Brief History of the IDEA

The federal law that supports special education and related service programming for children and youth with disabilities is called the **Individuals with Disabilities Education Act (IDEA)**, [formerly the Education for the Handicapped Act (EHA)]. Public Law 94-142 (the Education of All Handicapped Children Act), which was originally enacted in 1975 to establish grants to States for the education of children with disabilities, has been amended several times. Under Part B of the IDEA, all eligible school-aged children and youth with disabilities are entitled to receive a free appropriate public education (FAPE).

In 1986, the EHA was amended by P.L. 99-457 to provide special funding incentives for States that would make a free appropriate public education available for all eligible preschool children with disabilities ages three through five. Provisions were also included to help States develop early intervention programs for infants and toddlers with disabilities; this part of the legislation has become known as the Part H program.

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The EHA was amended again in 1990 by P.L. 101-476, which, among other things, changed the name of the legislation to the **Individuals with Disabilities Education Act (IDEA)**, or IDEA. The IDEA has been amended once, by P.L. 102-119.

The IDEA is an important federal law, because it requires that a free appropriate public education, which includes special education and related services, be available to children and youth with disabilities in mandated age ranges. This **News Digest** provides a general overview of the IDEA and its regulations as they relate to school-aged children. The **News Digest** is intended to help you understand the law and how it mandates services for your school-aged child with a disability. Information about services available to infants and toddlers through the Part H program and for preschool children with disabilities is presented in a separate NICHCY document entitled **A Parent's Guide: Accessing Programs for Infants, Toddlers, and Preschoolers With Disabilities**.

The information provided in this issue of the **News Digest** is drawn specifically from the IDEA, as passed by Congress in 1990 and codified into law under 20 United States Code (USC), **Chapter 33**. Final regulations for this Act were published in the **Federal Register** on September 29, 1992, with a correction appearing in the **Federal Register** on October 27, 1992. These final regulations replace those developed for the EHA (authority: 20 U.S.C. §1401, 1411-1420) and are currently guiding school systems in how they design and implement their special education and related services programs.

**Obtaining a Copy of the IDEA and Its Regulations**

Because states base their programs upon the law and its implementing Federal regulations, it is useful for parents to read and become familiar with the law itself. Generally speaking, you can obtain the latest copy of Federal regulations for laws through the U.S. Government Printing Office. At the time of this writing, however, one unified version of the regulations for the IDEA has not been compiled.
Part II:

Purposes and Promises of the IDEA—Who is Eligible for Services and What Types of Services Are Provided?

What is the purpose of the IDEA?

The major purposes of the IDEA are:

- to ensure that all children with disabilities have available to them a “free appropriate public education” that includes special education and related services designed to meet their unique needs;
- to ensure that the rights of children and youth with disabilities and their parents are protected;
- to assist states and localities in providing for the education of all children and youth with disabilities;
- and
- to assess and ensure the effectiveness of efforts to educate children with disabilities. (§300.1)

What is a free appropriate public education?

Under the law, a free appropriate public education (FAPE) means special education and related services that:

- are provided to children and youth with disabilities at public expense, under public supervision and direction, and without charge;
- meet the standards of the State Education Agency (SEA), including the requirements of the IDEA;
- include preschool, elementary, or secondary school education in the State involved; and
- are provided in conformity with requirements for the individualized education program. (§300.8)

There requirements for an individualized education program (IEP) are found in §300.340 - §300.350 and are discussed in greater detail in Part IV of this document.

Who is eligible for services under the IDEA?

The regulations for IDEA define “children with disabilities” as including children (a) who have been evaluated in accordance with IDEA’s evaluation requirements (§300.530 - §300.534) (see page 4 in this document); (b) who have been determined, through this evaluation, to have one or more of the disabilities listed below; and (c) who, because of their disability, need special education and related services. The disabilities listed by IDEA are:

- mental retardation;
- hearing impairments, including deafness;
- speech or language impairments;
- visual impairments, including blindness;
- serious emotional disturbance;
- orthopedic impairments;
- autism;
- traumatic brain injury;
- other health impairments;
- specific learning disabilities;
- deaf-blindness; or
- multiple disabilities. (§300.7)

Attachment A of this News Digest provides the complete description of each of these disabilities, as contained in the regulations for IDEA.

For children aged 3-5, “children with disabilities” may include, at the State’s discretion, those children who are experiencing developmental delays, as defined by the State and as measured by appropriate diagnostic instruments and procedures, in one or more of the following areas:

- cognitive development;
- physical development;
- communication development;
• social or emotional development; or
• adaptive development; and
• who need, for that reason, special education and related services. (§300.7)

From birth through age 2, children may be eligible for services through The Infants and Toddlers with Disabilities Program (Part H) of the IDEA. This News Digest does not discuss the Part H program. For information about how to access services for infants and toddlers, contact NICHCY and request a copy of A Parent's Guide: Accessing Programs for Infants, Toddlers, and Preschoolers With Disabilities.

What is special education?

