



**CENSUS REPORTING FORM**

(Please Print)

Today's Date:	Name of the Reporter:
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**CHILD/STUDENT INFORMATION**

Last Name:	First Name:	MI:
Gender: ___M ___F	Date of Birth: ___/___/___ <small>MM DD YEAR</small>	Age:
Street Address:	City:	State:
		Zip Code:
County:	Current School:	

**Parent 1/Guardian Information**

Last Name:	First Name:	Phone:
		Home: ( ) _____
		Cell: ( ) _____
		Work:( ) _____
Street Address:	City:	
State:	Zip Code:	
E-mail:	County:	
Relationship to the Child/Student:	<input type="checkbox"/> Parent	<input type="checkbox"/> Guardian

**Parent 2/Guardian Information (If Different from Above)**

Last Name:	First Name:	Phone:
		Home: ( ) _____
		Cell: ( ) _____
		Work:( ) _____
Street Address:	City:	
State:	Zip Code:	
E-Mail:	County:	
Relationship to the Child/Student:	<input type="checkbox"/> Parent	<input type="checkbox"/> Guardian

Living Setting:

___ 1. Home: Birth/Adoptive Parents	___ 5. Private Residential Facility	___ 8. Apartment
___ 2. Home: Extended Family	___ 6. Group Home (less than 6 residents)	___ 9. Pediatric Nursing Hm.
___ 3. Home: Foster Parents	___ 7. Group Home (6 or more residents)	___ 555. Other
___ 4. State Residential Facility		

Ethnicity(Please check one)

___ 1. American Indian/Alaskan Native	___ 4. Hispanic/Latino	___ 7. Two or More Races
___ 2. Asian	___ 5. White	___ 8. Middle Eastern/North Africa
___ 3. Black/African American	___ 6. Native Hawaiian/ Pacific Islander	

**CHILD/STUDENT'S MEDICAL BACKGROUND/HANDICAPPING CONDITION**

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 (no light perception)
6. Diagnosed Progressive Vision Loss
7. Further Testing Needed
9. Documented Functional Vision Loss

Cortical Visual Impairment  
 Does the individual have Cortical Visual Impairment (CVI)?

\_\_\_ Yes \_\_\_ No \_\_\_ Unknown

Primary Classification of **Hearing** Impairment (select 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 Hearing Loss
7. Further Testing Needed
8. Documented Functional Hearing Loss

Auditory Neuropathy  
 Does the individual have Auditory Neuropathy?

\_\_\_ Yes \_\_\_ No \_\_\_ Unknown

Central Auditory Processing Disorder (CAP-D): Does the individual have Central Auditory Processing Disorder?

\_\_\_ Yes \_\_\_ No \_\_\_ Unknown

**Other Impairments:** (Check all that apply)

Physical Impairment            \_\_\_ Yes \_\_\_ No  
 Cognitive Impairment        \_\_\_ Yes \_\_\_ No  
 Behavioral Disorder         \_\_\_ Yes \_\_\_ No  
 Complex Health Care Needs \_\_\_ Yes \_\_\_ No  
 Communication/Speech/Lang \_\_\_ Yes \_\_\_ No

Other

**Assistive Technology:**

Corrective Lenses            \_\_\_ Yes \_\_\_ No  
 Cochlear Implant            \_\_\_ Yes \_\_\_ No  
 Assistive Listening Devices   \_\_\_ Yes \_\_\_ No

Additional Assistive Technology?

\_\_\_\_\_

**ETIOLOGY INFORMATION**

Please indicate the ONE etiology from the list below (if known) 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 & Disorders**

- |   |  |
|---|--|
| <input type="checkbox"/> 101 Aicardi Syndrome                               | <input type="checkbox"/> 130 Marshall Syndrome                             |
| <input type="checkbox"/> 102 Alport Syndrome                                | <input type="checkbox"/> 131 Maroteaux-Lamy Syndrome (MPS VI)              |
| <input type="checkbox"/> 103 Alstrom Syndrome                               | <input type="checkbox"/> 132 Moebius Syndrome                              |
| <input type="checkbox"/> 104 Apert Syndrome (Acrocephalosyndactyly, Type 1) | <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 Syndrome                                | <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 Cornelia de Lange                              | <input type="checkbox"/> 140 Prader-Willi                                  |
| <input type="checkbox"/> 112 Cri du chat Syndrome (Chromosome 5p)           | <input type="checkbox"/> 141 Pierre-Robin Syndrome                         |
| <input type="checkbox"/> 113 Crigler-Najjar Syndrome                        | <input type="checkbox"/> 142 Refsum Syndrome                               |
| <input type="checkbox"/> 114 Crouson Syndrome (Craniofacial Dysotosis)      | <input type="checkbox"/> 143 Scheie Syndrome (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 21)                     | <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 Syndrome Type I                         |
| <input type="checkbox"/> 123 Kearns-Sayre Syndrome                          | <input type="checkbox"/> 152 Usher Syndrome Type II                        |
| <input type="checkbox"/> 124 Klippel-Feil Sequence                          | <input type="checkbox"/> 153 Usher Syndrome Type III                       |
| <input type="checkbox"/> 125 Klippel-Trenaunay-Weber Syndrome               | <input type="checkbox"/> 154 Vogt-Koyanagi-Harada Syndrome                 |
| <input type="checkbox"/> 126 Kniest Dysplasia                               | <input type="checkbox"/> 155 Waardenburg Syndrome                          |
| <input type="checkbox"/> 127 Leber Congenital Amaurosis                     | <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   |

