

Office Use Only: Name Code: _____ Code Number: _____

New Update No Change

Child Count Form
Nebraska Deaf-Blind Project
NDE, SPED
301 Centennial Mall South
Lincoln, NE 68509

PH: 402-471-4310 or teresa.coonts@nebraska.gov
Website: www.nedbp.org

updated 10-18

Student First Name: Student Middle Initial: Gender
 Male
 Female

Student Last Name: DOB: / /

Parent First Name: Parent Last Name:

Address:

City: State: Zip Code:

Telephone: Email:

Race/Ethnicity: check one only

- American Indian or Alaska Native Asian Hispanic/Latino White
 Black or African American Native Hawaiian or Other Pacific Islander
 Two or more races (does not include persons of Hispanic/Latino ethnicity)

Etiology Select **ONE** from the list below

Hereditary/Chromosomal Syndromes and Disorders

- Aicardi syndrome
- Alport syndrome
- Alstrom syndrome
- Apert Syndrome (Acrocephalosyndactyly Type 1)
- Bardet-Biedl syndrome (Laurence Moon-Biedl)
- Batten disease
- CHARGE association
- Chromosome 18, Ring 18
- Cockayne syndrome
- Cogan syndrome
- Cornelia de Lange
- Cri du chat syndrome (Chromosome 5p-syndrome)
- Crigler-Najjar syndrome
- Crouzon syndrome (Craniofacial Dysostosis)
- Dandy Walker syndrome
- Down syndrome (Trisomy 21 syndrome)
- Goldenhar syndrome
- Hand-Schuller-Christian (Histiocytosis X0)
- Hallgren syndrome
- Herpes-Zoster (or Hunt)
- Hunter syndrome (MPS II)
- Hurler syndrome (MPS I-H)
- Kearns-Sayre syndrome
- Kippel-Feil sequence
- Kippel-Trenaunay-Weber syndrome
- Kniest Dysplasia
- Leber congenital amaurosis
- Leigh Disease
- Marfan syndrome
- Marshall syndrome
- Maroteaux-Lamy syndrome (MPS-VI)
- Moebius syndrome
- Monosomy 10p
- Morquio syndrome (MPS IV-B)
- NF1-Neurofibromatosis (von Recklinghausen disease)
- NF2-Bilateral Acoustic Neurofibromatosis
- Norrie Disease
- Optico-Cochleo-Dentate Degeneration
- Pfeiffer syndrome
- Prader-Willi
- Pierre-Robin syndrome
- Refsum syndrome
- Scheie syndrome (MPS I-S)
- Smith-Lemli-Opitz (SLO) syndrome
- Stickler syndrome
- Sturge-Weber syndrome
- Treacher Collins syndrome
- Trisomy 13 (Trisomy 13-15, Patau syndrome)
- Trisomy 18 (Edwards syndrome)
- Turner syndrome
- Usher I syndrome
- Usher II syndrome
- Usher III syndrome
- Vogt-Koyanagi-Harada syndrome
- Waardenburg syndrome
- Wildervanck syndrome
- Wolf-Hirschhorn syndrome (Trisomy 4p)
- Other

(specify in space provided)

Pre-Natal/Congenital Complications

- Congenital Rubella
- Congenital Syphilis
- Congenital Toxoplasmosis
- Cytomegalovirus (CMV)
- Fetal Alcohol syndrome
- Hydrocephaly
- Maternal Drug Use
- Microcephaly
- Neonatal Herpes Simplex (HSV)
- Other

(specify in space provided)

Post-Natal/Non-Congenital Complications

- Asphyxia
- Direct Trauma to the eye and/or ear
- Encephalitis
- Infections
- Meningitis
- Severe Head Injury
- Tumors
- Stroke
- Chemically Induced
- Other

(specify in space provided)

Related to Prematurity

- Complications of Prematurity

Undiagnosed

- No Determination of Etiology

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Educational Setting: (Ages 3-21)

ECSE (3-5) Settings:

- Attending a regular early childhood program at least 80% of the time
- Attending a regular early childhood program 40% to 79% of the time
- Attending a regular early childhood program less than 40% of the time
- Attending a separate class
- Attending a separate school
- Attending a residential facility
- Service provider location
- Home

School Aged (6 – 21) Settings:

- Inside the regular class 80% or more of day
- Inside the regular class 40% to 79% of day
- Inside the regular class less than 40% of day
- Separate school
- Residential facility
- Homebound/Hospital
- Correctional facilities
- Parentally placed in nonpublic schools (private)

Participation in Statewide Assessments:

- Regular grade-level assessment
- Regular grade-level assessment with accommodations
- Alternate assessments aligned with standards
- Not yet required at age or grade level

Special Education Status (Exiting Part C):

- In a Part C early intervention program
- Completion of IFSP prior to reaching maximum age for Part C
- Eligible for IDEA, Part B
- Not eligible for Part B, exit with referrals to other programs
- Not eligible for Part B, exit with no referrals
- Deceased
- Moved out of state
- Withdrawal by parent (or guardian)
- Part B eligibilty not determined

Special Education Status (Exiting Part B):

- In ECSE or school-aged special education program
- Transferred to regular education
- Graduation with regular diploma
- Received a certificate
- Reached maximum age
- Moved, known to be continuing
- Dropped Out
- Died
- No longer receives Sp. Ed., but still receiving State DB Project Services

Living Setting:

- Home: With Parents
- Home: Extended Family
- Home: Foster parents
- State Residential Facility
- Private Residential Facility
- Group Home (less than 6 residents)
- Group Home (6 or more residents)
- Apartment (with non-family person(s))
- Pediatric Nursing Home
- Other _____

(Check one for each area):

Corrective Lens:

- Yes
- No
- Unknown

Assistive Listening Devices:

- Yes
- No
- Unknown

Additional Assistive Technology:

- Yes
- No
- Unknown

Intervener Services:

- Yes
- No
- Unknown

Contact Person:

Name: _____

Title: _____

School District: _____

Address: _____

City: _____ Zipcode: _____

Date Completed: ____/____/____

Telephone: _____

Email: _____

Return Form To:

Teresa Coonts, Director
Nebraska Deaf-Blind Project
301 Centennial Mall South
Lincoln, NE 68509
Teresa.coonts@nebraska.gov