Special education is defined as specially designed instruction, at no cost to you, to meet your child's or youth's unique needs. Special education can include classroom instruction, home instruction, instruction in hospitals and institutions, or other settings. It can also include instruction in physical education and vocational education. (§300.17)

Special education also includes speech pathology or any other related service, if the service consists of instruction that is specially designed, at no cost to you, to meet your child's unique needs and is considered by the State as special education rather than a related service. (§300.17)

Where is special education instruction provided?

As listed above, special education instruction can be provided in a number of settings, such as: instruction in regular classrooms, special classrooms, special schools, home instruction, instruction in hospitals and institutions, and instruction in other settings (§300.17). Schools must ensure that a continuum of alternative placements is made available to meet the needs of children with disabilities for special education and related services (§300.551(b)(1)). Unless a child's individualized education program (IEP) requires some other arrangement, the child must be educated in the school he or she would attend if he or she did not have a disability (§300.552(c)).

Special education instruction must also be provided to students with disabilities in what is known as the Least Restrictive Environment, or LRE. Both the IDEA and its regulations have provisions that require that children with disabilities are educated with nondisabled children, to the maximum extent appropriate. The IDEA'S LRE requirements apply to students in public or private institutions or other care facilities as well (§300.550(b)(1)). Each State must further ensure that special classes, separate schooling, or other removal of children with disabilities from the regular educational environment occurs only when the nature or severity of the disability is such that education in regular classes with the use of supplementary aids and services cannot be achieved satisfactorily (§300.550(b)(2)).

What are related services?

Related services are defined in the regulations as transportation and such developmental, corrective, and other supportive services as are required to assist a child with a disability to benefit from special education. Related services may include:

• audiology;
• psychological services;
• physical therapy;
• occupational therapy;
• medical services for diagnostic or evaluation purposes only;
• school health services;
• recreation, including therapeutic recreation;
• counseling services, including rehabilitation counseling;
• early identification and assessment of disabilities in children;
• social work services in schools;
• transportation;
• speech pathology; and
• parent counseling and training. (§300.16)

The list of related services identified in the IDEA regulations is not intended to be exhaustive and could include other developmental, corrective, or support services if they are required to assist a child with a disability to benefit from special education.

You should inform yourself fully about the related services that are listed (and described in some detail) in the law. NICHCY's News Digest called Related Services for School-Aged Children can also provide information and guidance about related services. This News Digest (Volume 1, Number 2, 1991) is available free by contacting NICHCY.
Part III: 
Accessing Services

What is the first step in obtaining special education and related services?

Before a child can receive special education and related services for the first time, an initial evaluation of the child, also known as a preplacement evaluation, must be conducted (§300.531). This evaluation must meet the requirements stated in §300.532 of the IDEA's regulations. Parental permission—that is, written consent—must be obtained before this evaluation can be conducted (§300.504(b)(1)(i)).

The law requires that no single procedure he used as the sole criterion for determining an appropriate educational program for a child...

Further, the child must be assessed in all areas related to the suspected disability.

How do I receive an evaluation of my child?

There are at least two ways in which your child may be selected to receive an evaluation:

(1) You may request that your child be evaluated. You can call or write to the Director of Special Education or the Principal of your child's school. If the school district suspects that your child has a disability, your child must be evaluated. If the school refuses to evaluate your child because it does not suspect that your child has a disability, you must be given written notice of the refusal and a full explanation that explains the reasons for the refusal [§300.504(a)(2) and §300.505(a)(2)]. This notice must also give you a full explanation of the procedural safeguards available to parents under the Individuals with Disabilities Education Act, including your right to challenge the school district's refusal by requesting an impartial due process hearing [§300.505(a)(1)].

(2) The school may initiate a request to evaluate your child. Based on a teacher's recommendation, or observations or results from tests given to all children in a particular grade, a school may recommend a child for further screening or assessment to determine if he or she needs special education and related services. If your child is thought to need further testing, the school must tell you. You, as a parent, must give written permission before the school can evaluate your child for the first time [§300.504(b)(1)].

It is important for you to know that the law requires that parents be notified in writing before the school district evaluates their child for the first time. All written communication from the school must be in a form that parents can understand (e.g., in their native language if they do not read English, or in the mode of communication normally used by the individual, such as Braille or large print). It must state the action that is proposed or refused; the reasons for the proposal or refusal; the evaluation procedures, tests, or records used to support the proposal or refusal; and an explanation of the parents' rights if they should disagree with the suggested actions or refusal to act, as well as any other options the agency considered and the reasons why those options were rejected (§300.505). Parents must give their consent in writing before an initial evaluation can be conducted. It is important to know that school districts must have procedures by which they can override a parental refusal of consent, which they may use if they believe a child should be evaluated. These are discussed on page 10.

What does the evaluation process involve?

The regulations are very specific about how school districts must conduct evaluations of children and youth thought to have a disability. For one, evaluations must be conducted by a group of persons, called a multidisciplinary team. This team must include at least one teacher or other specialist who is knowledgeable about the area of the child's suspected disability [§300.532(e)]. Thus, the multidisciplinary team conducting the assessment may involve, at the State's discretion, professionals such as the following: school psychologist, speech and language pathologist, occupational or physical therapist, adaptive physical education therapist, medical specialists, educational diagnosticians, classroom teachers, or others.

