

# 2023 California Deafblind Census

Return to: California Deafblind Services, SF State University  
Department of Special Education, 1600 Holloway Avenue  
San Francisco, CA 94132-4158 | (415) 405-7560

Check box if there are NO CHANGES from last year --->

<b>Name</b>			<b>Birth Date</b>			<b>Gender</b>		<b>Box below for staff use only.</b>	
First	MI	Last	Month	Day	Year	<input type="radio"/> Male	<input type="radio"/> Female	<input type="radio"/> Other/Non-Binary	created <input type="text"/>
<b>County of Residence</b>			County #			modified <input type="text"/>	ID Code <input type="text"/>	Kid Code <input type="text"/>	Child Code <input type="text"/>
<b>Guardian(s)</b>			<b>City</b>						
<b>Guardian 2 city if other address</b>			<b>City</b>			<b>State</b>			

**Ethnicity:** Please answer BOTH Hispanic origin & Race for each

**Hispanic/Latino:** Includes all individuals who identify with one or more nationalities or ethnic groups originating in Mexico, Puerto Rico, Cuba, Central and South America, and other Spanish cultures, regardless of race. If a person is not of Hispanic, Latino, or Spanish origin, answer "No, not Hispanic, Latino". **Race:** The general racial category that most clearly reflects individuals' recognition of their community or with which the individual most identifies

**Pt 1: Is this student Hispanic/Latino?**  0 No, not Hispanic/Latino  1 Yes, Hispanic/Latino

**Pt 2 RACE:** If more than 1, select #7 "Two or More"

1 American Indian or Alaska Native  2 Asian  3 Black or African American  5 White  6 Native Hawaiian or Other Pacific Islander  7 Two or more  999 Unknown/Missing

**Primary Language**

1 English  2 Spanish  3 Sign Language  9 Arabic  9 Cambodian  9 Cantonese  9 Chinese  9 Gujarati  9 Hmong  9 Ibo  9 Japanese  9 Khmer  9 Korean  9 Mandarin  9 Non-verbal  9 Punjabi  9 Pashto  9 Romanian  9 Russian  9 Tagalog  9 Tamil  9 Telugu  9 Urdu  9 Vietnamese  9 Other\*  999 Unknown

