



## Genetic Fact Sheets for Parents

# Fatty Acid Oxidation Disorders

Screening, Technology, and Research in Genetics is a multi-state project to improve information about the financial, ethical, legal, and social issues surrounding expanded newborn screening and genetic testing – <http://www.newbornscreening.info>

**Disorder Name: Medium chain acyl-CoA dehydrogenase deficiency**

**Acronym: MCADD**

- What is MCADD?
- What causes MCADD?
- If MCADD is not treated, what problems occur?
- What is the treatment for MCADD?
- What happens when MCADD is treated?
- What causes the MCAD enzyme to be absent or not working correctly?
- How is MCADD inherited?
- Is genetic testing available?
- What other testing is available?
- Can you test during pregnancy?
- Can other members of the family have MCADD or be carriers?
- Can other family members be tested?
- How many people have MCADD?
- Does MCADD happen more often in a certain ethnic group?
- Does MCADD go by any other names?
- Where can I find more information?

This fact sheet has general information about MCADD. Every child is different and some of this information may not apply to your child specifically. Certain treatments may be recommended for some children but not others. Children with MCADD should be followed by a metabolic doctor in addition to their primary doctor.

## What is MCADD?

MCADD stands for “medium chain acyl-CoA dehydrogenase deficiency”. It is one type of fatty acid oxidation disorder. People with MCADD have problems breaking down fat into energy for the body.

## Fatty Acid Oxidation Disorders:

Fatty acid oxidation disorders (FAODs) are a group of rare inherited conditions. They are caused by enzymes that do not work properly.

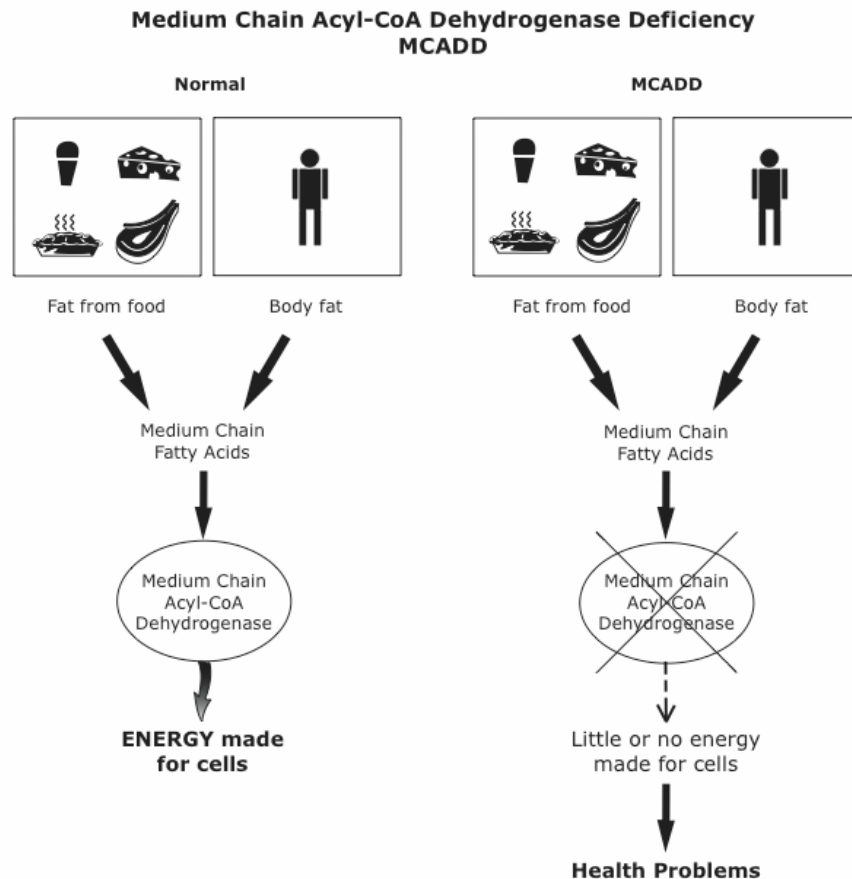
A number of enzymes are needed to break down fats in the body (a process called fatty acid oxidation). Problems with any of these enzymes can cause a fatty acid oxidation disorder. People with FAODs cannot properly break down fat from either the food they eat or from fat stored in their bodies.

The symptoms and treatment vary between different FAODs. They can also vary from person to person with the same FAOD. See the fact sheets for each specific FAOD.

FAODs are inherited in an autosomal recessive manner and affect both males and females.

## What causes MCADD?

MCADD occurs when an enzyme, called “medium chain acyl-CoA dehydrogenase” (MCAD), is either missing or not working properly. This enzyme’s job is to break down certain fats in the food we eat into energy. It also breaks down fat already stored in the body.



