

# Brain Injury: A Guide for Families About School





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This publication was developed by the Children and Adolescents Committee of the Brain Injury Alliance of New Jersey (BIANJ), and adapted, with permission, by the Brain Injury Alliance of Connecticut.

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# Introduction

Members of the Children and Adolescents Committee, formerly known as the Education System Advisory Group, of the Brain Injury Alliance of New Jersey (BIANJ) wrote this guide to help educators identify, understand and respond to the special needs of students with brain injuries. The writers and members of the Committee include professionals who work in the field of brain injury, cognitive rehabilitation, general and special education, school nursing and parents. The guide was adapted, with permission, by the Brain Injury Alliance of Connecticut.

This guide is for families who are looking for information and resources to help their child return to school and negotiate the academic system throughout their education. The authors hope you find this book useful and informative.

The terms, as used in this text, “parents” and “families” include biological, adoptive, single parent, extended and any other familial arrangement. This resource may be valuable for anyone who cares for a child with brain injury and may need some help understanding and negotiating the school re-entry process.

There are many people in schools who have contact with your child. The term “educators” is used to include anyone working within an educational setting, including administrators, teachers, aides, therapists, psychologists, counselors, coaches and many others. They all influence your child’s education in some way.

Brain injury affects each child differently. It is impossible to give all the answers for each situation. By familiarizing yourself with the educational system and learning what resources are available, you will be better able to find and choose those that best meet your child’s needs in school. This guide focuses on educational issues, but there are many other topics that can affect your child after a brain injury. There are references and resources at the end of this guide to help you find more information.

# Brain Injury Alliance of Connecticut (BIAC)

## **Our Mission**

The Brain Injury Alliance of Connecticut is Connecticut's partner in brain injury prevention and recovery.

BIAC offers the following programs and services in support of brain injury survivors, their families and caregivers, professionals and the community: To learn more about BIAC's free programs and services, visit [www.biact.org](http://www.biact.org) or call 860.219.0291.

- **Information & Resources Helpline**

Brain injury specialists provide support and resources for more than 4,000 inquiries from survivors, caregivers and professionals each year to ensure that the often complex and overwhelming challenges they face are negotiated more easily and effectively. Contacts are made through BIAC's toll-free Helpline, email, social media and individual in-person consultations.

- **Outreach & Education**

Through its Prevention & Education Programs, BIAC works to increase awareness and understanding of brain injury and its prevention through school, professional and community presentations. BIAC's Annual Conference is Connecticut's premiere brain injury education program, drawing more than 275 professionals each year to learn the latest developments in the field.

- **Support Groups & Social Events**

BIAC sponsors numerous support groups throughout Connecticut that provide information, support and encouragement to survivors, their loved ones and caregivers, while offering social events and opportunities such as picnics and our Valentine's Day Dance.

- **Legislative Advocacy**

BIAC advocates for legislation to improve the quality of life for people living with brain injuries while supporting initiatives that help to prevent brain injuries from occurring.

# Chapter 1

## What is Brain Injury?

The definition of brain injury has changed over the past 10 years. Brain injuries are described as traumatic or acquired based on the cause of the injury. Knowing what causes a brain injury is very important for researching effective treatment and developing prevention programs.

*Traumatic brain injury (TBI) is an insult to the brain, not of a degenerative or congenital nature, which is caused by an external physical force that may produce a diminished or altered state of consciousness, and which results in an impairment of cognitive abilities or physical functioning. It can also result in the disturbance of behavioral or emotional functioning.*

Traumatic brain injuries can happen during everyday activities. The most common causes of brain injuries among young children are falls, such as a fall from a changing table or down the stairs in a walker. Car crashes where the child is a passenger or is struck by a car are also major causes of brain injury. Physical abuse, such as hitting or shaking a child, is another. Skate boarding, rollerblading, falling off a bike or scooter, falling from a slide, and sports injuries are other common causes of brain injuries in older children.

*Acquired brain injury (ABI) is an injury to the brain that is not hereditary, congenital or degenerative.*

Acquired brain injuries are caused by some medical conditions, such as stroke, encephalitis, aneurysm, anoxia (lack of oxygen during surgery, drug overdose, or near drowning), metabolic disorders, meningitis, or brain tumors.

Although the causes of brain injury differ, the effects of these injuries on a child's life are quite similar. This guide generally refers to children with traumatic and acquired brain injuries as children with a brain injury. The term "traumatic brain injury" is used throughout the text when information provided is specific to traumatic injuries.

### **Why is brain injury called the silent epidemic?**

Brain injury is called the "silent epidemic" because of the staggering number of people who are injured each year. As many as 2.8 million people in the United States experience traumatic brain injuries each year, but only one sixth of them are admitted to hospitals. This does not even count the number of people with acquired brain injuries.

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**Brain injury is the leading cause of death and disability in children and young adults.**

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The majority of children who have mild or even moderate brain injuries may not even be hospitalized after being hurt. That means that many children and their families are not aware that a child with a brain injury may need special attention. Family members, school personnel, and even medical professionals may have trouble figuring out why a child's behavior or abilities have changed when symptoms finally appear. Often what is really a brain injury is diagnosed as a behavioral or learning problem. Thus, the "silent epidemic" includes thousands of undiagnosed brain injuries.

Medical technology has advanced so greatly in the past thirty years that many people with brain injury, who formerly would have died, are now saved. Rehabilitation programs used to be primarily for restoring physical functioning – helping people walk and talk again. The increase in the number of "survivors" of brain injury has led to new and specialized treatment in hospitals and rehabilitation programs as well as services in the community. They are all part of the road to recovery after brain injury.

Brain injury is also "silent" because most children with brain injuries look "normal". Difficulties caused by the injury may not become apparent for months or years after the injury. As a child tries to learn new and more complex information in school, or make complicated and important moral decisions, difficulties may appear over time.

### **Degrees of brain injury**

This guide has information about children with mild, moderate and severe brain injuries. If your child has been hospitalized for any length of time - a week to months - you probably have heard the terms "moderate" or "severe" to describe the injury. You have seen and experienced the physical, cognitive, and emotional effects of these injuries. As you have watched your child recover, there have likely been reports and explanations about the injury's effects.

For many children who have what is called a "mild brain injury" or concussion, there may have been no explanations. Often, these youngsters only visit the Emergency Department or a doctor's office and may be told that their symptoms will "go away". Sometimes, these difficulties continue and parents and their child become confused and frustrated by changes in mood, memory, personality, or headaches. Given that the effects of a mild brain injury most often appear in the school setting, information about mild brain injury, or concussion, is included in this guide.

### **What is mild brain injury or concussion?**

You begin to see unexplained changes in your child. The teacher calls and asks if something is happening at home to explain your child's lack of attention, forgetfulness, and impulsive calling out in class. You have seen behavioral changes as well, but cannot explain why. You remember the fall your child took when roller blading or skate boarding and the bump on the head. Your child was shaken up, but seemed okay and went back to playing. Could there be a connection... Maybe?

*My son fell off the monkey bars and hit his head. That was six months ago. It seems like now I'm on the phone every day about a problem.*

A child with a mild brain injury or concussion may not have lost consciousness. Children are less likely to lose consciousness from a brain injury than an adult. Loss of consciousness is not the best indication for mild brain injury or concussion. As a result, the injury often goes unnoticed or is not diagnosed. A few days or months later, there are some changes in the child's behavior. The child gets frustrated easily, has trouble sticking to tasks and gets distracted, or grades drop. More problems may result from a child's inability to understand these difficulties. A child may have trouble getting along with other children and become aggressive or depressed.

These behaviors can get worse when family, educators and friends do not understand these changes and don't know what to do.

Common causes of mild brain injury or concussion are:

- falls off bikes, swings, or skateboards
- collisions during sports activities such as soccer, football or baseball
- throwing or swinging a young baby
- whiplash injuries in a motor vehicle crash

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**You do not have to hit your head to sustain a mild brain injury or concussion**

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A mild brain injury or concussion usually results in a temporary change in the way the brain works. There may be a short loss of consciousness, but often there is no loss of consciousness. Rather, a child may feel confused, groggy or disoriented for a while. For example, a child may not remember what was happening just before the injury.

Some of the common symptoms or after effects include:

- headaches, nausea
- tinnitus or ringing in the ear
- slower processing of information or slow thinking
- trouble understanding written and spoken information
- distractibility or jumping from one thing to another
- difficulty changing from one activity to another
- sensitivity to lights and/or sounds
- changes in sleep patterns

A child may have one or more of these symptoms or show other changes not in this list. Every mild brain injury, or concussion, should be measured, monitored and managed for optimal recovery. Premature return to activity may prolong the recovery process and increases the risk of a second concussion. In a small percentage of children, symptoms continue for a long time and have serious effects at home and in school.

*Over the summer, our family was in a minor car crash. My daughter hit her head and the doctor in the emergency room said she might have a mild concussion. He sent us home and told us to watch her for 24 hours. After three months, her school grades dropped dramatically although she was studying more. She was socializing less. She seems to be getting more depressed every day. What should I do? Who should I call?*

Many times a child with a mild brain injury or concussion may appear to be fine until problems show up in school. Grades may drop for no apparent reason. Often, doctors are unable to find anything physically wrong, because mild brain injury is difficult to diagnose. There are usually no findings on a CT Scan or MRI to explain the symptoms. This is why education about mild brain injury is so important for your child, family and school staff.