<b>ETIOLOGY</b> <i>specify ONE only, from one of the 5 subsections:</i>	<b>Hereditary/Chromosomal Syndromes and Disorders</b>	<b>Pre-Natal/Congenital Complications</b>	
	<input type="radio"/> 101 Aicardi syndrome <input type="radio"/> 102 Alport syndrome <input type="radio"/> 103 Alstrom syndrome <input type="radio"/> 104 Apert syndrome (Acrocephalosyndactyly) <input type="radio"/> 105 Bardet-Biedl syndrome (Laurence Moon-Biedl) <input type="radio"/> 106 Batten disease <input type="radio"/> 107 CHARGE syndrome <input type="radio"/> 108 Chromosome 18, Ring 18 <input type="radio"/> 109 Cockayne syndrome <input type="radio"/> 110 Cogan syndrome <input type="radio"/> 111 Cornelia de Lange <input type="radio"/> 112 Cri du chat syndrome (Chromosome 5p) <input type="radio"/> 113 Crigler-Najjar syndrome <input type="radio"/> 114 Crouzon syndrome (Craniofacial Dysostosis) <input type="radio"/> 115 Dandy Walker syndrome <input type="radio"/> 116 Down syndrome (Trisomy 21) <input type="radio"/> 117 Goldenhar syndrome <input type="radio"/> 118 Hand-Schuller-Christian (Histiocytosis X) <input type="radio"/> 119 Hallgren syndrome <input type="radio"/> 120 Herpes-Zoster (or Hunt) <input type="radio"/> 121 Hunter syndrome (MPSII) <input type="radio"/> 122 Hurler syndrome (MPS I-H) <input type="radio"/> 123 Kearns-Sayre syndrome <input type="radio"/> 124 Klippel-Feil sequence <input type="radio"/> 125 Klippel-Trenaunay-Weber syndrome <input type="radio"/> 126 Kniest Dysplasia <input type="radio"/> 127 Leber congenital amaurosis <input type="radio"/> 128 Leigh disease <input type="radio"/> 129 Marfan syndrome	<input type="radio"/> 130 Marshall syndrome <input type="radio"/> 131 Maroteaux-Lamy syndrome (MPS VI) <input type="radio"/> 132 Moebius syndrome <input type="radio"/> 133 Monosomy 10p <input type="radio"/> 134 Morquio syndrome (MPS IV-B) <input type="radio"/> 135 NF1 - Neurofibromatosis <input type="radio"/> 136 NF2- Bilateral Acoustic Neurofibromatosis <input type="radio"/> 137 Norrie disease <input type="radio"/> 138 Optico-Cochleo-Dentate Degeneration <input type="radio"/> 139 Pfeiffer syndrome <input type="radio"/> 140 Prader-Willi <input type="radio"/> 141 Pierre-Robin syndrome <input type="radio"/> 142 Refsum syndrome <input type="radio"/> 143 Scheie syndrome (MPS I-S) <input type="radio"/> 144 Smith-Lemli-Optiz (SLO) syndrome <input type="radio"/> 145 Stickler syndrome <input type="radio"/> 146 Sturge-Weber syndrome <input type="radio"/> 147 Treacher Collins syndrome <input type="radio"/> 148 Trisomy 13 (Patau syndrome) <input type="radio"/> 149 Trisomy 18 (Edwards syndrome) <input type="radio"/> 150 Turner syndrome <input type="radio"/> 151 Usher I syndrome <input type="radio"/> 152 Usher II syndrome <input type="radio"/> 153 Usher III syndrome <input type="radio"/> 154 Vogt-Koyanagi-Harada syndrome <input type="radio"/> 155 Waardenburg syndrome <input type="radio"/> 156 Wilderervanck syndrome <input type="radio"/> 157 Wolf-Hirschhorn syndrome (Trisomy 4p) <input type="radio"/> 199 Other hereditary *	<input type="radio"/> 201 Congenital Rubella <input type="radio"/> 202 Congenital Syphilis <input type="radio"/> 203 Congenital Toxoplasmosis <input type="radio"/> 204 Cytomegalovirus (CMV) <input type="radio"/> 205 Fetal Alcohol syndrome <input type="radio"/> 206 Hydrocephaly <input type="radio"/> 207 Maternal drug use <input type="radio"/> 208 Microcephaly <input type="radio"/> 209 Neonatal Herpes Simplex (HSV) <input type="radio"/> 299 Other pre-natal *
	<input type="radio"/> 130 Marshall syndrome <input type="radio"/> 131 Maroteaux-Lamy syndrome (MPS VI) <input type="radio"/> 132 Moebius syndrome <input type="radio"/> 133 Monosomy 10p <input type="radio"/> 134 Morquio syndrome (MPS IV-B) <input type="radio"/> 135 NF1 - Neurofibromatosis <input type="radio"/> 136 NF2- Bilateral Acoustic Neurofibromatosis <input type="radio"/> 137 Norrie disease <input type="radio"/> 138 Optico-Cochleo-Dentate Degeneration <input type="radio"/> 139 Pfeiffer syndrome <input type="radio"/> 140 Prader-Willi <input type="radio"/> 141 Pierre-Robin syndrome <input type="radio"/> 142 Refsum syndrome <input type="radio"/> 143 Scheie syndrome (MPS I-S) <input type="radio"/> 144 Smith-Lemli-Optiz (SLO) syndrome <input type="radio"/> 145 Stickler syndrome <input type="radio"/> 146 Sturge-Weber syndrome <input type="radio"/> 147 Treacher Collins syndrome <input type="radio"/> 148 Trisomy 13 (Patau syndrome) <input type="radio"/> 149 Trisomy 18 (Edwards syndrome) <input type="radio"/> 150 Turner syndrome <input type="radio"/> 151 Usher I syndrome <input type="radio"/> 152 Usher II syndrome <input type="radio"/> 153 Usher III syndrome <input type="radio"/> 154 Vogt-Koyanagi-Harada syndrome <input type="radio"/> 155 Waardenburg syndrome <input type="radio"/> 156 Wilderervanck syndrome <input type="radio"/> 157 Wolf-Hirschhorn syndrome (Trisomy 4p) <input type="radio"/> 199 Other hereditary *	<b>Post-Natal/Non-Congenital Complications</b> <input type="radio"/> 301 Asphyxia <input type="radio"/> 302 Direct Trauma to the eye and/or ear <input type="radio"/> 303 Encephalitis <input type="radio"/> 304 Infections <input type="radio"/> 305 Meningitis <input type="radio"/> 306 Severe head injury <input type="radio"/> 307 Stroke <input type="radio"/> 308 Tumors <input type="radio"/> 309 Chemically induced <input type="radio"/> 399 Other post-natal *	

<b>Documented Vision Loss</b>	<b>Cortical Vision Impairment (CVI)</b>	<b>Documented Hearing Loss</b>	<b>Central Auditory Processing disorder</b>
<input type="radio"/> 1 Low Vision (20/70 to 20/200) <input type="radio"/> 2 Legally Blind (20/200) or Field Restricted <input type="radio"/> 3 Light Perception Only <input type="radio"/> 4 Totally Blind <input type="radio"/> 6 Diagnosed Progressive Loss <input type="radio"/> 7 Further Testing Needed <input type="radio"/> 9 Documented Functional Vision Loss	<input type="radio"/> 0 No <input type="radio"/> 1 Yes <input type="radio"/> 2 Unknown <b>Corrective Lenses</b> <input type="radio"/> 0 No <input type="radio"/> 1 Yes <input type="radio"/> 2 Unknown	<input type="radio"/> 1 Mild (26-40dB loss) <input type="radio"/> 2 Moderate (41-55dB loss) <input type="radio"/> 3 Moderately Severe (56-70dB loss) <input type="radio"/> 4 Severe (71-90dB loss) <input type="radio"/> 5 Profound (91+ loss) <input type="radio"/> 6 Diagnosed Progressive Loss <input type="radio"/> 7 Further Testing Needed <input type="radio"/> 9 Documented Functional Hearing Loss	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> 2 Unknown <input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> 2 Unknown <input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> 2 Unknown <input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> 2 Unknown

