity	501 No Determination of Etiology	

RACE ETHNICITY: CHECK ONE BOX ONLY

- 1. American Indian or Alaska Native \bigcirc 3. Black or African American \bigcirc 5. White \bigcirc 2. Asian Ο
 - ○4. Hispanic/Latino

 \bigcirc 7. Two or more races \bigcirc 6. Native Hawaiian or Pacific Islander

VISUAL IMPAIRMENT: PRIMARY CLASSIFICATION OF VISUAL IMPAIRMENT		
CHECK ONE BOX ONLY	CHECK IF APPLICABLE	
1. Low Vision (visual acuity of 20/70 to 20/200 or more <i>in the better eye with correction</i> .)	Cortical Vision Impairment	
○ 2. Legally Blind (visual acuity of 20/200 or less <i>or</i> field restriction of 20 degrees or less <i>in the</i>	0 1 V.	
better eye with correction.)	\bigcirc 1. Yes	
○ 3. Light Perception Only		
○ 4. Totally Blind	○0. No	
○ 6. Diagnosed Progressive Loss		
7. Further Testing Needed to Determine Visual Impairment	\bigcirc 2. Unknown	
 9. Documented Functional Vision Loss 		

HEARING IMPAIRMENT: PRIMARY CLASSIFICATION OF HEARING IMPAIRMENT		
CHECK ONE BOX ONLY	CHECK EITHER "NO," "YES," OR "UNKNOWN"	
 1. Mild (26-40 dB loss) 2. Moderate (41-55 dB loss) 3. Moderately Severe (56-70 dB loss) 4. Severe (71-90 dB loss) 5. Profound (91+ dB loss) 6. Diagnosed Progressive Loss 7. Further Testing Needed to Determine Hearing Impairment 9. Documented Functional Hearing Loss 	Central Auditory Processing Disorder 1. Yes 0. No 2. Unknown Auditory Neuropathy 1. Yes 0. No 2. Unknown Cochlear Implant 1. Yes 0. No 2. Unknown	

Other Impairments: Indicate impairments, in addition to the individual's hearing and visual impairments, that have a significant impact on the individual's development or educational progress.

CHECK ALL CATEGORIES AS EITHER "NO" OR "YES"

1. Orthopedic/Physical Impairments	$\bigcirc 1. $ Yes $\bigcirc 0. $ No	,
2. Cognitive Impairments	$\bigcirc 1. $ Yes $\bigcirc 0. $ No	
3. Behavioral Disorders	$\bigcirc 1. $ Yes $\bigcirc 0. $ No	
4. Complex Health Care Needs	$\bigcirc 1. $ Yes $\bigcirc 0. $ No	
5. Communication, Speech and/or Language	$\bigcirc 1. $ Yes $\bigcirc 0. $ No	
6. Other Impairment(s)	$\bigcirc 1.$ Yes $\bigcirc 0.$ No	(Specify)

PART C (BIRTH THROUGH 2 YRS.) CATEGORY CODES: CHECK ONE BOX ONLY

○ 1.At-risk for developmental delays (as defined by the state's Part C Lead Agency) ○ 888. Not Reported under Part C of IDEA ○ 2. Developmentally Delayed

PART B (3 THROUGH 21 YRS.) CATEGORY CODES*: CHECK ONE BOX ONLY

0	1.	Intellectual	Disability
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- 2. Hearing Impaired/Deaf \bigcirc
- 3. Speech or Language Impairment \bigcirc
- 4. Visually Impaired/Blind Ο
- 5. Emotional/Behavioral \bigcirc
- 6. Orthopedic Impairment 0
- Other Health Impairment \bigcirc 7.
- 8. Specific Learning Disability Ο

- 9. Deaf-Blindness \bigcirc
- 10. Multiple Disabilities \bigcirc
- 11. Autism \bigcirc
- 12. Traumatic Brain Injury Ο
- 13. Developmentally Delayed (age 3 to 9) \bigcirc
- 14. Non-Categorical \bigcirc
- 888. Not Reported under Part B of IDEA \bigcirc