*My child was diagnosed as having Attention Deficit Disorder (ADD) in the first grade. After falling off his bike a few weeks ago, he complained of headaches. He had been doing very well in the program developed by his school. Now he seems to be getting forgetful and is always fighting with someone.*

You can help by getting information about brain injury from the Brain Injury Alliance of New Jersey. You can also help by working closely with school staff to put together a plan to help your child adjust to these changes. For example, your child may need shorter homework assignments or extra time on a test. Support and information can help you and your child develop skills to manage any physical, cognitive and social/emotional changes at home and school. It may also prevent or minimize behavioral difficulties such as depression, temper tantrums, or acting out.

*Jack sustained a concussion at soccer practice on Monday and sat for his SAT's on Saturday. He had difficulty focusing. He managed to continue, even with an increasingly painful headache and was unable to complete the exam. Jack and his family were dissappointed with his scores as they did not match his PSAT projection - and influenced what colleges he would apply to.*

Just as accomodations are made for a sprained ankle, likewise a student with a mild brain injury or concussion needs support and assistance until the symptoms subside. A short term 504 plan may be appropriate.

Weeks or months of a school year can span more than one marking period. This can have devastating effects on your child's ability to develop the knowledge base for the entire school year. There may be rippling effects on the next year.

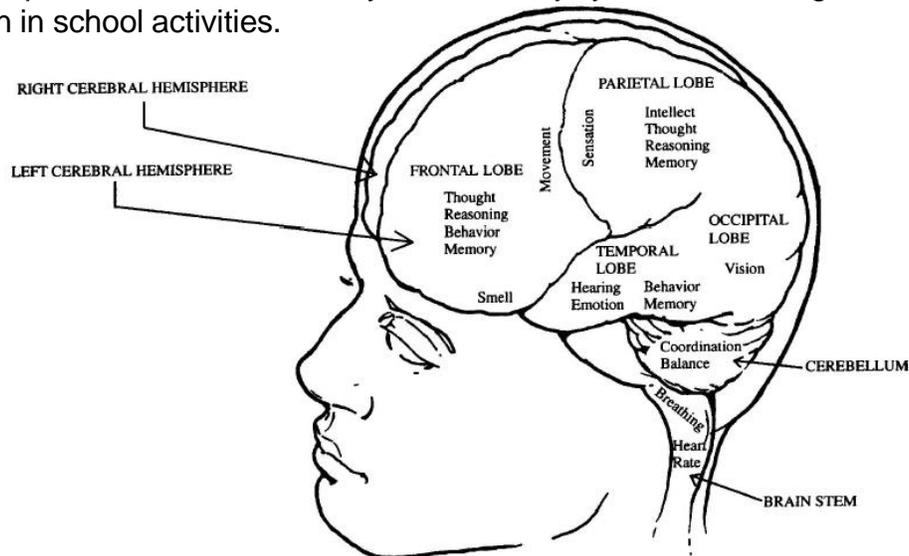
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**Education about brain injury and close contact with school staff are important for ANY student with a mild brain injury. They can reduce difficulties and failure in school and help avoid negative behavioral, emotional and academic consequences.**

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## Anatomy of the brain

When your child's brain injury is described, you may hear about different areas of the brain. Each area is associated with functions or abilities that affect how your child navigates through daily life. Basic knowledge about what these terms mean can be helpful for understanding tests, such as neuropsychological evaluations. It can also help you better explain to school staff how your child's injury affects learning and participation in school activities.



This diagram reprinted with permission from: Acute Brain Injury: A Guide for Family and Friends The University of Iowa 2000

## Common changes following a brain injury

Some of the most common changes or complaints that are reported after a brain injury are listed below. Many of these symptoms lessen over time, but some may be permanent. It is likely that your child has some of these symptoms, but it is unlikely that your child has them all. Use the checklist on the following page to describe your child.

Physical changes - does your child have ...

- fatigue or tiredness
- ringing in the ears
- blurred or double vision
- trouble walking or grasping objects
- slurred speech or trouble finding words when speaking
- unsteady balance (falls or bumps into things more)
- dizziness
- headaches
- sensitivity to lights and/ or sounds
- changes in sense of taste or smell
- nausea
- changes in sleep patterns
- endocrine or hormonal changes

Cognitive or thinking changes - does your child find it hard to...

- remember things
- concentrate
- do two things at once
- follow directions
- understand what is going on
- get the same grades as before
- solve problems and think abstractly
- get organized
- do things consistently
- read and understand at same speed as before
- recall previously learned information
- learn new things

Psychological or behavioral changes – is your child...

- irritable
- moody
- showing flat affect or little change in emotion even if very happy or sad
- self-centered or finding it hard to take another's point of view
- depressed, sad or with low energy
- fidgety
- impulsive, doing things without thinking them through

It is important that information about the cause, date, description and consequences of your child's brain injury be included in both medical and educational records. Because your child's brain is still developing, the effects of the brain injury may not be evident until your child is older and has difficulty performing more advanced tasks. Symptoms may change as a child passes through developmental milestones, such as puberty, or with transitions from one level to another, such as transition from middle school to high school. Should problems arise in the future, documentation of the injury can help educators identify causes of difficulty with learning in order to give help and support.

### **Brain injury affects the whole family**

This is a very difficult time for you as you strive to learn about the education system, advocate for your child, and deal with the varied emotions that come along with a diagnosis of brain injury. When a child sustains a brain injury, the impact is felt throughout the family and can be expressed in different ways. Other children may begin to act out, marital relations can be strained and friends and community support can taper off, leaving a parent overwhelmed, anxious, and depressed. It is important to note that these and other emotions are normal and healthy and part of your own recovery and adjustment to brain injury.

There are supportive services available to you and your family to aid in the adjustment process. Family and/or individual counseling, sibling and family support groups and/or informal school supports are often positive steps in helping family members cope. If you do have other school age children, it is important to maintain contact with their teachers and guidance counselors so

they can receive extra support and understanding.

You are receiving a lot of information within this guide and probably throughout the onset of the brain injury. There are many people and agencies, including the Brain Injury Alliance of Connecticut, listed in the back of this guide, that are available to assist you in understanding brain injury and how it relates to you.

There are also glossaries at the back of this guide that define many medical and educational terms which you may be seeing or hearing.

# Chapter 2

## Going Back to School

Each family and child faces a different situation. Your child may have been in the hospital, been transferred to inpatient rehabilitation, or received outpatient rehabilitation. Your child may have only been treated in the emergency department and sent home. As a parent reading this manual, you may be trying to figure out what to do next. Although each of you has traveled different roads, you share the common goal of ensuring that your child goes back to school with an effective educational program.

### **How do I know what my child needs to successfully return to school?**

There are many valuable types of assessments and evaluations that can be useful when your child returns to school. The type of evaluation depends on the severity of the injury and how it has affected your child. For instance, a speech evaluation is done when a student with a brain injury has problems using or understanding language; an occupational therapy evaluation when there is difficulty with fine motor movements (picking things up with hands); and a physical therapy evaluation when there is difficulty with gross motor movements (walking).

Planning & Placement Teams (PPTs) will conduct testing that measure what academic and skill level a student is able to work at. There are also other tests that examine “cognitive” or “thinking” abilities. These are important for understanding how to address weaknesses in the brain’s ability to process information. These tests also give information on how to use the student’s strengths in cognitive abilities that remain intact after a brain injury. A neuropsychological evaluation is the primary method for assessing cognitive abilities after a brain injury.

### **Neuropsychological evaluation**

It is important for a child with a brain injury to have a neuropsychological evaluation one or more times during the course of recovery. A neuropsychological evaluation is a comprehensive assessment of your child’s thinking skills (cognition), behavior, and emotional status. The testing takes between six and twelve hours to complete. This evaluation gives information on how your child is learning and identifies your child’s strengths and weaknesses. The results give information that is critical for developing an effective educational program.

A neuropsychologist is a doctorate-level psychologist with extensive training and experience in brain-behavior relationships. When a child is being evaluated, the neuropsychologist must be familiar with developmental neuropsychology. This is the study of how behavior and thinking develop and change as the brain matures over time. When choosing a neuropsychologist, it is important to find one who has experience with children and brain injury.

Most school districts do not have a neuropsychologist on staff. However, school districts are required by law to assess your child in all areas of suspected disability. A Planning & Placement Team and, later on, the Child Study Team, typically becomes involved to determine what special services are needed. These teams may be uncomfortable authorizing and funding a neuropsychological evaluation due to the cost. They also may not understand the difference

between a neuropsychologist and the school psychologist. However, as a parent, you have the right to ask for a neuropsychological evaluation.

The cost of a full neuropsychological evaluation varies, depending on geographic location and the length of testing. Because the neuropsychological evaluation gives information that is critical to planning the educational program, the school may pay for it. It is important that you know your child's rights under the law of special education and all possible sources of funding. A neuropsychological evaluation may be paid by the school, your child's medical insurance, or auto insurance (if the brain injury was due to a motor vehicle incident). The Brain Injury Alliance of New Connecticut can help you with information and resources regarding a neuropsychological evaluation.