The law also requires that no single procedure be used as the sole criterion for determining an appropriate educational program for a child (§300.532(d)). Further, the child must be assessed in all areas related to the suspected disability. This includes, where appropriate, evaluating a child's:

- health;
- vision;
- hearing;
- social and emotional status;
- general intelligence;
- academic performance;
- communicative status; and
- motor abilities. [§300.532(b)]

School districts must also be sure that the tests they select and use are not racially or culturally discriminatory (biased) against the child [§300.530(b)]. This provision of the law is meant to protect children of different racial or cultural backgrounds from misdiagnosis. For example, children's cultural backgrounds may affect their behavior or test responses in ways that teachers or other personnel do not understand. Similarly, if a child speaks another language or has limited English
proficiency, he or she may not understand directions or words on tests and may be unable to respond correctly. As a result, a child may mistakenly appear to be a slow learner or to have a hearing or communication problem. In order to prevent such mistakes, the law states that tests and other evaluation materials given to a child during the evaluation process must be provided and administered in the child's native language or through another mode of communication, unless it is clearly not feasible to do so [§300.532(a)(1)].

In keeping with the requirements mentioned above, then, the multi-disciplinary team must collect information about your child from a variety of sources, which could include you, the parent. In addition to testing, the evaluation process may include:

- observations by professionals who have worked with your child;
- your child's medical history, when it is relevant to his or her performance in school; and
- information and observations from you about your child's school experience, abilities, needs, and behavior outside of school, and his or her feelings about school.

What happens if I don't agree with the evaluation results?

If parents of a child with a disability disagree with the results of the evaluation performed for or by the school, they have the right to obtain an Independent Educational Evaluation (IEE) (§300.503). This means that you can request that a qualified professional examiner or qualified evaluation clinic conduct another evaluation of your child. The examiner may not be employed by or involved in any way with the school. School districts generally maintain a list of qualified examiners and clinics who meet the State's standards, because they are required to tell parents, upon request, where an IEE may be obtained [§300.503(a)(2)]. If the public agency maintains such a list, parents are not limited to the examiners on the list, if they can demonstrate that unique circumstances justify selection of another evaluator.

Who pays for the independent evaluation? The answer is that some IEEs are at public expense and others are paid for by the parents. For example, if you are the parent of a child with a disability and you disagree with the school's evaluation, you may request an IEE at public expense. The school district may grant this request and pay for the IEE, or it may initiate a hearing to show that its own evaluation was appropriate. If the final decision of the hearing officer is that the school's evaluation was appropriate, then you still have the right to an IEE but not at public expense [§300.503(b)]. As part of a hearing, a hearing officer may also request an IEE; if so, that IEE must be at public expense. Of course, you have the right to arrange to have your child independently evaluated at any time at your own expense. (Note: When the same tests are repeated within a short time period, the validity of the results can be seriously weakened.) Whenever an IEE is publicly funded, that IEE must meet the same criteria that the school district uses when it initiates an evaluation.

Regardless of who pays for the IEE, the school district is obligated to consider the results of this evaluation when making any decisions about the educational program of the child in question. They are not required to invite the parent to this meeting — often called an eligibility meeting. However, if you do attend, your child's assessment results should be explained. The specialists who assessed your child should explain what they did, why they used the tests they administered, the results of your child's tests, and what your child's scores mean when compared to other children of the same age and in the same grade. Regardless of whether you attend the meeting, you must be notified as to its outcome.

If the evaluation results indicate that your child meets the definitions of one or more of the disabilities listed under IDEA (see page 13) and needs special education and related services, the results will form the basis for development of your child's Individualized Education Program (IEP).

Will my child be evaluated again in the future?

In most cases, yes. The first time your child is evaluated is called a preplacement evaluation (§300.531). Evaluations must also be conducted at least every three years (called a triennial re-evaluation) after your child has been placed in special education (§300.534(b)). Re-evaluations can also occur more frequently if conditions make it necessary, or if you or your child's teacher request(s) a re-evaluation (§300.534(b)).
Part IV:
The Individualized Education Program

What is an Individualized Education Program (IEP)?

An IEP is a written statement of the educational program that is designed to meet a child's unique needs. Among the purposes of the IEP are:

- to establish learning goals for the child; and
- to state the services that the school district will provide.

The law requires that every child receiving special education services have an IEP (§300.342). The IEP must include statements about:

- the child's current levels of educational performance. This may include information concerning his or her: academic achievement, social adaptation, prevocational and vocational skills, sensory and motor skills, self-help skills, and speech and language skills;

Appendix C of the regulations for IDEA contains additional guidance on the IEP process. Serving as a "Notice of Interpretation" and using a question and answer format, Appendix C can assist the team developing a child's IEP. Excerpts of this Notice of Interpretation — relating to how teams should describe a student's present levels of educational performance, and develop and state annual goals and short-term objectives for that student — are presented in Attachment B of this document.