Energy from fat keeps us going whenever our bodies run low of their main source of energy, a type of sugar called glucose. Our bodies rely on fat when we don't eat for a stretch of time – like when we miss a meal or when we sleep.

When the MCADD enzyme is missing or not working, the body cannot use certain types of fat for energy, and must rely solely on glucose. Although glucose is a good source of energy, there is a limited amount available. Once the glucose has been used up, the body tries to use fat without success. This leads to low blood sugar, called hypoglycemia, and to the build up of harmful substances in the blood.

## **If MCADD is not treated, what problems occur?**

MCADD can cause bouts of illness called metabolic crises. Children with MCADD often show effects for the first time between 3 months and 2 years of age. Some of the first symptoms of a metabolic crisis are:

- extreme sleepiness
- behavior changes
- irritable mood
- poor appetite

Some of these other symptoms may also follow::

- fever
- diarrhea
- vomiting
- hypoglycemia

If a metabolic crisis is not treated, a child with MCADD can develop:

- breathing problems
- seizures
- coma, sometimes leading to death

Between episodes of metabolic crisis, people with MCADD are usually healthy. However, repeated episodes can cause permanent brain damage. This may result in learning problems, mental retardation or spasticity.

Symptoms often happen after having nothing to eat for more than a few hours. Hypoglycemia can occur, with or without other symptoms of metabolic crisis, just by going too long without food. Hypoglycemia can cause a person to feel weak, shaky or dizzy, and have clammy, cold skin. If not treated, hypoglycemia can lead to coma and even death.

Hypoglycemia and metabolic crises are also more likely to occur when a person with MCADD gets sick or has an infection.

Some children with MCADD never have symptoms and are only found to be affected after a brother or sister is diagnosed.

## **What is the treatment for MCADD?**

Your baby's primary doctor will work with a metabolic doctor to care for your child. Your doctor may also suggest that you meet with a dietician familiar with MCADD.

Certain treatments may be advised for some children but not others. When necessary, treatment is usually needed throughout life. The following are treatments often recommended for children with MCADD:

### **1. Avoid going a long time without food**

Babies and young children with MCADD need to eat often to avoid hypoglycemia or a metabolic crisis. They should not go without food for more than 4 to 6 hours. Some babies need to eat even more often than this. It is important that babies be fed during the night. They need to be woken up to eat if they do not wake up on their own.

Young children with MCADD may need to have a starchy snack before bed and another during the night. They may need another snack first thing in the morning. Raw cornstarch mixed with water, milk or other drink is a good source of long-lasting energy. This is sometimes suggested for children older than one year of age. Your dietician can give you ideas for good night-time snacks.

When they are well, most teens and adults with MCADD can go without food for up to 12 hours without problems. They do need to continue the other treatments throughout life.

### **2. Diet**

Sometimes a low fat, high carbohydrate diet is recommended. Carbohydrates give the body many types of sugar that can be used as energy. In fact, for children needing this treatment, most food in the diet should be carbohydrates (bread, pasta, fruit, vegetables, etc.) and protein (lean meat and low-fat dairy foods). Your dietician can create a food plan with the correct type and amount of fat your child needs. Any diet changes should be made under the guidance of a dietician.

Ask your doctor whether or not your child needs to have any changes in his or her diet.

### **3. L-carnitine**

Some children may be helped by taking L-carnitine. This is a safe and natural substance that helps body cells make energy. It also helps the body get rid of harmful wastes.

Your doctor will decide whether or not your child needs L-carnitine. Unless you are advised otherwise, use only L-carnitine prescribed by your doctor. Do not use any medications without checking with your doctor.

#### **4. Call your doctor at the start of any illness**

Always call your health care provider when your child has any of the following:

- poor appetite
- low energy or excessive sleepiness
- vomiting
- diarrhea
- an infection
- a fever

People with MCADD need to eat extra starchy food and drink more fluids during any illness - even if they may not feel hungry – or they could develop hypoglycemia or a metabolic crisis. Children who are sick often don't want to eat. If they won't or can't eat, they may need to be treated in the hospital to prevent problems.

Ask your metabolic doctor if you should carry a special travel letter with medical instructions for your child's care.

### **What happens when MCADD is treated?**

With prompt and careful treatment, children with MCADD usually live healthy lives with typical growth and development. The goal of treatment is to prevent long-term problems. However, children who have repeated metabolic crises may have life-long learning disabilities, spasticity, or other effects.