### **Cognitive rehabilitation**

Cognitive rehabilitation may be recommended after the neuropsychological evaluation. Cognitive rehabilitation can help a child process and use information more effectively. Information from the neuropsychological evaluation helps identify areas where help is needed. For example, family and teachers may help a student with very short attention learn how to recognize distractions and how to get back on task. A student with poor short-term memory might learn to use a daily checklist or log to help with recall.

Cognitive rehabilitation may be delivered in the home or in a rehabilitation center by specialized therapists. There are a number of creative ways that school and cognitive rehabilitation may be combined. For example, a child might attend a cognitive rehabilitation program in the morning and school in the afternoon. Ongoing communication among family, the cognitive therapist, and the student's teacher can help the student carry over the skills learned in therapy to home and school.

### **What kinds of services are available through IDEA and 504?**

Creativity and flexibility at school can make the difference between your child's success or failure. The ongoing changes in your child's brain after an injury require flexible programs. The confines of the regular education system may limit the scope of services available. Differing degrees of supports or academic assistance can be obtained through the federal Individuals with Disabilities Education Act (IDEA) or Section 504 of the Vocational Rehabilitation Act.

The wording and laws about special education may be new to you and may feel overwhelming at times. As you continue reading this guide, you will develop a clearer understanding of what the terms and laws are and how they relate to your child's needs. The contact resources in the back of this guide can help you with questions and information about the educational system.

### **Understanding IDEA**

The Individuals with Disability Education Act, commonly referred to as IDEA, is the federal law which provides for a free and appropriate education for all students from birth through 21 in the least restrictive environment.

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**IDEA is the United States federal law that governs how states and public agencies provide early intervention, special education, and related services to children with disabilities from birth through age 21**

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The Connecticut special education regulations include “traumatic brain injury” as one of the specific disabilities that may be covered under IDEA.

### **Understanding 504 plans**

Section 504 of the Vocational Rehabilitation Act-1973 was established before IDEA. It was the first anti-discrimination legislation that addressed people with disabilities.

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**Section 504 protects an individual who has a physical or mental impairment that substantially limits one or more life activities such as caring for oneself, performing manual tasks, walking, seeing, hearing, speaking, breathing, learning, or working. It protects persons of all ages, not only school aged children.**

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Almost every school in Connecticut receives federal funding of some type and is therefore obligated to adhere to Section 504 of the Vocational Rehabilitation Act. The Act also includes day care programs, before and after school programs, and recreational programs.

The spirit of the law is to provide access through reasonable accommodations so all students benefit from their education.

An example of a reasonable accommodation....

*After Tom's brain injury, he found it hard to keep up with homework because he processes information slowly. When material is read to him, he understands it and can learn at grade level. Under his 504 Plan, teachers break his homework reading assignments into smaller amounts per night and extend them over the weekend. Tom also has books on tape. He reads while listening to the tape for multisensory input. It also helps pace his reading.*

There are many possible accommodations depending on what your child needs. Commonly, accommodations are used for test taking and work done in the classroom as shown by the following lists.

#### Test-taking accommodations

- allowing extra time
- dictating responses
- taking tests in a quiet setting
- having directions read, repeated or re-worded
- alternate formats, depending on a students specific needs

## Classroom accommodations

- special seating
- use of an FM system (headphones that directly link the teacher to the child and screen out extraneous noise)
- equipment access (use of calculator, laptop computer, tape recorder, or note taker)
- modified assignments (shortening classwork and homework tasks)
- enlarged print
- assistive devices to assist with memory, organization, planning, etc.

## What decisions need to be made?

Your school district will determine whether or not your child meets the criteria to be eligible for IDEA or Section 504 services. Deciding whether or not you want your child to receive these services can be a difficult decision. Fear of classification, labeling, stigma, and concerns about being in the special education system are just a few of the concerns expressed by parents.

*My child had a moderate traumatic brain injury last year and went through outpatient rehabilitation and cognitive remediation for six months. She lost a year of school and they say she needs to be classified. I don't want her classified, but I don't know what to do.*

Some adolescents may object to appearing “different” from their peers, particularly if they feel that the environmental accommodations of Section 504 or IDEA draw attention to them. Focusing on the extent of your child’s injury and its impact on your child’s ability to function in school can help you decide whether or not IDEA or 504 accommodations are necessary to meet your child’s specific educational needs.

Additional support services, such as counseling, and a sensitive approach by school personnel can help you and your child develop a hopeful and positive approach. Children may be classified as eligible for special education and related services and receive special education supports and services in general education classrooms when appropriate.

The Individualized Education Programs (IEPs) that are required by IDEA and plans designed under 504 increase the responsibility and accountability of the school to provide specific services. The required documentation of your child’s program is an advantage of these plans. This makes the school legally accountable for providing an appropriate educational program as developed in the IEP or 504 Plan. This also gives an organized way of passing on important information from year to year. It avoids depending on “word of mouth”, thus insuring that valuable information about your child is not lost.

A student who is determined, through the evaluation process, to be eligible for IDEA services is entitled to receive related or ancillary services that are necessary to meet educational needs. These services may include, but are not limited to, physical therapy, occupational therapy, speech therapy, counseling and special transportation. The need for and frequency of related services are assessed based on how the brain injury affects your child’s ability to participate in activities at school. While the services discussed above may be provided under IDEA, they are not typically provided as part of a 504 Plan. A 504 Plan focuses on adaptations

to a regular classroom, such as having tests without a time limit or taking a test orally instead of writing it. Please refer to the chart later in this chapter for some differences between IDEA and Section 504. It is important to learn everything you can about special education in your district, and the services available. You may want to speak to people who have experience with school reentry after brain injury and find out what others have done. Call the Brain Injury Alliance of Connecticut or the CT Parent Advocacy Center (CPAC). Speak to other parents whose children have faced similar situations.

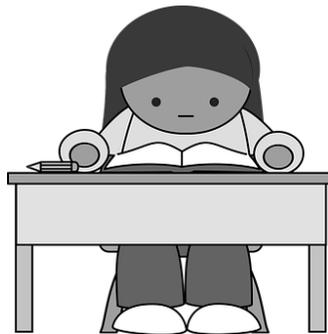
The following chart may help you understand how decisions are made regarding IDEA and Section 504. Keep in mind that no decision is absolutely final. If you start with a plan and later find that it is not working, you can revise it. It is vital to have frequent and accurate communications with teachers and school liaisons so that needed changes are not delayed. If you wait until the end of a marking period, catching up will be more difficult.

If any services become unnecessary, they can be stopped. One of the most important considerations for returning to school is to work or build from real successes, not failures. Many times a student with a brain injury has changes in what they can and cannot do; focusing on *what cannot be done* can lead to a “deficit identity.” Letting your child try school without any supports and waiting to “see what happens” should be considered with caution. While this may seem like a supportive approach as a start, it can backfire if your child has difficulties and then feels like a failure. A successful return to school requires a thorough assessment of a child’s coping abilities, academic strengths, and weaknesses.

## How does IDEA work?

The basic steps to get special education services for your child are:

- referral
- identification
- evaluation
- determination of eligibility
- development of the Individualized Education Program (IEP)
- review



	<b>IDEA</b>	<b>Section 504</b>
Purpose	To provide a free, appropriate public education (FAPE) in the least restrictive environment	To provide people with disabilities, to the maximum extent possible, the opportunity to be fully integrated into mainstream American life.
Scope	Applies to all public schools (and charter schools)	Applies to any program or activity that receives federal financial assistance
Protection	Children and youth from birth through high school graduation who fall within one or more of the specific categories of qualifying conditions –Traumatic Brain Injury (TBI) is included as a category	Any person, child or adult, who (1) has a physical or mental impairment that substantially limits one or more major life activities, (2) has a record of such an impairment, or (3) is regarded as having such an impairment – Individuals with brain injury are protected in and out of school
Legal	Procedures for ensuring enforcement are specified under the federal Individuals with Disabilities Education Improvement Act regulations	Procedures for ensuring enforcement are specified under federal 504 and Americans with Disabilities Act regulations.
Eligibility	Requires that a child's disability affects his/her educational performance - All students who are eligible under IDEA are also eligible for Section 504	Not all students who are eligible for Section 504 are also eligible for IDEA
Evaluation	Requires that a child be comprehensively evaluated by a multidisciplinary team	Evaluation draws on information from a variety of sources and is documented
Resulting Document	Requires an Individualized Education Program (IEP)	Does not require an IEP, but does require a written plan
Annual Review	IEP must be reviewed once a year at minimum	504 plan must be reviewed periodically, but it is up to the educational agency or institution to establish procedures for review
Reevaluation	Required every three years or more frequently if needed, or at request of the parent, but not more than once a year, unless the district and parent agree	Required before significant change in placement for school-aged children – Post-secondary education requirements vary from school to school

## **The referral**

The parent or a teacher generally requests the referral. This is the first step toward obtaining special education services for your child. You will be asked to sign a release form so that your child can be evaluated. A pamphlet describing the special education laws and your rights as a parent usually accompanies the release form.