Your child's need for assistive technology devices or services, or both, may also be included in the IEP. Assistive technology devices are defined as any item, piece of equipment, or product system that is used to increase, maintain, or improve the functional capabilities of individuals with disabilities (§300.5). Assistive technology devices can be acquired commercially off the shelf, modified, or customized. Since the explosion of technology in our country, assistive technology devices have become more widely available and have been shown to dramatically improve the functional capabilities of individuals with disabilities in terms of mobility, communication, employment, and learning. Many of the devices have been instrumental in allowing students with disabilities to be educated in regular classrooms, working and learning alongside of their nondisabled peers. Some examples of these devices are: electronic communication aids, devices that enlarge printed words on a computer screen, devices that facilitate communication for individuals with hearing impairments, prosthetic devices, braille writers, and keyboards adapted for fist or foot use.

Assistant technology services are any services that directly assist an individual with a disability to select, acquire, or use an assistive technology device. This includes evaluating the needs of the child, including a functional evaluation in the child's customary environment. The term also includes such services as:

- purchasing, leasing, or otherwise providing for the acquisition of assistive technology devices;
- selecting, designing, fitting, customizing, adapting, applying, retaining, repairing, or replacing assistive technology devices;
- coordinating and using other therapies, interventions, or services with assistive technology devices (such as those associated with existing educational and rehabilitation plans and programs);
- providing training and technical assistance to the individual with disabilities and his or her family, if appropriate;
- providing training and technical assistance to professionals, employers, or others who provide services to, employ, or are substantially involved in the major life functions of children with disabilities. (§300.6)

Regulations for the IDEA state that schools must make assistive technology devices and/or services available to a child with a disability, if required as part of that child's special education, related services, or supplementary aids and services (§300.308). A recent policy letter from the Office of Special Education Programs (OSEP) states that "consideration of a child's need for assistive technology must occur on a case-by-case basis in connection with the development of a child's Individualized Education Program (IEP)" (Goodman, 16 Education of the Handicapped Law Report 1317, OSEP 1990). Thus, when the IEP of a student is...
Transition services can be provided as special education if they are specially designed instruction or as related services if they are required to assist a student with a disability to benefit from special education.

Because of the importance of transition services to youth with disabilities, you may wish to contact NICHCY and ask for a free copy of NICHCY’s Transition Services in the IEP. This document looks at transition services in detail.

Who develops the IEP?

The IEP is developed by a team whose members meet, review the assessment information available about the child, and design an educational program to address the child’s educational needs. This meeting, called an IEP meeting, must be held within 30 calendar days after the school district determines, through a multidisciplinary evaluation, that a child has a specified physical or mental impairment and needs special education and related services. A child’s IEP review must occur at least annually thereafter.

According to the regulations, the following people must be invited to attend the IEP meeting:

- One or both of the child’s parents (subject to requirements in §300.345);
- The child’s teacher(s). If the child has more than one teacher, State policy or law may specify which teacher should participate;
- A representative of the school other than the child’s teacher (this person must be qualified to provide special education or supervise its provision);
- Other individuals, at the discretion of the school or the parents; and
- The child, when appropriate.

Depending on the purpose of the meeting, other participants may be involved. For example, when a child has been evaluated for the first time, the school must also ensure that a member of the evaluation team participates in the IEP meeting, or that someone knowledgeable about the evaluation procedures and results (e.g., a representative of the school or the child’s teacher) is present. [§300.344(b)]

If one of the purposes of the meeting is the consideration of transition services for the student, then the school must also invite the student and a representative of any other agency that will be responsible for providing or paying for the transition services. [§300.344(c)]. "For all students who are 16 years or older, one of the purposes of the annual meeting will always be the planning of transition services, since transition services are a required component of the IEP for these students" (§300.344, Note 2).

So we, as parents, are involved in developing our child’s IEP?

Yes. The law is very clear that parents have the right to participate in the meeting where their child’s IEP is developed. Therefore, your school district must ensure that you have the opportunity to participate in your child’s IEP meeting and to contribute to the development of his or her IEP. This means that:

- The school district must notify you of the meeting early enough to ensure that you will have an opportunity to attend. [§300.345(a)(1)]
- The notice given you by the school district must indicate the purpose, time, and location of the meeting. If one of the purposes is to consider transition services, the notice must indicate this. [§300.345(b)(1) and §300.345(b)(2)]
- The notice must also identify who will participate in the meeting. If transition services are to be discussed, the notice must indicate that the school will invite your child and identify any other agency that will be invited to send a representative. [§300.345(b)(1) and §300.345(b)(2)]
- The school must attempt to schedule the meeting at a time and place agreeable to you and the school. [§300.345(a)(2)]
- The school must take whatever action is necessary to ensure that you understand the proceedings at the meeting, including arranging for an interpreter for you, if you are deaf or if your native language is other than English. [§300.345(e)]
The regulations state that the school may hold the IEP meeting without the parents if they are unable to attend. However, the school must have a record of its attempts to arrange a mutually-agreed-upon time and place. This can be accomplished by keeping detailed records of telephone calls made or attempted and the results of those calls, copies of all letters sent to the parents and any responses received, and a record of any visits made to the parents' home or work and the results of those contacts. [§300.345(d)] The regulations also state that if neither parent can attend the IEP meeting, the school must use other methods to ensure their participation, including individual or conference telephone calls [§300.345(c)].

What occurs during an IEP meeting?

As has been said, the purpose of the IEP meeting is to develop a child’s IEP. The meeting can serve as an excellent communication vehicle between parents and the school. It enables the parents and the school to decide what the child’s needs are, what services will be provided, and what outcomes can be anticipated, and to specify these in the IEP.