*As child was reported on December 1 Child Count

EDUCATIONAL SETTING: CHECK ONE BOX ONLY		
Birth through Age 2		
○ 1. Home ○ 2. Community-based settings ○ 3. Other settings (specify):		
Ages 3 - 5		
 1. Attending a regular early childhood program at least 80% of the time 2. Attending a regular early childhood program 40% to 79% of the time 3. Attending a regular early childhood program less than 40% of the time 4. Attending a separate class 		
Ages 6 – 21		
 9. Inside the regular class 80% or more of day 10. Inside the regular class 40% to 79% of the 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, including home schooled 		
PARTICIPATION IN STATEWIDE ASSESSMENTS: CHECK ONE BOX ONLY ~ TAKEN FROM THE LAST STATEWIDE ASSESSMENT		
 1. Regular grade-level state assessment 2. Regular grade-level state assessment with accommodations 3. Alternate assessments aligned with grade-level achievement standards 4. Not Used 5. Not Used 6. Not yet required 7. Parent Opt Out 		
PART C (BIRTH THROUGH 2 YRS.) EXITING: CHECK ONE BOX ONLY		
NOTE: CHILDREN WHO HAVE TURNED AGE 3 AND TRANSITIONED FROM PART C TO PART B DURING THE REPORTING PERIOD MAY BE REPORTED UNDER BOTH PART C AND PART B.		
0. In a Part C early intervention program5. Part B eligibility not determined1. Completion of IFSP prior to reaching maximum age for Part C6. Deceased2. Eligible for IDEA, Part B7. Moved out of state3. Not eligible for Part B, exit with referrals to other programs8. Withdrawal by parent (or guardian)4. Not eligible for Part B, exit with no referrals9. Attempts to contact parent and/or child were unsuccessful		
PART B (3 YRS. THROUGH 21 YRS.) EXITING: CHECK ONE BOX ONLY		
NOTE: CHILDREN WHO HAVE TURNED AGE 3 AND TRANSITIONED FROM PART C TO PART B DURING THE REPORTING PERIOD MAY BE REPORTED UNDER BOTH PART C AND PART B.		
 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. Deceased () 6. Moved, known to be continuing. School district moved to: () 8. Dropped out 		
DEAF-BLIND PROJECT EXITING STATUS: CHECK ONE BOX ONLY		
○1. Eligible to receive services from the deaf-blind project ○2. No longer eligible to receive services from the deaf-blind project		
LIVING SETTING: CHECK ONE BOX ONLY		
1. Home: Birth/Adoptive Parents 6. Group Home (less than 6 residents) 2. Home: Extended Family 7. Group Home (6 or more residents) 3. Home: Foster Parents 8. Apartment (with non-family person(s)) 4. State Residential Facility 9. Pediatric Nursing Home 5. Private Residential Facility 555. Other (Specify)		
Assistive Devices: Check either "no," "Yes," or "Unknown"		
Glasses or contact lenses used: Additional assistive technology other than corrective lenses or assistive listening devices: 1. Yes 0. No 2. Unknown Hearing aids, FM system or other listening device used: 1. Yes 0. No 2. Unknown 1. Yes 0. No 2. Unknown 1. Yes 0. No 2. Unknown		

Student's Name:	Date Completed:
	Intervener Services: Check either "NO," "YES," or "UNKNOWN"
	Intervener*: 0. No () 1. Yes () 2. Unknown

*Intervener services are provided by an individual, typically a paraeducator, who has received specialized training in deaf-blindness and the process intervention.

Person Completing Form: (Please fill out #1 and #2 below.)

1. Your Name:	Title:	
Agency Name:		
Address:	City:	
Phone: ()	Email:	
District:	ESD:	
Home School District (if different from serving district):		
2. Teacher's Name:	Title:	
School Name:		
School Address:	City:	Zip:
Teacher's Phone: ()	Email:	
District:	ESD:	
Home School District (if different from serving district): Revised 01/15/2016		

RETURN FORM TO:

Washington State School for the Blind (WSSB) **c/o WA Deafblind Project** 2214 E 13th St, Vancouver WA 98661 Phone: (360) 947-3297

- OR –

FAX to: (833) 903-0338 EMAIL to: khanh.huhtala@wssb.wa.gov