The district will convene a Planning and Placement Team (PPT). The purpose of the PPT is to review the referral to special education, current evaluations and information, and to determine if additional information is needed to determine eligibility for special education. The Planning & Placement Team includes:

- The child's parent or parents;
- At least one regular education teacher if the child is being considered for regular education setting;
- At least one special education teacher;
- A representative from the district knowledgeable of the availability of resources;
- An individual who can interpret evaluation results;
- Others at the discretion of the parent or agency; and
- When appropriate, the child.

The PPT will determine eligibility for special education and related services, and also develop, review and/or revise a child's Individualized Education Program (IEP).

## **Evaluation and eligibility determination**

The purpose of the evaluation is to gather information on your child's functioning in many areas including the following type of skills:

- academic
- physical
- cognitive
- social/behavioral

The information gathered during the evaluation is used to determine if your child needs special education and related services. Your child may have been tested during rehabilitation. Often the PPT will accept some or all of these tests, including neuropsychological evaluations, which can reduce a great deal of stress and repeated testing.

A second PPT will be scheduled to review the results of the evaluation. Parents will receive a written copy of the evaluation results. The information will be reviewed to determine:

- Does the child have a disability?
- Does the disability have an adverse affect on the child's education?
- Does the child require special education and related services?

The evaluation must be completed, and for children who are determined eligible for special education, an IEP developed within 45 school days from the date of the written referral (not including time needed to obtain consent for evaluation).

It is very important for parents and educators to view test results cautiously. Brain injury is unique. A child with a brain injury typically can still recall and use knowledge and concepts learned prior to the injury. Scores on knowledge based tests, such as achievement tests, given after the injury may be similar to testing scores before the injury, as a student's old learning may remain intact. The ability to learn new information is often impacted after brain injury

Educational evaluations from the PPT often do not challenge the student's ability to learn new information. Yet this is the ability that is most often affected by a brain injury. Specialized neuropsychological testing can detect changes in your child's new learning style. If you feel that the school's evaluation does not accurately describe your child, then you have the right as a parent or guardian to request an independent or second evaluation at the school's expense. If the school believes its evaluation is appropriate, it can deny the evaluation and must initiate a due process hearing rather than pay for the independent evaluation. Parents can choose to seek an independent evaluation on their own. The school district must consider results of any independent evaluation, however they are not required to agree with or implement all or any of the results or recommendations.

When the evaluation is completed, the PPT will determine whether the student is eligible for special education services. If so, your child will be classified as "eligible" for special education and related services based on the criteria for a disability category that best fits your child's condition. Eligibility under IDEA is a tool to get services to meet your child's needs, not a lifelong label.

### **Knowing the differences in IDEA categories**

Students with brain injury are unique in many ways. It is important to be aware of the many differences between your child with a brain injury and others who are eligible for special education. Eligibility based on traumatic brain injury (TBI) should be used if your child meets the criteria for the federal definition of TBI used by the CT Department of Education.

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**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. Traumatic brain injury 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. Traumatic brain injury does not apply to brain injuries that are congenital or degenerative, or to brain injuries induced by birth trauma.**

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Some students with brain injuries are classified as eligible for special education and related services based on the criteria for TBI. However, within the special education system in Connecticut, students with brain injury may be categorized under other disabling conditions that also qualifies them for special services under IDEA, such as other health impairment or multiple disabilities if there are physical impairments as a result of the injury.

The lack of good statistical information on children and brain injury is not unique to the Connecticut Department of Education. The incidence of TBI compared to the number of students classified under the category of TBI bears a significant discrepancy in many states. A 2010 report, *Traumatic Brain Injury in Children: Connecticut's Silent Epidemic*, from the State of Connecticut's Commission on Children stated:

*One sign of underreporting of TBIs is that statistics strongly suggest that many more children's TBIs should have been identified. Each year in the United States, 630,000 brain injuries occur in children under 20 years of age.<sup>39</sup> Assuming that approximately 7 percent of these children have a long-term disability, then approximately 44,000 children in the U.S. – including 500 children in Connecticut – likely become disabled by TBI every year.<sup>40</sup> However, only 24,446 children are classified with TBI under the Individuals with Disabilities Education Act (IDEA), and this number includes all children served at a point in time, not just those who had been classified in one particular year. Based on the IDEA data, it would appear that thousands of Connecticut students are disabled by a TBI. However, a recent report from the CT Department of Education reported the statewide number at 122.*

These figures, and those available from the pediatric rehabilitation hospitals that specialize in brain injury rehabilitation, tell us that thousands of children with brain injuries severe enough to require inpatient rehabilitation are hidden among the population of students receiving special education services. These numbers do not reflect the many thousands more students with mild brain injuries who never received rehabilitation or other medical interventions. Part of the process of establishing an accurate census for TBI within the education system is making sure that students, who are categorized under some other condition but actually meet the criteria for TBI, have their eligibility criteria changed.

Despite meeting the TBI criteria, some districts may identify a student with a brain injury under categories such as "other health impairment", "multiple disabilities" or "emotional disturbance". Further, determining eligibility of such students on the basis of their brain injury helps ensure that statistics on the incidence or numbers of students with traumatic brain injuries are accurate. This can lead directly to more appropriate funding, research, prevention strategies, and better ability to develop effective school programs. Although students with brain injury may have characteristics similar to other disabilities they also have unique challenges that require specific strategies and accommodations.

### **Communication is the key**

Communication is an important part of a successful return to school, especially if your child has a severe brain injury. Your child may still be receiving therapies outside of school, such as occupational therapy, physical therapy, speech therapy, and counseling. As you work on developing an effective educational program, remember that communication among you, the teachers, and therapists is critical for effectiveness and consistency at home and in school.

*My son spent several months in a rehabilitation hospital after his brain surgery. We've been through so much. He's getting special services but he is very unhappy in school. The teacher doesn't seem to know what to do with him and he is just getting more frustrated and depressed. What can I do?*

You may need to initiate communication and stay involved to be sure that communication is established and continued. You might suggest that monthly progress reports with specific goals and strategies be exchanged between therapists and school personnel. By encouraging ongoing cooperation among everyone, progress can be regularly reviewed and any necessary changes can be made.

If your child is more severely injured, early contact with the school, while your child is still in the hospital, helps set up an open line of communication between the hospital and school. This also starts the process of educating the school about your child's needs. Once your child is ready to begin schoolwork, the hospital social worker or case manager can be a liaison or link between the hospital and school. As a parent or guardian, you are the primary contact with everyone. Your role is vital in the coordination of services and the communication of information.

### **Individualized Education Program (IEP)**

Some key areas addressed in the IEP include:

- present levels of academic achievement and functional performance
- Impact of the student's disability on involvement in the general curriculum
- a statement of measurable academic and functional goals and benchmarks or short-term objectives
- Programmatic accommodations and modifications, including nonacademic and extracurricular activities
- collaboration/supports for school personnel
- transition planning and related goals and objectives (beginning at the annual review following the student's 15<sup>th</sup> birthday)

The IEP also includes any program modifications or supports for school personnel that help your child reach the stated goals and objectives. The supports are to help your child participate in the general, extracurricular and other nonacademic activities. The supports also help your child be educated with other children in the general education class.

The IEP is a legal document that the school must implement. Parents are an important part of the team that develops this document. The Connecticut Parent Advocacy Center (CPAC) provides trainings and periodic workshops on IEP development to help families work with members of the PPT and to develop an effective IEP document.

The first part of the IEP states that your child is eligible for special education and related services based on the criteria for TBI. The IEP will state the dates for beginning the program and any related services (such as occupational, physical or speech therapy). It also states how often, how long and where these services will be provided.

The IEP also includes a list of annual measurable academic and functional goals and objectives for each academic subject and other areas to be addressed. Many schools use a menu type of computerized IEP form where goals and objectives are chosen from a pool or list. Sometimes these goals and objectives do not meet the unique needs of your child. If you have a goal for your child that is not in the pool of computerized goals, it is within your rights to ask the IEP Team to write in the goal.

The IEP must state specific strategies, techniques, and activities. It also lists any special equipment or instructional materials that your child needs. Any special equipment or materials (such as a computer or communication device) that must be provided by the school should be specifically written into the IEP so that the school's responsibilities are clear.

You may feel intimidated, anxious or uneasy about the IEP process the first few times you participate. While it is very familiar to school staff, it is very new to you.

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**You can invite somebody to attend an IEP meeting with you. You do not need to attend alone.**

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It is a good idea to invite an advocate who is familiar with the special education system or another helpful professional (such as a rehabilitation staff member or neuropsychologist) to go with you. They can help you decide if the program being offered is the most appropriate for your child. If you can not arrange for an advocate to go with you, you can bring anyone who can provide you with moral support and act as a notetaker. If no one is available to accompany you to the meeting you can request to change the date of the IEP meeting. If you must go to the meeting alone, it is a good idea to bring a tape recorder so that you can review the meeting at home when you are more relaxed. Be sure to notify the district prior to the meeting that you plan to tape record the meeting.

In most cases, you may agree with the content of the IEP. You do not have to sign the IEP at the meeting. Never sign a blank IEP form that is to be filled in later. Wait until the goals and objectives and instructional supports and accommodations are written, then read it and sign if you agree. It is always a good idea to bring the document home and review it thoroughly. If you agree with the IEP after carefully reviewing it, you can sign it and return it to your child's case manager.

