**ELIGIBILITY APPLICATION PART II: ADDITIONAL STUDENT INFORMATION**

Student: Date: Contacted by:

1. **PRIMARY IDENTIFIED ETIOLOGY** **To be completed by VSAP personnel. Circle only one.**

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| **Hereditary/Chromosomal Syndromes and Disorders** | |
| 101 Aicardi Syndrome | 130 Marshall Syndrome |
| 102 Alport Syndrome | 131 Maroteaux-Lamy Syndrome (MPS VI) |
| 103 Alstrom Syndrome | 132 Moebius Syndrome |
| 104 Apert Syndrome (Acrocephalosyndactyly, Type 1) | 133 Monosomy 10p |
| 105 Bardet-Biedl Syndrome (Laurence Moon-Biedl) | 134 Morquio Syndrome (MPS IV-B) |
| 106 Batten Disease | 135 NF 1 - Neurofibromatosis-(von Recklinghausen Disease) |
| 107 CHARGE Association | 136 NF 2 - Bilateral Acoustic Neurofibromatosis |
| 108 Chromosome 18, Ring 18 | 137 Norrie Disease |
| 109 Cockayne Syndrome | 138 Optico-Cochleo-Dentate Degeneration |
| 110 Cogan Syndrome | 139 Pfieffer Syndrome |
| 111 Cornelia de Lange | 140 Prader-Willi |
| 112 Cri du Chat Syndrome (Chromosome 5p Syndrome) | 141 Pierre-Robin Syndrome |
| 113 Crigler-Najjar 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 (Trisomy 13-15, Patau Syndrome) |
| 120 Herpes-Zoster (or Hunt) | 149 Trisomy 18 (Edwards Syndrome) |
| 121 Hunter Syndrome (MPS II) | 150 Turner Syndrome |
| 122 Hurler Syndrome (MPS I-H) | 151 Usher I Syndrome |
| 123 Kearns-Sayre Syndrome | 152 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 (describe) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ |

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| **Pre-Natal/Congential Complications** | **Post-Natal/Non-Congenital Complications** |
| 201 Congenital Rubella | 301 Asphyxia |
| 202 Congenital Syphilis | 302 Direct Trauma to the Eye and/or Ear |
| 203 Congenital Toxoplasmosis | 303 Encephalitis |
| 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 |

**B. PART C EXITING STATUS (BIRTH THROUGH 2)**  Please **check one**.

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| 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, exit with referrals to other  programs  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 (or guardian)  9. Attempts to contact the parent and/or child were  unsuccessful |

**C. PART B EXITING** Please **check one**. NOTE: Item 7 is intentionally not used or available.

|  |  |
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| 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 |  |
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**D. DEAF-BLIND PROJECT EXITING STATUS** Please **check one**.

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| 0. Eligible to receive services from the deaf-blind  project | 1. No longer eligible to receive services from the  state deaf-blind project |

**E. NOTES**