The Individuals with Disabilities Education Act (IDEA) ensures states and public agencies provide early intervention, special education, and related services to more than 7.2 million eligible children and youth with disabilities across the country, including infants and toddlers. Programs for young children with disabilities play a significant role in early childhood education and development, which is particularly salient when it comes to promoting school readiness. The connections in a baby’s brain are most adaptable in the first three years of life. These connections, which are the foundation for learning, behavior, and health, become harder to change over time. Ensuring that young children have access to inclusive early interventions during this critical period of development means that children will get the appropriate services and supports they need to be equipped for success in school and avoid costly interventions later in life.

**Background**

Since 1975, the federal government has sought to uphold the fundamental principle of educational equality for children with disabilities through a permanent, broad-scale federal assistance program. IDEA provides federal funding for the education of children with disabilities and, as a condition of the funding, requires states to provide a free appropriate public education specifically

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**At a Glance**

- **Early Intervention Programs Created:**
  - 1986

- **Who this Program Serves:**
  - Children birth through age 5 with established physical or mental conditions.

- **Number of Children Served in the 2020-2021 School Year:**
  - Part C: 786,275 children
  - Part B, Sec. 619: 502,391 children

- **Delivery Method:**
  - Part C provides early intervention services to infants and toddlers with disabilities and their families.
  - Part B, Sec. 619 provides special education and related services to children ages 3 through 5.

- **FY 2023 Federal Funding:**
  - Part C: $540 million;
  - Part B, Sec. 619: $420 million

Since its inception, IDEA has never been fully federally funded. Currently, the federal contribution is around 14%, meaning states are required to make up a significant portion of the funding. There have been legislative efforts to fully fund at 40%, however, they have not yet been realized.

1 U.S. Department of Education (Part C; Part B, Sec. 619)
designed to meet the needs of children with disabilities at no cost to their parents. Recent data from the Centers for Disease Control and Prevention found that approximately one in six children in the United States have a developmental disability ranging from mild disabilities, such as speech and language impairments, to serious developmental disabilities, such as intellectual disabilities, cerebral palsy, and autism.

**Early Intervention and Preschool Special Education Programs**

IDEA Part B covers special education for children and youth with disabilities between the ages of three and twenty-one. IDEA also contains two programs specifically designed to serve younger children with disabilities: IDEA Part C, which authorizes federal funding for early intervention services to infants and toddlers with disabilities ages birth to three years; and IDEA Part B, Section 619, which authorizes supplementary grants to states for preschool programs serving children with disabilities ages three through five.

Part C aids states in creating and maintaining “a statewide, comprehensive, coordinated, multidisciplinary, interagency system that provides early intervention services for infants and toddlers with disabilities and their families.” Services focus on children experiencing, or who have a high probability of experiencing, “developmental delay” and are outlined for each child and his or her family in an Individualized Family Service Plan (IFSP). Early intervention services may include audiology, vision, speech-language pathology, and psychological services; physical therapy; provision of assistive technology devices; and social work, referral, and coordination services. To the maximum extent feasible, services are to be provided in “natural environments,” including the home, with other infants and toddlers who are not disabled. This element of inclusion helps children and their families feel like important and accepted members of their community and leads to positive social relationships and friendships, as well as development and learning to reach their full potential. If the developmental delay has not been resolved by age 3, a child may continue receiving services under Part B, Sec. 619.