Each local child study team identifies one person to be the child's case manager. This person is the liaison between the school and the parents and ensures that everyone involved in implementing the IEP gets the appropriate information. The case manager could be the school counselor or school psychologist.

If you disagree with your child's school district about any part of your child's evaluation,

educational program, or placement, try to work out the problem informally by talking with staff. If you can not work it out, you can have your case formally reviewed by a third party. You can begin the process by filing a complaint. If you are unsure how to file a complaint refer to special education law or call an advocate for further help. If needed, call BIAC or CPAC for more information.

## **Placement in the Least Restrictive Environment**

The PPT is responsible for where your child's education program will be delivered, based on the results of the evaluation and what is written in the IEP. By regulation, placements should be in the "least restrictive" settings. For one student, the "least restrictive" setting may be in a general education class without any supports or accommodations. On the other end of the continuum, the services that another student needs to be educated may necessitate a residential, out of district setting.

The optimal placement is to give your child the opportunity to participate with the regular school population as much as possible with reasonable help or supervision. Generally a student receives services through one, or a combination of, the following programs:

**Inclusion in a general education class...** a child with special needs participates in general education class with or without additional in class support detailed in the IEP.

**Resource programs...** individual and small group instruction provided to students with disabilities by a certified teacher of students with disabilities. Resource programs may be provided in general education class or in a pull-out session.

**Special class program...** Small group setting within the local school district. Student usually has most if not all subjects in this classroom.

**Approved Private School for Students with Disabilities...** An incorporated entity approved by the NJ Department of Education to provide special education and related services to students with disabilities placed by the district board of education responsible for providing their education

**Home instruction...** One-to-one tutoring at home or in the hospital.

**Residential...** An out of home and district placement that provides highly specialized services such as behavior modification for students with severe impairments.

As a member of the IEP team, you have an important role in determining your child's placement, but you do not do it alone. Consideration of all the information about the student, plus creativity and flexibility, are crucial for a successful placement for any child with a brain injury.

If you and the team cannot find or develop an appropriate placement in your school district that can meet your child's needs, then there are private out-of-district schools that may or may not

be suitable for your child. Deciding on an out of district placement can be a difficult process. If an out-of-district placement is appropriate for the student, it is important for you to make appointments and visit any school that is recommended. Make sure to visit when school is in session so that you can observe students and staff together. It helps to make a list of questions for staff ahead of time.

Although a private out-of-district placement may be needed for your child initially, your child may be able to enter a less restrictive placement months or years later. It is also sometimes appropriate for a child with a severe brain injury to be placed in the home neighborhood school. This placement almost always requires that the school be educated about your child's specific injury and needs. The school can have an in-service training for staff to educate them about brain injury and to prepare them to meet and understand your child's needs.

A placement within your home school district can still be flexible and creative. Reviewing your child's needs that have been outlined in the IEP can help you figure out if the suggested placement will meet those needs in the least restrictive manner. You can meet the teachers and staff that would be working with your child. Your informed participation in the placement decision-making process is very important.

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**Remember that you know your child best.**

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### **How can the school nurse help?**

The school nurse can be an important person in your child's reentry to school. The nurse's medical background helps monitor your child's recovery. Your child may see the nurse regularly for medication or if other ailments occur. Your child may become anxious and develop stomachaches or headaches. Your child may even visit the nurse when feeling cognitively overwhelmed, fatigued or having trouble with a specific subject.

By being included in the communication network, the school nurse can spot a problem early and help address it. It is always advisable to keep the nurse informed and request the nurse's presence at the transition meeting when your child reenters school. You may wish to request that your school nurse attend IEP meetings, given their important role in your child's school routine.

Whenever there is a change in your child's medication (even if the medication is not given at school), or a change in your child's medical condition, it is important to contact the school nurse.

If your child sustains a fall, injury or any change in behavior, it is important to keep in touch with the nurse regularly and ask that this information be given to other staff. If you want the school nurse to communicate directly with your child's doctor, you must give written consent. You can limit this consent to information about your child's injury or any illness that affects abilities and activities in school.

### **Sports and prevention of brain injuries**

When a student is preparing to return to school after a brain injury, a common question is about participation in gym, playing sports and other physical activities such as biking, rollerblading,

and skiing. Your child's doctor will be able to recommend what activities your child can engage in, based on your child's medical condition.

Concern about preventing an additional brain injury will influence your child's choice of activities, particularly in the first year of recovery. Multiple brain injuries, even mild ones, or one concussion after another, can have a cumulative negative effect on the brain. If your child was actively involved in sports prior to the injury, it may be difficult to accept limits on activities.

You are in the difficult position of enforcing those limitations. Deciding whether to deny your child an activity that brings pleasure and builds self-esteem versus protecting your child from the chance of another injury can be a difficult task. A second injury can worsen the effects of the first brain injury. To help make this decision and to help your child adjust to any changes, talk with doctors and therapists treating your child and discuss the level of injury and your child's current abilities. There are non-physical ways for your child to participate on a team (keeping score). It is also helpful to gather information about protective gear and how it works.

Bicycle helmets are a well-known form of protective head gear. CT law requires that children under the age of 16 wear them while riding. Helmets also protect children while skiing, rollerblading, skateboarding and riding a scooter. Statistics have shown that, when worn correctly, a properly made helmet can significantly reduce the severity of a brain injury. This can mean the difference between a fatal or severe injury and a milder one.

A helmet alone will not completely prevent an injury. Children and adolescents need to learn safety skills and how to reduce risky behaviors. For bicycle safety, the Centers for Disease Control recommend a combined approach of wearing appropriate protective gear, teaching safety skills, and developing safe riding environments. These principles can be applied to any sport or physical activity that carries the risk of sustaining a brain injury from an impact or collision.

As awareness of the serious effects of sports-related brain injuries has increased, many elementary and high school physical education teachers and coaches are learning more about the prevention and screening of injuries. If the teachers and coaches in your school have not had this training yet, you may want to let them know that such training is available.

The Brain Injury Alliance of Connecticut can help you find speakers on the prevention of sports injuries to educate school personnel. It is a lot easier for any child with a brain injury to fit in when the whole team or class plays safely. You can contact BIAC for more information about how to bring prevention into your child's school.

### **Important things to know about the Special Education Law**

*Private or parochial schools have more limited special education services.*

*Children from birth to age 3 years old are entitled to services through Connecticut's Birth to Three Program. Referrals are made through the Child Development InfoLine (CDI). The mission of the Connecticut Birth to Three system is to strengthen the capacity of Connecticut's families to meet the developmental and health-related needs of their infants and toddlers who have delays or disabilities.*

*Children ages 3 – 5 years old* can receive special education and related services through your local public school. To be eligible for these programs, you must request that your local public school district evaluate your child for eligibility for special services.

*Children ages 5 to 21* are serviced through your local school district. The principal of your assigned school can direct you to the Special Services Department.

*For adolescents ages 14 years and above*, the IEP must reflect goals for the transition from school to adult life. It is important to plan and prepare for what the adolescent will do after high school. Two key agencies that help with adult services are the Division of Developmental Disabilities (DDD) and the Division of Vocational Rehabilitation Services (DVRS).

*Delaying graduation* makes it possible for your child to receive important special educational services through the school year in which your child turns 21 in some instances. When a student graduates, the responsibility of the school system for providing special education services is over, and thus, there may be a gap in services. Graduation should be considered very carefully. Your child may want to graduate at age 18 years. You can explore having your child participate in commencement exercises without actually graduating or receiving a diploma. This may ease your child's anxiety or sense of loss by including your child in this important event.

Thoroughly investigate all program options to consider what each has to offer your adolescent educationally, vocationally, emotionally, and socially in order to prepare for adulthood. If you have questions about delaying graduation, be sure to talk to everyone involved.

*It is a parental right to have an advocate who* can help you prepare for and attend school meetings. An advocate can help you understand information, provide support, and ask critical questions about special education laws and your specific situation. Your local child study team is required to provide you with a copy of your parental rights.

*You also have the right to have a translator present* who is fluent in your native language. The translator can help you participate in the IEP process and read and explain the IEP to you in your native language.

*Considering assistive technology is a required part of the IEP process.* Assistive technology is a device or service your child needs to access or benefit from educational programs. It goes from low technology to high technology and includes assessment, purchasing, training, and follow-up.

Your school district cannot require you to pay for the device and services, but you can share the cost if you choose. By considering options and benefits from alternate sources such as Medicaid and health insurance, you can share the cost among several payers. BRS is another possible source of funding for assistive technology, but only during the transition to work or post high school education/training and only for school or work use.

When the school district pays the entire cost of assistive technology, the district owns the equipment. This does not mean that your child has to keep the devices at school. It can be worded in the IEP that your child can move the assistive technology from class to class and to home. During the transition process, parents can ask the school district to discuss transferring ownership or selling the assistive equipment or services to the new provider of services for your child that may be funded, if appropriate, by BRS or DDS.

Some examples of assistive technologies include:

*Audiobooks...* Your child can read along or just listen. Audiobooks give multisensory input, compensate for slower reading speed, low reading level, or difficulties with sounding out words (decoding).

*Text reader programs...* These may have study skills options. They include computer programs that read material. Your child can highlight the main points, vocabulary etc., to be printed out as notes. They can help your child compensate for physical issues affecting writing speed or legibility and also help with slow speed of processing.