Many school districts conduct the IEP meeting separately from the meeting where the child’s eligibility for services was determined. Other school systems combine the eligibility meeting and the IEP meeting, moving directly into developing the IEP once a child has been determined eligible for services. Whichever approach the school district uses, once your child’s eligibility for services has been determined, the focus of discussion will be on developing his or her IEP. You and the other members of the team will discuss:

- what educational goals and objectives are appropriate for your child;
- what type of special education your child needs;
- which related services are necessary to ensure your child benefits from his or her special education;
- what assistive technology devices or services (if any) your child needs to benefit from special education;
- what transition services are necessary to prepare your child for life after completing high school (this applies if your child is age 16 or older, or younger in some circumstances); and
- what placement alternatives exist (see discussion of LRE on page 3), and which are most appropriate.

You, as the parent, may wish to provide information on your child’s educational (and, when appropriate, transition) needs, offer suggestions for the services appropriate for meeting those needs, and help select an appropriate program. At any point during the IEP meeting, don’t hesitate to ask questions until you are sure that you understand what is being said.

Following the team’s discussion, decisions will be made about the educational program and related services that best suit your child’s needs. These decisions are then specified in the IEP. You are entitled to receive your own copy of this document [(§300.345(f)].

It is important to understand that the IEP sets out the individualized instruction and related services to be provided to the child but it is not a contract. While the school, teacher, and others are not liable if a child does not achieve the growth projected in the annual goals and objectives, the school is responsible for providing the instructional and related services written into the IEP (§300.350). In order to check on your child’s progress, you will find it helpful to have a copy of your child’s IEP.

The IEP constitutes the basis for your child’s special education placement. It is important to know that your written consent must be obtained before your child is placed in a special education program for the first time [§300.504(b)(ii)]. School districts do have procedures they can follow to override parents’ unwillingness to give consent; these are discussed in Part V.

How often is my child’s IEP revised?

The law clearly makes provisions for the growth and changing needs of children. At least once a year, whether you request it or not, a meeting must be scheduled with you to review your child’s progress and to develop a new IEP for the upcoming year [§300.343(d)]. In addition, you may request a review or revision of the IEP at any time. However, for any changes in your child’s special education program after the initial placement, your written consent is not required under Federal law(§300.504, Note 1). What is required is that the school district give you prior written notice, within a reasonable time, of any changes it intends to make regarding your child’s education program [§300.504(a)(1) and Note 1].

How can I involve myself after my child’s IEP is developed?

It is in the best interests of everyone— the parents, the school district, and the child with a disability — that the school and the parents maintain a good working relationship. The following are some suggestions that parents can use to develop and maintain a positive working relationship with professionals who work with their child.

- Let your child’s teacher(s) and therapists know that you are interested in playing an active role in your child’s educational program. Plan and schedule times to talk with the professionals working with your child and, if possible, visit the classroom or program.
- Offer to explain any special equipment, medication, or medical problem which your child has.
- Ask that samples of your child’s work be sent home. If you have questions, make an appointment with your child’s teacher(s) or therapists to discuss new strategies to meet your child’s goals.
- Ask for suggestions of how you can continue, expand, and reinforce your child’s educational activities at home.
- Volunteer to be a classroom or program parent. In this way, you can observe how things work in your child’s program or school and how your child interacts with others.
- Let the school or program know that you may be called if you are needed.
- Remember that both you and the school or program in which your child is enrolled want the best for your child. Working together can make this happen.
The IDEA includes an entire section entitled "Procedural Safeguards." These safeguards are designed to protect the rights of parents and their child with a disability, as well as to give families and schools a mechanism for resolving disputes.

Procedural safeguards under the IDEA (many of which have been described previously in this News Digest) include the right of parents to inspect and review their child's education records; the right to obtain an independent educational evaluation (IEE); the right to written prior notice on matters regarding the identification, evaluation, or educational placement of their child, or the provision of FAPE to their child; the right to request a due process hearing on these matters; which hearing must be conducted by an impartial hearing officer; the right to appeal the initial hearing decision to the State Education Agency (SEA) if the SEA did not conduct the hearing; the right of the child to remain in his or her present educational placement, unless the parent and the agency agree otherwise, while administrative or judicial proceedings are pending; the right to bring a civil action in an appropriate State or Federal court to appeal a final hearing decision; the right of the parent to request reasonable attorney's fees from a court for actions or proceedings brought under the IDEA under the circumstances described in §615(e)(4) of the IDEA; and the right of parents to give or refuse consent before their child is initially evaluated or placed in a special education program for the first time.

We will look at some of these areas in this section, including: the confidentiality of education records; access to education records and the parents' right to request that records be amended; and what parents can do if they do not agree with an educational decision made by the school system regarding their child.

Are my child's records confidential?

There are provisions under the IDEA (and other Federal laws as well) that protect the confidentiality of a child's education records. These safeguards address three issues: (a) the use of personally identifiable information; (b) who may have access to a child's records; and (c) the parents' right to request their child's records be amended.

