

Juvenile Myoclonic Epilepsy

What Is Juvenile Myoclonic Epilepsy?

Patients with juvenile myoclonic epilepsy have myoclonic, absence and generalized tonic-clonic seizures with onset during adolescence.

Seizures are usually controlled by antiseizure medication but it is a lifelong condition, and most patients will need medication for several years.

What Are the Signs & Symptoms of a Juvenile Myoclonic Seizure?

Different types of seizures can happen in juvenile myoclonic epilepsy.

Myoclonic Seizures

In a myoclonic seizure, a child:

- Has brief muscle twitches or jerks in the upper arms, shoulders, or neck
- Usually is awake and can think clearly during and right after the seizure

Absence Seizures

An absence seizure starts suddenly in the middle of activity and ends abruptly. During the seizure, a child:

- Blanks out or has staring spells that last 5-20 seconds
- May flutter their eyes or look upward
- Is unaware of what is going on during the seizure
- Returns to normal activity and won't remember having the seizure after it's over
- Has lip-smacking, chewing movements

Tonic-Clonic Seizures

In this type of seizure, a child:

- Has convulsions, or rigid muscles and rhythmic body jerks
- Rolls the eyes back
- Cries out
- May pee or poop
- Can't respond during seizure
- Is confused and sleepy after the seizure

Seizures in juvenile myoclonic epilepsy typically happen within 30 minutes of waking up in the morning or after a nap. They're more likely to happen when someone is tired, stressed, or didn't get enough sleep.

What Causes Juvenile Myoclonic Epilepsy?

The cause of juvenile myoclonic epilepsy isn't known, but it tends to run in families. Genetic mutations have been associated with the condition.

How Is Juvenile Myoclonic Epilepsy Diagnosed?

If your child had a seizure, the doctor probably will want you to see a pediatric neurologist. The neurologist will ask questions about what happened during the seizure, do an exam, and order an EEG.

How Is Juvenile Myoclonic Epilepsy Treated?

Seizures are usually well-controlled with medication, which many people will need to take for life.

How Can Parents Help?

To help your child live better with epilepsy, be sure they follow the neurologist's recommendations about:

- Taking any medicines
- Avoiding triggers (such as excessive stress, lack of sleep, blinking lights, drinking alcohol, some types of medicines)
- Taking precautions while swimming or bathing
- Knowing whether it's okay for your teen to drive
- Knowing whether your child should wear a medical ID bracelet

To keep your child safe during a seizure, make sure that other adults and caregivers (family members, babysitters, teachers, coaches, etc.) know what to do. Your doctor may prescribe an emergency medicine to give if your child has a long seizure or many seizures in a short amount of time. Be sure to ask your doctor about a seizure rescue plan for your child.

Juvenile myoclonic epilepsy is a lifelong condition. But many kids go on to live a normal life with a few extra steps taken to keep them safe.

What Else Should I Know?

If your child has epilepsy, reassure them that they're not alone. Your doctor and the care team can answer questions and offer support. They also might be able to recommend a local support group. Online organizations can help too, such as:

- Epilepsy Foundation
- <u>CDC Managing Epilepsy</u>