Other technology may include a voice recorder to put assignments in a notebook, tablet word prediction programs or a note taker.

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**The Connecticut State Department of Education provides Assistive Technology Guidelines for students 3-21 and for infants and toddlers at <http://www.sde.ct.gov/sde/cwp/view.asp?a=2663&q=334974>.**

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# Chapter 3

## Coordinating Your Child's Services

Over time, the professional coordinator of your child's services will change. If your child has a severe brain injury, a case manager from the hospital or a rehabilitation nurse provided by the insurance company may initially help you coordinate services. With the increased use of managed care plans through health insurers you may be drawn into the often confusing domain of accessing benefits very early in your child's hospitalization. Financial counselors are also available at most acute care and rehabilitation hospitals to gather information on your benefits and to explain the needed steps to acquire non-emergency services such as Primary Care Physician (PCP) referrals.

You may hear the term case manager in many settings, for example, case manager of medical insurance, case manager of rehabilitation services and /or case manager within the school. The term, case management refers to the coordination of services; in these cases, within the specific environments where the case manager works.

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**You are always the best advocate for your child. You know your child the best and are the only constant among the many professionals involved in your child's recovery.**

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If your child has a mild brain injury, there may be a lag time between an accurate diagnosis and assignment of a case manager. You may be the only person who knows all the people and issues involved in your child's case. This may put you in the position to communicate information to and from all parties. Coordinating this information makes you a case manager.

If your child has a moderate or severe injury and is eligible for special education through IDEA, a member of the child study team will be assigned as the educational case manager. If your child is receiving accommodations in school through a Section 504 plan, the guidance counselor may be the contact person for your child's needs at school.

There is a great deal for you to learn about your child's changing medical, emotional, and intellectual needs. You are the one constant in a parade of professionals. You knew your child before the brain injury. You have overseen the entire recovery process. You know how your child is functioning today. You are the most qualified to advocate and coordinate services for your child. Case management is something you and many parents have been doing all along. You have just not been called the "case manager".

You develop and plan activities for your child and help with schoolwork. You keep files on doctor visits and school progress reports. Case management for a child with a brain injury is the same basic process. It is the newness and unfamiliarity of the medical and educational information that can be intimidating, not the case management. The number and type of

decisions you are making since your child's brain injury may be the most challenging part of being a case manager. To simplify the process, it is helpful to become organized early and keep important information where it can be easily found.

A useful way to begin organizing information about your child's brain injury and school reentry is to get a large 3 ring binder (with some blank paper for note taking), a 3-hole punch, and a calendar. Add documents as they arrive to the binder in order by date. Use the calendar to note important meetings and phone calls. Keep notes from important phone conversations (including records of all attempted calls or messages) and notes from all meetings.

As the information in the binder grows and you gather more information, it is wise to keep files of the following information along with your binder:

- A list of everyone (people from school, rehabilitation services, hospital or doctor's office) involved at any time with your child. Include the name, address, and telephone number for easy reference. You can later transfer these names into a separate telephone directory and cross-reference them with a description. For example you can enter the neuropsychologist in 3 or 4 places; the name, practice or agency name, under the heading "neuropsych", or under any other name that helps you find the person quickly.
- Copies of all "written consent to release information" forms.
- A list of your child's testing with dates and names of who did the testing.
- Copies of all written reports and medical tests, rehabilitative progress reports and evaluations, school reports including IEP's, etc., filed under corresponding categories.

You can get copies of reports from professionals over the course of your child's recovery. Having copies available in one place eliminates a great deal of stress when someone asks for a copy of a specific report.

- Keep an ongoing list of questions regarding you child's medical condition, medication, benefits, programming etc. Be prepared to ask them on the next contact or meeting.
- Keep a separate file where you put written explanations, informative guides and pamphlets, even if you are not ready for them yet. By collecting and filing this information now, it will be easier for you to find it when you need it later.

### **Learn the lingo to get through the system**

To be an effective advocate for your child, it is important to understand clearly the terms and conditions of your insurance and any other sources of funding for your child's care. The following are some of the frequent issues that parents encounter when trying to access services for their child after a brain injury.

## **Know the rules**

Be sure that you are clear about the conditions and limits of your insurance policies, particularly which services require precertification or preapproval from the case manager. If you have an HMO, be sure you know when a referral is needed for services.

Ask about the appeals process if a therapy or program has been denied. Initial denials can be based on the premise that the service is “not medically necessary”. However, appeals can be pursued through the insurance company and, depending on the type of insurance, through independent reviewers outside of the insurance company.

You may have questions or feel errors have been made on financial matters. To avoid this, ask for explanations in writing from insurance companies and other funding sources so that you have a direct source to refer to when pursuing resolutions.

## **Medically necessary**

The phrase “medically necessary” is a term often used by payers. What seems so vitally medically necessary to you, as a parent of an injured child, may not seem medically necessary to the insurance representative. That is why it is very important to ask for decisions in writing, to consult with medical professionals, and not to be afraid to seek an appeal. Seeking the advice of an attorney may also be advisable if you feel that your child has been inappropriately denied the benefits/ coverage stipulated in your insurance policy. Often, an initial denial of a service is overturned during an appeals process as more information is reviewed about the case.

## **Importance of 60 days**

Health insurance policies generally have limits on the number of days and/or sessions of rehabilitation therapies covered under the policy, particularly those provided on an outpatient basis. Ask to have terms such as “60 days” clearly explained.

For example, does 60 days mean 60 consecutive days of treatment or 60 sessions. Is the limit of sessions per calendar year or for the lifetime of the policy? Knowing the conditions of your insurance is extremely important so that you can coordinate therapies provided by school and insurance to maximize benefits for your child.

## **Special education law**

A free and appropriate education must be provided to students under the special education law. This is not the same as providing rehabilitation to reach maximum recovery from an injury or illness. This is illustrated in the difference between “medically necessary” versus “educationally related” services.

Physical, occupational, and speech therapies can be provided as related services when included in the IEP, if your child has been found eligible for special education and related services within your local school district. Augmentative and assistive devices (see glossary) along with transportation to and from the school program can also be provided through your school district.

Special education regulations state that the requested therapies must be “required for the student to benefit from the educational program” in order to receive these services through a school district in Connecticut. This means that the disability or impairment must impede or affect your child’s ability to participate in the learning environment.

Example of coordinating school and health insurance resources:

*Your child’s physician prescribes physical therapy three times a week to improve independent walking. The school says it is not necessary to provide the therapy at this frequency if the child can sit in class and attend to task. The child study team recommends PT only once a week to improve movement throughout the school and to increase inclusion. You choose to advocate for more therapy time in school and/or use your health insurance to provide the additional sessions. This is an example of how important it is to know the limits of your benefits and how to coordinate legally mandated educational services with insurance benefits. As a result, your child has the greatest likelihood of receiving all the assistance needed.*

### **Advocacy groups and alternative funding sources**

There are parent founded advocacy groups as well as many relevant services that are available through state and county departments of human services in New Jersey. There are also free and low cost legal services.

Suggestions for resources....

- go to the Resources section in this guide.
- contact the Brain Injury Alliance of Connecticut’s Information and Resources Family Helpline at 1-800-278-8242.
- explore resources for parents of children with disabilities through the internet and social media.
- contact the Department of Education for a free booklet, *A Parent’s Guide to Special Education in Connecticut*, that lists advocacy services and resources.
- attend trainings on special education and advocacy, such as through CPAC. These are opportunities to obtain current information on education, funding options, and resources, and to network with other family members of children with special needs.
- join an advocacy organization and receive their informative newsletters and e-mail communications.

# Chapter 4

## Personal stories

### How life changed for Katie, Kevin, Eric and their families

Three personal stories show different options and interventions for children with mild, moderate and severe brain injuries. They show how information in this guide can be used to make a difference.

#### Katie's mild brain injury

Katie was in second grade when she brushed up against a moving school bus. She bounced back from the bus, bumping her head on the curb. Katie bruised her ribs and shoulders and lost consciousness for about one minute. Afterwards, she was dazed, held her head, and kept saying that her head ached.

She was taken to the local hospital's emergency department and had skull and chest x-rays. The results were normal and she was sent home. Katie continued to complain of neck and head pain, nausea, and a ringing or buzzing noise in her head. Her family took her to their chiropractor who performed some spinal adjustments.

Katie was out of school for one week. Upon returning to school, she still complained of head pain and ringing in her ears. She reported that the pain and noise distracted her and made her a little tired. These symptoms lessened over the first three months following the accident.

In first grade before her injury, Katie scored between the 75th and 98th percentile ranks on her achievement testing and had been a good student. Three weeks following the accident, she took her second grade achievement tests and scored much lower, between the 30th and 50th percentile. Reading and arithmetic scores had the most dramatic decline. The teacher noticed that Katie found it hard to retain new information and that she read more slowly. She often had to reread material, sometimes several times, to understand the information. Katie also had some difficulty with writing.

Her family had noticed changes too. They noticed that Katie was unhappy, irritable and more withdrawn. Her family had consulted an attorney about the accident. As the attorney gathered information about Katie's injuries, she recognized the symptoms of post-concussive syndrome. She referred Katie to a neuropsychologist, who confirmed the presence of cognitive slowing and attention difficulties due to a traumatic brain injury.