**Pre-Natal/Congenital Complications**

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

**Post-Natal/Non-Congenital Complications**

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

**Related to Prematurity**

- |   |
|---|
| <input type="checkbox"/> 401 Complications of Prematurity |
|---|

**Undiagnosed**

- |   |
|---|
| <input type="checkbox"/> 501 No Determination of Etiology |
|---|

### Attending School District

**Funding Category** (Please indicate the funding category under which the individual was receiving services on January 1, 2017.):

- \_\_\_\_\_ 1. IDEA Part B (ages 3 through 21 years)
- \_\_\_\_\_ 2. IDEA Part C (ages birth through 2 years)
- \_\_\_\_\_ 3. Not reported under Part B or Part C

**Part C Special Education Status**(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 maximum age for Part 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 referrals
- \_\_\_\_\_ 5. Part B eligibility not determined
- \_\_\_\_\_ 6. Deceased
- \_\_\_\_\_ 7. Moved out of state
- \_\_\_\_\_ 8. Withdrawal by parent/guardian
- \_\_\_\_\_ 9. Attempts to reach parent and/or 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 one only.):

- \_\_\_\_\_ 1. At risk for developmental delay
- \_\_\_\_\_ 2. Developmentally Delayed
- \_\_\_\_\_ 888. Not reported under Part C of IDEA

**Part B Special Education Status**(Please indicate the one code that best describes the individual's special education program status.):

- \_\_\_\_\_ 0. In ESCE or school-aged Special Education Program
- \_\_\_\_\_ 1. Transferred to regular education
- \_\_\_\_\_ 2. Graduated with regular diploma
- \_\_\_\_\_ 3. Received a certificate of completion
- \_\_\_\_\_ 4. Reached maximum age
- \_\_\_\_\_ 5. Deceased
- \_\_\_\_\_ 6. Moved out of state, known to be continuing
- \_\_\_\_\_ 7. (Space intentionally left blank)
- \_\_\_\_\_ 8. Dropped out of school

**Part B Category Code** (Please indicate the primary category code under which the individual was reported on The IDEA, Part B Child Count – Select one only).

- \_\_\_\_\_ 1. Intellectual Disabilities
- \_\_\_\_\_ 2. Hearing Impaired (including deafness)
- \_\_\_\_\_ 3. Speech or Language Impairment
- \_\_\_\_\_ 4. Visual Impairment (including blindness)
- \_\_\_\_\_ 5. Emotional Disturbance
- \_\_\_\_\_ 6. Orthopedic Impairment
- \_\_\_\_\_ 7. Other Health Impairment
- \_\_\_\_\_ 8. Specific Learning Disability
- \_\_\_\_\_ 9. Deafblindness
- \_\_\_\_\_ 10. Multiple Disabilities
- \_\_\_\_\_ 11. Autism
- \_\_\_\_\_ 12. Traumatic Brain Injury
- \_\_\_\_\_ 13. Developmentally Delayed (age 3 through 9 years)
- \_\_\_\_\_ 14. Non-Categorical
- \_\_\_\_\_ 888. Not reported under Part B of IDEA

**Participation in Statewide Assessments**

- \_\_\_\_\_ 1. Regular grade-level Statewide Assessment
- \_\_\_\_\_ 2. Regular grade-level Statewide Assessment with accommodations
- \_\_\_\_\_ 3. Alternate Assessment aligned with grade-level achievement standards
- \_\_\_\_\_ 4. Alternate Assessment based on alternate achievement standards
- \_\_\_\_\_ 5. Modified achievement standards
- \_\_\_\_\_ 6. Not yet required

**Educational Setting** Indicate the one educational setting code from the appropriate age subcategory that best describes the individual’s educational setting. Specify “other” if none of the provided codes apply.)

**Early Intervention Settings** (Birth through 2 years of age):

- \_\_\_\_\_ 1. Home
- \_\_\_\_\_ 2. Community-Based Settings
- \_\_\_\_\_ 3. Other settings

**ECSE Settings** (Ages 3 through 5 years of age):

- \_\_\_\_\_ 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 Age Setting** (Ages 6 through 21 years of age)

- \_\_\_\_\_ 9. Inside the regular class 80% or more of the school day
- \_\_\_\_\_ 10. Inside the regular class 40% to 79% of the school day
- \_\_\_\_\_ 11. Inside the regular class less than 40% of the school day
- \_\_\_\_\_ 12. Separate school
- \_\_\_\_\_ 13. Residential facility
- \_\_\_\_\_ 14. Homebound / Hospital setting
- \_\_\_\_\_ 15. Correctional facility
- \_\_\_\_\_ 16. Placed by parents in private school

<b>School Placement Information</b>			
Name of School or Agency:			County:
Street Address	City	State	Zip
Phone #:	Fax#:	Email:	
Contact Person / Position:			

Home School District (LEA) Information			
Name of School District:		County	
Street Address	City	State	Zip
Phone #:	Fax#:	Email:	
Contact Person / Position:			

Is this child registered with the New Jersey Commission for the Blind and Visually Impaired (NJCBVI)? \_\_\_\_\_ Yes \_\_\_\_\_ No

If the response is "Yes" what is the name of the NJCBVI Teacher? \_\_\_\_\_

**THANK YOU VERY MUCH FOR PROVIDING THE INFORMATION ABOVE.**