<b>Other Impairments or Conditions</b>	<b>Orthopedic / Physical</b>	<b>Cognitive</b>	<b>Behavioral</b>	<b>Complex Health Care</b>	<b>Communication, Speech or Language</b>	<b>Other Impairments*</b>	<b>* If Other, specify:</b>
	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	<input type="radio"/> 0 No <input type="radio"/> 1 Yes	

**Intervener and One-on-One Services:** Does the student receive One-on-One Support from someone with the function or title of an Intervener?  0 = No  1 = Yes  2 = Unknown  888 = N/A

Intervener services provide access to information and communication and facilitate the development of social and emotional well-being for children who are deafblind. In educational environments, intervener services are provided by an individual, typically a paraeducator, who has received specialized training in deafblindness and the process of intervention. An intervener provides consistent one-to-one support to a student who is deafblind (age 3 through 21) throughout the instructional day.

# 2023 California Deafblind Census

Check box if there are NO CHANGES from last year --->

## IDEA Funding / Educational Placement: Must Correlate to the Child's Age

DOB:

Funding Category →  1 IDEA Part B (ages 3-21)  2 IDEA Part C (ages birth - 2)  3 Not reported under Parts B or C

### Part B Category *specify ONE only*

- 1 Intellectual Disability
- 2 Hearing Impairment (includes deafness)
- 3 Speech or Language Impairment
- 4 Visual Impairment (includes blindness)
- 5 Emotional Disturbance
- 6 Orthopedic Impairment
- 7 Other Health Impairment
- 8 Specific Learning Disability
- 9 Deaf-blindness
- 10 Multiple Disabilities
- 11 Autism
- 12 Traumatic Brain Injury
- 13 Developmental Delay
- 14 Non-Categorical
- 888 Not Reported under Part B of IDEA
- 999 Unknown

### Educational Setting (ages 3-21) *specify ONE only*

- ages 3 - 5
- 301 Services in Regular Early Childhood Program (10+ hours)
  - 302 Other Location Regular Early Childhood Program (10+ hours)
  - 303 Services in Regular Early Childhood Program (<10 hours)
  - 304 Other Location Regular Early Childhood Program (<10 hours)
  - 305 Attending a Separate Class
  - 306 Attending a Separate School
  - 307 Attending a Residential Facility
  - 309 Home, at public expense
  - 310 Home, not at public expense
  - 888 N/A Not Served Under Part B
  - 999 Unknown/Missing

- ages 6 - 21
- 610 Inside the regular class 80% or more of day
  - 611 Inside the regular class 40% to 79% of day
  - 612 Inside the regular class less than 40% of day
  - 613 Separate school
  - 614 Residential facility
  - 615 Homebound/Hospital
  - 616 Correctional facilities
  - 617 Parentally placed in private schools
  - 620 Home school/remote learning, at public expense
  - 621 Home school/remote learning, NOT at public expense
  - 888 N/A Not Served Under Part B
  - 999 Unknown/Missing

### Part B Exiting Status *specify ONE only*

- 0 Not exited — In Special Education Program
- 1 Transferred to regular education
- 2 Graduated with regular diploma
- 22 Graduated with alternate diploma
- 3 Received a certificate
- 4 Reached Maximum Age
- 5 Died
- 6 Moved, Known to be Continuing\*
- 8 Dropped out

### \*Known info. on students who have moved:

### Participation in Statewide Assessments: *specify most recent one only*

- 1 Regular grade-level State assessment
- 2 Regular grade-level State assessment w/accommodations
- 3 Alternate assessments aligned with grade-level achievement standards
- 6 Not required at age or grade level
- 7 Parent Opt-Out

### Deaf-Blind Project *FOR STAFF USE ONLY. PLEASE LEAVE BLANK.*

Currently eligible to receive services from the deaf-blind project? 0=YES, 1=NO

### Part C Category — (Ages Birth to 2 yrs) *specify ONE only*

- 1 At-risk for Developmental Delays (as defined by the state's Part C Lead Agency)
- 2 Developmentally Delayed
- 888 N/A Not reported under Part C of IDEA

### Part C Exiting Status *specify ONE only*

- 0 = Not Exited - Currently in 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
- 888 = NA Not Part C

### Early Intervention Setting

- 1 Home  3 Other \*
- 2 Community-based  888 NA / Not under Part C

\* If "Other", please specify:

### Placement: School or Site where services are received

School/Site Name

---

Street Address

---

City Zip Code

---

Phone Fax

---

Teacher Teacher Phone

---

Teacher Email

---

Co. # District # School #

---

District/LEA Name

---

District Type

---

School Type

## Best Service Provider Contact for the Child OR Person Completing Form to Contact

.....  
Contact Name

.....  
Phone 1 Fax

.....  
Title/Position

.....  
Phone 2

.....  
Organization/Agency

.....  
e-mail address

.....  
Street Address

.....  
Signature - (Please also print name if different from Contact Name)

.....  
City Zip Code

.....  
Date