The neuropsychologist recommended to Katie's family that they write a letter to the Planning & Placement Team (PPT) at her school to request an evaluation for special services to help her with schoolwork. The neuropsychologist also recommended that Katie receive outpatient cognitive rehabilitation services. Specifically, rehabilitation was suggested to help Katie understand and deal with the changes she was going through from the concussion. She also needed help to develop compensatory strategies to cope with these changes in order to be successful at school again.

The PPT evaluation was completed and determined that Katie did not require services under IDEA. They suggested that the rehabilitation center and school work together to develop a 504 plan. The PPT gave information about 504 to Katie's family. They discussed how the strategies that Katie needed could be spelled out in a 504 plan and carried out by the class-room teacher.

Katie's parents were confused and sought help to understand what was the best thing to do for their daughter. They contacted the Brain Injury Alliance of Connecticut and were put in touch with an advocacy group. An advocate attended the school meeting with Katie's parents and helped them to work out a 504 plan that was clear, but flexible. This seemed important because Katie's needs were not completely understood but she was continually improving.

Many aspects of the first month were rocky. Katie did not follow through with homework for school and was tired by the time she got to her twice-weekly rehabilitation appointments in the afternoon. A meeting was called where the rehabilitation therapist attended and brainstormed with Katie, her parents and the school staff. A modified plan was developed that increased Katie's rehabilitation hours. Even though this sounded like a step backwards to Katie's parents, they agreed to try it because the rationale made sense. The rehabilitation therapist's plan was to use Katie's homework as part of the therapy content and to develop routines, organizational study skills and attention and learning strategies within the sessions. Recognizing that Katie experienced periods of mild fatigue at school, she was scheduled to rest in the nurse's office twice a day. Communication between the rehabilitation therapist and school staff was set up on a regular basis to address problems immediately.

The 504 plan was updated and revised, initially every six weeks and then once each marking period after Katie's performance improved and held steady. The same was true with the rehabilitation services. After one month, Katie's time in rehabilitation was cut back to two hours a week. Then it was weaned to once a week, every other week, and finally down to telephone check-ins over a period of three months. What had felt like a lifetime was a school year. Katie and her parents still notice some minor cognitive changes that remain, but Katie is active and successful in school after a year. The 504 plans were discontinued in mid-third grade, with the knowledge that reevaluation and rewriting of a 504 plan would be done if any problems occurred.

## **Comments**

This case shows the importance of not being afraid to seek out and use community resources to help a student's reentry to school after an injury. Parents often hesitate to disagree with a school's recommendation or are afraid to bring others into the school's territory. Here, the combination of an informed lawyer, a neuropsychologist's recommendations, help from an advocate, and the rehabilitation staff's involvement were all important pieces in the success of Katie's return. Even though information is available about brain injury, not all school systems have expertise in this area. Most school staff will work cooperatively with community resource people because it is in the best interest of a student.

## **Kevin's moderate brain injury**

At the time of his injury, Kevin was 16 years old. He was a junior in high school and was considered by his family and teachers as a "good kid" who took a lot of risks. Kevin was a C student and had run-ins with other kids now and then, as well as detention for minor offenses. Kevin was a passenger in a car driven by his best friend when they were involved in a motor vehicle crash. Kevin was thrown several feet after the crash. His friend died.

Kevin was in the hospital's Intensive Care Unit for one week, spent two weeks in a Brain Trauma Unit and then had outpatient rehabilitation for 8 weeks. He was unconscious for 20 hours and had seven days of post-traumatic amnesia. The CT scans showed bruising and swelling of frontal, temporal, and parietal lobes in his brain.

After he was discharged from the hospital, Kevin found it hard to remember things, follow directions, organize his thoughts, express himself, and plan his time. It took him longer to process information and respond to people than before his brain injury.

In addition, his parents worried about how he was reacting to the death of his friend. Because expressing himself was harder for Kevin after his injury, it was difficult to know how he felt about both the loss of his friend and his own loss of memories and abilities.

Kevin's parents contacted the school right after the accident. They gave permission to the staff at the brain trauma unit and cognitive rehabilitation program to have ongoing contact with the school to provide updates on his progress and plan for his return to school. Rehabilitation staff went to the school and provided brain injury education for Kevin's peers.

It was suggested to Kevin's parents that he be classified to receive special education services to better meet his needs. Kevin's parents sought other professional opinions and were put in touch with the Brain Injury Alliance of Connecticut and with a parent advocacy group, CPAC, to help them. They made the decision to take advantage of special education services and they felt that turned out to be a positive move. As part of the Individualized Education Program (IEP) process, an initial transitional plan was also proposed.

Kevin's school reentry began with home tutoring in combination with outpatient rehabilitation. (He began tutoring the second week he was in the rehabilitation program.) The rehabilitation team and the tutor met weekly to exchange information. It was during this stage of recovery that Kevin also began individual counseling with a psychotherapist to help him cope with loss and grief issues and to adapt to internal changes. He was also receiving medication for depression.

Kevin progressed to a half day at school followed by outpatient rehabilitation therapies (OT, PT, and Speech) in the afternoon. After two months, Kevin was ready to increase his time at school. He began to stay for lunch and gym class.

Staff at school began to see that Kevin was having more difficulty with social interactions. He was experiencing altercations with peers and sometimes with teachers and other school staff. Especially when fatigued, he became verbally abusive. School staff met with Kevin and his parents to explore strategies to help him with the extended school day and how to handle the less structured activities that had been added to his schedule. Kevin's psychotherapist was also consulted.

Together with the child study team, Kevin, and his parents, a plan was developed to address his behaviors. Kevin's behavior began to improve with his increased awareness of what he was doing and consistent feedback about his behavior. At the close of Kevin's junior year, he was in school two-thirds time. He attended an extended school year program to increase his skill level and complete credits for his junior year. Kevin's goal for his senior year was to develop his vocational direction and be a full-time student with a half-day of academics and a half-day of workstudy. These goals were added to his IEP and transition plan as part of the ongoing revision process. Kevin's child study team also contacted the Bureau of Rehabilitation Services to consult with them about post high school services available to Kevin.

Although Kevin was making good progress in the vocational program, it was evident that he was not ready, vocationally or academically, to enter the work world. After reviewing his skills and needs, Kevin, his parents, and the child study team decided to hold off graduation for one year. Special arrangements were made for Kevin to attend the graduation ceremony and related activities without receiving a diploma. This meant he could be part of this big event but not lose his educational benefits.

Kevin, currently 19 years old, attends two academic classes and a vocational training program each day. His social skills and ability to self-monitor his behavior have continued to improve as well as his reading and math skills. While continuing to feel sadness at times over the effects of his own injury and the loss of his friend, Kevin has been able to express that he is not overwhelmed by this and has "graduated" from seeing a counselor once a week for support. He has referrals to brain injury support groups as well as a "clubhouse" type social group if he wishes to explore these resources.

## **Comments**

Kevin's experience illustrates the steps involved in going through the rehabilitation stage, school reentry, and transition processes with sensitivity to the effects that cognitive impairments, grief, and personal adjustment can present for an adolescent with a brain injury. This example demonstrates good communication among the family, rehabilitation staff, school personnel, and BRS. Delaying graduation allowed Kevin time to develop his academic and vocational skills in order for him to be better equipped to enter the work world. It also demonstrates how a coordinated effort to address behavioral issues in a positive way can have a beneficial outcome for the student.

## **Eric's severe brain injury**

As a result of an auto accident, 14 year old Eric, was propelled from a car and had a traumatic brain injury and severe chest trauma. He was hospitalized for several months and spent six months in a low-level coma. After his transfer to a rehabilitation hospital, Eric received physical, occupational and speech therapies, and later cognitive rehabilitation.

He regained his speech eight months post injury and started academic tutoring shortly thereafter. A child study team from his local school district became involved in his case once tutoring was requested. At that time, Eric was deemed eligible for special education services under the category of traumatic brain injury. One year after his accident, Eric had progressed to reading, writing, and simple mathematics. He was walking with a walker and was able to eat enough on his own to have his feeding tube removed.

Progress continued, and as discharge from the rehabilitation hospital neared, Eric's child study team case manager began to attend the monthly case conferences to plan for the transition to school. Eric's parents obtained the services of a private consultant, specializing in brain injury. This consultant instructed his teachers about Eric's unique needs and helped develop an appropriate program at school. She helped school personnel understand the effects of Eric's brain injury and introduced strategies to maximize his classroom instruction.

Once discharged from the hospital, Eric attended an outpatient cognitive rehabilitation program four days a week, followed by cognitive/academic tutoring and therapies (OT, PT, and Speech.) On Fridays, he attended a full day at his home high school where he received all academic instruction in the resource room, plus some speech therapy. He had an instructional aide to help him navigate the halls and provide extra help in class.

Eric's IEP included assistive technology. His primary assistive device was a laptop computer with word prediction software, a screen reader and scheduling/date book software. Eric was mainstreamed into a computer class where he typed any notes or assignments necessary for his other courses into his laptop.