The IDEA guarantees you the right to inspect and review any records relating to your child that the school district... collects, maintains, or uses regarding the identification, evaluation, and educational placement of your child and the provision of FAPE to your child...

Personally identifiable information means information that includes: (a) the name of the child, parent, or other family member; (b) the address of the child; (c) a personal identification number (such as the child's social security number or student number); or (d) a list of personal characteristics or other information that would allow the child to be identified with reasonable certainty ($300.500). With a number of exceptions, you must give your consent before any personally identifiable information can be disclosed by the school district ($300.571). These exceptions are specified by your State or other participating agency's policy in keeping with the regulations of §99.31 of the regulations for the The Family Educational Rights and Privacy Act (FERPA), P.L. 93-380. (Regulations for the entire FERPA can be found in 34 CFR §99.1 through §99.67. IDEA's regulations on confidentiality — §300.560 through §300.576 — contain several references to FERPA.) You have and review any records relating to your child that the school district or other participating agency collects, maintains, or uses regarding the identification, evaluation, and educational placement of your child and the provision of FAPE to your child ($300.502 and §300.562). Should you ask to review your child's records, the school district or other participating agency must respond to the request without unnecessary delay and before your child's IEP meeting or a due process hearing involving your child, and in no case later than 45 days after the request. You also have the right to reasonable requests for explanations and interpretations of the records. You may ask the school district or other participating agency to provide you a copy of your child's records and you have a right to a copy if you do not live within commuting distance of your school district or participating agency. The school may charge you a reasonable fee for making
copies, as long as this fee does not effectively prevent you from inspecting and reviewing the records. Schools or other participating agencies may not charge you for searching for or retrieving the records [§300.566(b)]. Furthermore, you have the right to obtain from the school district or other participating agency a list of the types of education records that are collected, maintained, or used by the agency, and where these records are kept (§300.565).

In keeping with the requirements of the FERPA, only certain individuals besides you, as parents, may have access to your child's records. These individuals may include, for example, teachers or officials of the school or State who have a legitimate educational interest in the records. The school or other participating agency is required to maintain a record of all parties who obtain access to a child's educational records (with the exception of parents and authorized employees). This record should include the name of the person who accessed the records, the date, and the purpose for which the person was authorized to use the records (§300.563).

The right to request that records be amended is also given to parents under the law. If you believe the information in your child's records is inaccurate or misleading or that information in the records violates your child's right to privacy, you may request that the school district or other participating agency amend this information [§300.567(a)]. The district must then decide, within a reasonable amount of time, whether to comply with your request [§300.567(b)]. If the district or other participating agency decides to refuse your request, it must inform you of this decision, as well as advise you of your right to a hearing [§300.567(c)].

If you decide to challenge the school district's or other participating agency's refusal through a hearing, you have the right to present evidence showing why you feel the information in your child's records should be amended (§99.23). The hearing must be conducted by an individual who does not have a direct interest in its outcome, and the educational agency or institution must make its decision in writing within a reasonable amount of time after the hearing (§300.570 of IDEA refers readers to §99.23 of FERPA). Should the decision be in your favor, the district or other participating agency must amend your child's records and inform you in writing that it has done so (§300.569(a)). If, however, the decision supports the school district's refusal to amend the records, then you have the right to add a statement to your child's records authorizing the school to conduct the evaluation or place the child into a special education program [§300.504(b)(2) and Note 2]. If, however, the State has no legal requirement for parental consent outside of Federal regulations, then the school may use the IDEA due process procedures to obtain a decision that allows the initial evaluation or special education placement of a child to take place without parental consent [§300.504(b)(3)].

In any event, the school must notify the parents of its intended actions [§300.504(aXI) and Notes 1 and 2]. Parents have rights at such a due process hearing, as well as the right to appeal decisions made at a due process hearing [§300.504(b)(3)]. (See page II for a discussion of due process hearings and the rights of parents at these hearings.)

**What can I do if, at some point in time, I don't agree with decisions the school makes concerning my child?**

There are several, different procedures that you might want to use when you are not in agreement with the decisions recommended by the school with regard to your child's identification, evaluation, educational placement, or the provision of FAPE. Five commonly used procedures are listed below. You may want to use one or more of these approaches; some may be more appropriate than others at different times and for different reasons. Parents may also find it useful to involve an advocate in some of these approaches. An advocate can help parents to understand their rights and responsibilities fully, as well as those of the school district, and can assist in discussions concerning whatever differences exist between parents and the school system. (To find out more about advocates in your area, contact a local disability organization or parent group.)

**The school district does have steps it can take that may or may not result in an overriding of parents' refusal of consent**

There are two ways that parental consent can be withheld. One is that the parents simply do not respond to any of the school's communications, thereby withholding consent through silence. The other is explicit denial of consent, meaning that parents refuse orally or in writing to allow their child to be evaluated for or placed into a special education program.