At that point, Eric was considered a ninth grader operating on a 4th – 6th grade level. Before his accident, Eric had functioned well above grade level. Much time was taken to modify his curriculum, develop appropriate goals and strategies and put together a transition plan. Frequent meetings among his parents, child study team members, and the TBI consultant were held to review progress and update the IEP as needed.

The following September, almost two years after his accident, Eric attended a resource program science class daily in addition to his other classes. He continued to receive his related therapies after cognitive rehabilitation therapy sessions and attended school all day on Friday. After the second semester that year, Eric's inschool program was increased to three periods with two periods of resource program science and history and one period of computers. Transition planning was continually updated with special attention to services Eric might need after graduation.

After a year and a half of services in a rehabilitation center, Eric was discharged. He returned to school the following September as a full-time junior. He was included in general education classes for art and computers, and continued his resource program for academics. He also had three periods of one-on-one speech therapy in school per week. He received PT and additional academic support along with speech therapy at home. He became involved in community reentry activities with a recreational therapist. Again due to his progress, Eric's transition plan in his IEP was updated to include new possibilities after high school.

By mid-junior year, testing revealed that Eric's academic achievement scores were between the 7th and 12th grade level. His functional cognitive performance remained impaired, especially in memory, the ability to organize his thoughts, and problem solving. The school nurse identified absence seizure activity and Eric was subsequently placed on anti-seizure medication by the neurologist.

Programming in school had become a bit more problematic because his resource program curriculum generally tapped into Eric's prior knowledge and did not challenge him with new material. One regular academic class had been added to his schedule with the additional support of an aide and a supplemental class to reinforce and modify the regular class materials. Programming for his senior year included three general education classes with extra support built into his program. His IEP clearly stated what modifications would be made to the general education class curriculum, what accommodations would be required and who would be responsible for its day-to-day implementation. Accommodations written into the IEP included alternative test materials, modified assignments, tape-recorded classes for later review, study guides, and note takers.

As Eric's developing abilities changed his future educational needs, options were investigated at the post-high school level focusing on colleges and other post secondary schools with Section 504 accommodations and support programs. Eric, his parents and the rest of his child study team obtained information on possible programs and Eric and his parents visited various schools. They ended up choosing a county college with 504 accommodations.

## **Comments**

This case illustrates the slow, steady and creative school reintegration process. It also demonstrates the flexibility a school offered and the positive results for the student. By looking at Eric's unique needs, planning and following through, programming went relatively smoothly.

# Statistics

According to the Center for Disease Control, in the United States,

- 5.3 million Americans (approximately 2% of the U.S. population) live with disabilities resulting from brain injuries.
- A brain injury occurs every 23 seconds in this country, and sends 1.4 million people to the hospital each year.
  - Of this 1.4 million:
    - 235,000 will be hospitalized and survive
    - Approximately 80,000-90,000 will experience an onset of disabilities resulting from their brain injuries
    - 50,000 will die

## Traumatic brain injury and children – national statistics

- Brain injury is the most frequent cause of disability and death among children and adolescents in the United States (CDC, 1999; Lehr, 1990).
- Males are 1.5 times as likely to sustain a brain injury as females, due to differences in risk exposure and lifestyle. 14 to 24 year old males are at highest risk. (CDC, 2004).
- The leading causes of brain injury in children are falls, motor vehicle crashes and assaults (CDC, 2004).

## Traumatic brain injury and children - CT statistics

The Connecticut Department of Public Health prepared a special report on TBI data in 2005-2007 for the June 10 forum. Each year, on average, 29 Connecticut children and adolescents (birth to 19 years) die from a TBI. There are approximately 480 hospitalizations and 8,100 emergency department (ED) visits due to TBI among the birth -19 age group.

As notable as these data are, statistics underestimate the enormity of the TBI problem.<sup>17</sup> They include only those children who die, are hospitalized or who receive care in an emergency room.<sup>18</sup> An unknown number are not included in TBI statistics because they are not treated in hospitals, if they receive treatment at all.<sup>19</sup>

### References

Centers for Disease Control and Prevention. *Traumatic Brain Injury in the United States*, October, 2004

State of CT General Assembly – Commission on Children. *Traumatic Brain Injury in Children: CT's Silent Epidemic*, June 22, 2010.

## Resources

- **Children & Youth with Special Health Care Needs (CYSHCN)**  
Phone: 800-505-7000  
Website: [www.ctunitedway.org/cdi.html](http://www.ctunitedway.org/cdi.html)  
Children and Youth with Special Health Care Needs Program (ages birth -21) provides comprehensive and coordinated health care for income-eligible children who have, or are at elevated risk for chronic physical, developmental, behavioral or emotional conditions.
- **Parents Available to Help, Inc. (PATH) Parent-to-Parent & Family Voices of CT & CT Family to Family Health Information Center (F2F HIC)**  
Phone: 1-800-399- PATH (7284) or 203-234-9554  
Website: [www.pathct.org](http://www.pathct.org) Email: [pathp2pct@pathct.org](mailto:pathp2pct@pathct.org)  
Talk to a Health Information Specialist: They assist families in navigating health care systems, Information, education, training, support and referral services, One to One Connection to a family who has similar needs and experiences.
- **CT Family Support Network (CTFSN)**  
Phone: 1-877-FSN-2DAY (1-877-376-2329)  
Website: [www.ctfsn.org/](http://www.ctfsn.org/)  
The Family Support Network (CTFSN) is a network of families who have children with disabilities. They are parents interested in helping other parents to find the supports they need. Support can include information related to in home support, respite care, health care, child care, accessible transportation, parent networking and support, home and vehicle modifications, assistive technology, inclusive education programs, summer camp, recreation and after school programs.
- **Katie Beckett Waiver/Deeming Waiver/ 2176 Models Waiver Department of Social Services**  
Phone: 1- 800-445-5394 or 860-424-4904  
The Katie Beckett Waiver enables individuals with severe disabilities to be cared for at home. The income and assets of legally liable relatives (parent) are typically counted when the person with the disability is cared for at home. However, this waiver was developed to allow parents to access Medicaid for their child with a disability or medical needs without having to financially qualify for Medicaid. Eligibility: Individuals with severe disabilities age 21 or younger; Individual's income must be at or below 300% of SSI; Asset limit is \$1000; legally liable relative's income is waived.

### Educational Resources

- **CT Parent Advocacy Center (CPAC)**  
Phone: 1-800-445-2722 Email: [cpac@cpacinc.org](mailto:cpac@cpacinc.org)  
Website: [www.cpacinc.org](http://www.cpacinc.org)  
The CT Parent Advocacy Center, Inc. (CPAC) is a statewide nonprofit organization that offers information and support to families of children with any disability or chronic illness, age birth through 26 in addressing educational support. Services are free.

## **Educational Resources Cont.**

- **Disability Rights Connecticut,**  
Phone: 860-297-4300  
Case Services Advocates and Attorneys implement federal and state Protection & Advocacy mandates to protect the rights of adults and children with disabilities & to intervene on behalf of children with disabilities who are receiving special education services and are experiencing difficulty obtaining relevant educational supports.
- **State Education Resource Center (SERC)**  
Phone: 860-632-1485  
Website: <http://ctserc.org/s/>  
SERC is a nonprofit agency primarily funded by the CT State Department of Education. SERC provides information dissemination in the latest research and best practices to educators, service providers, and families throughout the state.
- **LEARNet**  
Website: [http://projectlearn.net.org/project\\_learn.html](http://projectlearn.net.org/project_learn.html)  
LEARNet is a resource for teachers, clinicians, parents, and students. LEARNet is an interactive web-based problem-solving program designed to enable families and school staff to create effective intervention and support plans for students with Acquired brain injury/ Traumatic brain injury.
- **Brain Injury Alliance of Connecticut (BIAC)**  
Phone: 1-800-278-8242 or 860-219-0291  
Website: [www.biact.org](http://www.biact.org)  
BIAC can assist in facilitating a smoother Hospital to School transition after a child sustains a brain injury. BIAC offers outreach and education and can do a general presentation in the school for the students in the classroom and/or for the educational professionals on brain injury. Call the BIAC Helpline at 800-278-8242.
- **Center on Post Secondary Education**  
Website: <http://cped.uconn.edu/>  
Lists Connecticut Resources for Assessment Services □ For a listing of Disability Services contacts on the CT college and universities go to:  
<http://cped.uconn.edu/learning-disability-ld-and-disability-services-ds-contact-personsfor-students-on-connecticut-college-and-university-campuses/>
- **Gateway Community College Step Forward Programs:**  
Contact for Step Forward is: Jaime French, Director 203-285-2505  
[jfrench@gatewayct.edu](mailto:jfrench@gatewayct.edu)  
A 10-month transition program for young adults (18-21) with mild cognitive disabilities. This non-credit program focuses on interpersonal communication skills, career exploration and workplace readiness. In addition to classroom instruction, students are placed in a community or campus-based work experience. Topics covered in the Step Forward I program may include vocational exploration & career planning, job search essentials, personal finances, interpersonal & workplace communications, self-advocacy & disability awareness, and nutrition & fitness. There is a follow up Step Forward 2.



**Brain Injury Alliance of Connecticut**  
**300 Day Hill Road • Suite 250 • Windsor, CT 06095**  
**1-800-278-8242 • [general@biact.org](mailto:general@biact.org) • [www.biact.org](http://www.biact.org)**