In both cases, the school district does have steps it can take that may or may not result in an overriding of parents' refusal of consent. If the district feels strongly that the child should be evaluated to determine if a disability exists, or if the district feels strongly that the child should be placed into special education, the district can pursue the matter through the procedures specified in Federal or State law. Where State law requires parental consent, the school must follow procedures developed by the State, such as obtaining a court order for or retrieving the records [§300.566(b)]. This statement would, in effect, comment upon the information in dispute and set forth reasons why you disagree with the school district. The district must then place your statement in the records, keep it there as long as the records are maintained, and share it with any party to whom the records are disclosed [§300.569(c)].
Due process hearing. You may request a due process hearing if you do not agree with your State educational agency (SEA). Once the complaint is received, the SEA must conduct an investigation to be necessary. The SEA must review all relevant information and make an independent determination as to whether a requirement of the IDEA has been violated. The SEA must conduct an on-site investigation, if it determines such an investigation to be necessary. The complainant also must be given the opportunity to provide information, either orally or in writing, regarding the complaint. Once the complaint resolution procedure is completed, the SEA must issue a written decision that addresses each of the allegations in the complaint and contains the reasons for the SEA's decision.

A party to the complaint who disagrees with the State's final decision has the right to appeal that decision by requesting review by the U.S. Secretary of Education. Secretarial review of complaints involving Part B is the responsibility of the Assistant Secretary for the Office of Special Education and Rehabilitative Services (OSERS). To find out more about your State's complaint procedures, contact the Director of Special Education at your SEA and request a copy of these procedures. Additional clarification is also available about how the due process procedure and the state complaint process operate as distinct and separate remedies; specific guidance is provided in this regard in the December 4,1991 letter of clarification from the United States Department of Education (Davila, 1991).
At the hearing, the arguments and evidence of both the parents and the school district are presented before the impartial hearing officer, who then gives a decision. The decision is considered final, unless one of the parties involved in the hearing appeals the decision to the SEA ($§300.510). (Appeal to the SEA is available only if the SEA did not conduct the hearing.) If the hearing decision is appealed, the SEA must then conduct an impartial review of the hearing, examining the entire hearing record and seeking additional information, if necessary. If the reviewing official wishes to, he or she can allow both parties to submit oral and/or written arguments. The matter must be resolved and a copy of the final decision must be mailed to both parties within 30 days of the request for a review ($§300.512).

Part VI:
Additional Resources

As was mentioned at the beginning of this News Digest, two of the best resources a parent can have in regard to the educational rights of their child with a disability are copies of the law itself and the Federal regulations written for the law. Refer to page 1 for information about where and how to obtain these resources. Information about how your State implements the law is also useful; State policies are generally available by contacting your school district or State's Director of Special Education Programs.

For additional information, you may find it helpful to contact NICHCY for other materials about the law and how to access special education services. Contact NICHCY for a Publications List, to talk to an Information Specialist about your question(s), or to locate other support groups in your area.

References


Goodman, 16 Education of the Handicapped Law Report (EHLR) 1317, OSEP 1990. (Available from LRP Publications, 747 Dresher Road, Suite 500, P.O. Box 980, Horsham, PA 19044-0980. Telephone: (215) 784-0860.)


Attachment A
IDEA's Definitions of Disabilities

The IDEA lists 13 separate categories of disabilities under which children may be eligible for special education and related services. This Attachment presents the IDEA's definitions verbatim, as found in §300.7(b)(1)-(13).

(1) Autism

"Autism" means a developmental disability significantly affecting verbal and nonverbal communication and social interaction, generally evident before age 3, that adversely affects a child's educational performance. Other characteristics often associated with autism are engagement in repetitive activities and stereotyped movements, resistance to environmental change or change in daily routines, and unusual responses to sensory experiences. The term does not apply if a child's educational performance is adversely affected primarily because the child has a serious emotional disturbance, as defined in paragraph (b)(9) of this section.

(2) Deaf-blindness

"Deaf-blindness" means concomitant hearing and visual impairments, the combination of which causes such severe communication and other developmental and educational problems that they cannot be accommodated in special education programs solely for children with deafness or children with blindness.

(3) Deafness

"Deafness" means a hearing impairment that is so severe that the child is impaired in processing linguistic information through hearing, with or without amplification, that adversely affects a child's educational performance.

(4) Hearing impairment

"Hearing impairment" means an impairment in hearing, whether permanent or fluctuating, that adversely affects a child’s educational performance but that is not included under the definition of deafness in this section.

(5) Mental retardation

"Mental retardation" means significantly subaverage general intellectual functioning existing concurrently with deficits in adaptive behavior and manifested during the developmental period that adversely affects a child’s educational performance.

(6) Multiple disabilities

"Multiple disabilities" means concomitant impairments (such as mental retardation-blindness, mental retardation-orthopedic impairment, etc.), the combination of which causes such severe educational problems that they cannot be accommodated in special education programs solely for one of the impairments. The term does not include deaf-blindness.

(7) Orthopedic impairment

"Orthopedic impairment" means a severe orthopedic impairment that adversely affects a child’s educational performance. The term includes impairments caused by congenital anomaly (e.g., clubfoot, absence of some member, etc.), impairments caused by disease (e.g., poliomyelitis, bone tuberculosis, etc.), and impairments from other causes (e.g., cerebral palsy, amputations, and fractures or burns that cause contractures).

(8) Other health impairment

"Other health impairment" means having limited strength, vitality or alertness, due to chronic or acute health problems such as a heart condition, tuberculosis, rheumatic fever, nephritis, asthma, sickle cell anemia, hemophilia, epilepsy, lead poisoning, leukemia, or diabetes that adversely affects a child's educational performance.

(9) Serious emotional disturbance

"Serious emotional disturbance" is defined as follows:

(i) The term means a condition exhibiting one or more of the following characteristics over a long period of time and to a marked degree that adversely affects a child's educational performance:

(A) An inability to learn that cannot be explained by intellectual, sensory, or health factors;
(B) An inability to build or maintain satisfactory interpersonal relationships with peers and teachers;

(C) Inappropriate types of behavior or feelings under normal circumstances;

(D) A general pervasive mood of unhappiness or depression; or

(E) A tendency to develop physical symptoms or fears associated with personal or school problems.

(ii) The term includes schizophrenia. The term does not apply to children who are socially maladjusted, unless it is determined that they have a serious emotional disturbance.

(10) **Specific learning disability**

"Specific learning disability" means a disorder in one or more of the basic psychological processes involved in understanding or in using language, spoken or written, that may manifest itself in an imperfect ability to listen, think, speak, read, write, spell, or to do mathematical calculations. The term includes such conditions as perceptual disabilities, brain injury, minimal brain dysfunction, dyslexia, and developmental aphasia. The term does not apply to children who have learning problems that are primarily the result of visual, hearing, or motor disabilities, of mental retardation, or emotional disturbance, or of environmental, cultural, or economic disadvantage.

(11) **Speech or language impairment**

"Speech or language impairment" means a communication disorder such as stuttering, impaired articulation, a language impairment, or a voice impairment that adversely affects a child's educational performance.

(12) **Traumatic brain injury**

"Traumatic brain injury" means an acquired injury to the brain caused by an external physical force, resulting in total or partial functional disability or psychosocial impairment, or both, that adversely affects a child's educational performance. The term applies to open or closed head injuries resulting in impairments in one or more areas, such as cognition; language; memory; attention; reasoning; abstract thinking; judgment; problem-solving; sensory, perceptual, and motor abilities; psychosocial behavior; physical functions; information processing; and speech. The term does not apply to brain injuries that are congenital or degenerative, or brain injuries induced by birth trauma.

(13) **Visual impairment, including blindness**

"Visual impairment including blindness" means an impairment in vision that, even with correction, adversely affects a child's educational performance. The term includes both partial sight and blindness.

Note: If a child manifests characteristics of the disability category "autism" after age 3, that child still could be diagnosed as having "autism" if the criteria in paragraph (b)(1) of this section are satisfied.
Attachment B:
Excerpts from Appendix C of the Regulations for the IDEA-Discussion of Selected Components of the IEP

Appendix C of the regulations for IDEA contains specific guidance for the team developing a student's IEP. Using a question and answer format, Appendix C adds more detailed information about the IEP process than is available in the main body of the regulations. Parts of this "Notice of Interpretation" are excerpted below.

Present Levels of Educational Performance

- "The statement of present levels of educational performance will be different for each child with a disability." (Question 36)

- In describing a child's present levels of educational performance, "the statement should accurately describe the effect of the child's disability on the child's performance in any area of education that is affected, including (1) academic areas (reading, math, communication, etc.), and (2) non-academic areas (daily life activities, mobility, etc.)." (Question 36)

- Moreover, the statement "should be written in objective, measurable terms, to the extent possible. Data from the child's evaluation would be a good source of such information...Whatever test results are used should reflect the impact of the disability on the child's performance. Thus, raw scores would not usually be sufficient." (Question 36)

Annual Goals

- The annual goals in the IEP are statements that describe what a child with a disability can reasonably be expected to accomplish within a twelve month period in the child's special education program." (Question 38)

Short-term Instructional Objectives

- "Short term instructional objectives (also called IEP objectives) are measurable, intermediate steps between the present levels of educational performance of a child with a disability and the annual goals that are established for the child. The objectives are developed based on a logical breakdown of the major components of the annual goals, and can serve as milestones for measuring progress toward meeting the goals." (Question 39)

- The short-term objectives are similar to those used in daily classroom instructional plans, in that both are used (a) to describe what a child is expected to accomplish in a particular area in a specified time period, and (b) to determine the extent to which the child is progressing towards those accomplishments. (Question 39)

However, the IEP objectives are different from those used in classroom plans, "primarily in the amount of detail they provide. IEP objectives provide general benchmarks for determining progress toward meeting the annual goals. These objectives should be projected to be accomplished over an extended period of time (e.g., an entire school quarter or semester). On the other hand, the objectives in classroom instructional plans deal with more specific outcomes that are to be accomplished on a daily, weekly, or monthly basis. Classroom instructional plans generally include details not required in an IEP, such as the specific methods, activities, and materials (e.g., use of flash cards) that will be used in accomplishing the objectives." (Question 39)

"There should be a direct relationship between the IEP goals and objectives for a given child with a disability and the goals and objectives that are in the special education instruction plans for the child. However, the IEP is not intended to be detailed enough to be used as an instructional plan. The IEP, through its goals and objectives, (1) sets the general direction to be taken by those who will implement the IEP, and (2) serves as the basis for developing a detailed instruction plan for the child." (Question 41)
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