### **What causes the MCAD enzyme to be absent or not working correctly?**

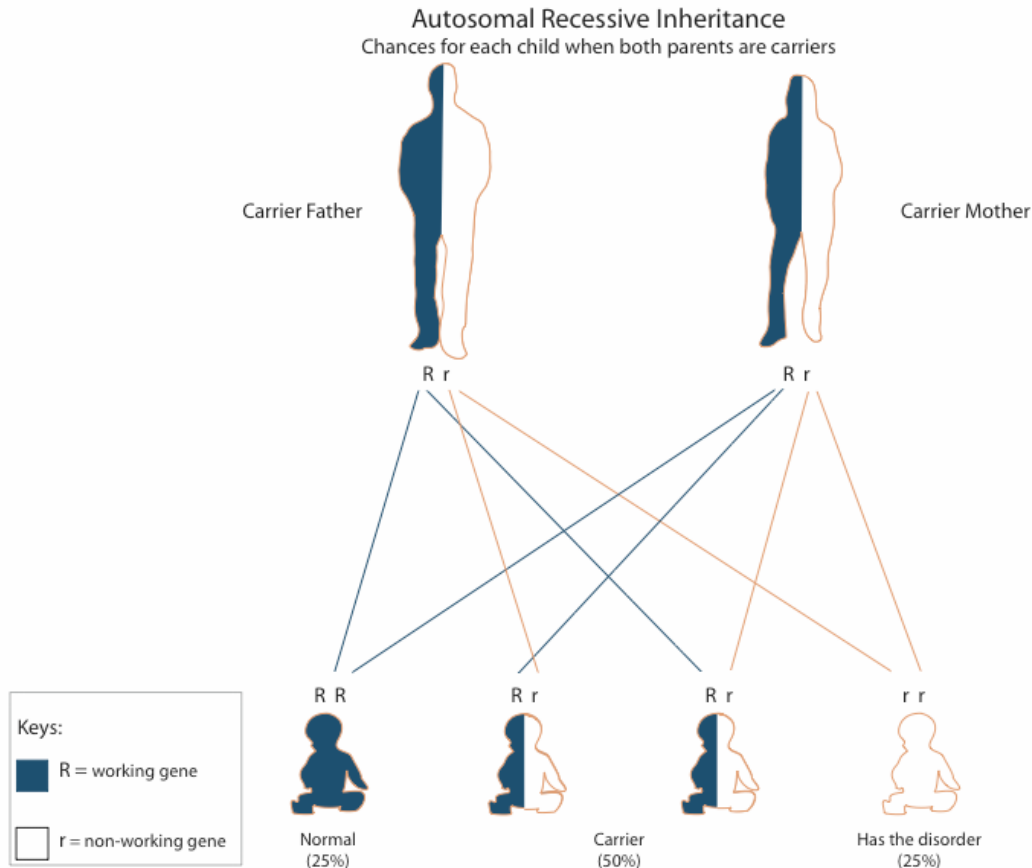
Genes tell the body to make various enzymes. People with MCADD have a pair of genes that do not work correctly. Because of the changes in this pair of genes, the MCAD enzyme either does not work properly or is not made at all.

### **How is MCADD inherited?**

MCADD is inherited in an autosomal recessive manner. It affects both boys and girls equally.

Everyone has a pair of genes that make the MCAD enzyme. In children with MCADD, neither of these genes works correctly. These children inherit one non-working gene for the condition from each parent.

Parents of children with MCADD rarely have the disorder. Instead, each parent has a single non-working gene for MCADD. They are called carriers. Carriers do not have MCADD because the other gene of this pair is working correctly. When both parents are carriers, there is a 25% chance in each pregnancy for the child to have MCADD. There is a 50% chance for the child to be a carrier, just like the parents. And, there is a 25% chance for the child to have two working genes.



Genetic counseling is available to families who have children with MCADD. Genetic counselors can answer your questions about how MCADD is inherited, choices during future pregnancies, and how to test other family members. Ask your doctor about a referral to a genetic counselor.

### Is genetic testing available?

Genetic testing for MCADD can be done on a blood sample. Genetic testing, also called DNA testing, looks for changes in the pair of genes that cause MCADD. In many children with MCADD, both gene changes can be found. However, in other

children, only one or neither of the two gene changes can be found, even though we know they are present.

DNA testing is not necessary to diagnose your child. It can be helpful for carrier testing or prenatal diagnosis, discussed below.

## **What other testing is available?**

MCADD can also be confirmed either by a blood test called an acylcarnitine profile or an enzyme test on a skin sample. Talk to your doctor or your genetic counselor if you have questions about testing for MCADD.

## **Can you test during pregnancy?**

If both gene changes have been found in your child with MCADD, DNA testing can be done during future pregnancies. The sample needed for this test is obtained by either CVS or amniocentesis.

If DNA testing would not be helpful, an enzyme test can be done during pregnancy on cells from the fetus. Again, the sample needed for this test is obtained by either CVS or amniocentesis.

Parents may either choose to have testing during pregnancy or wait until birth to have the baby tested. A genetic counselor can talk to you about your choices and answer questions about prenatal testing or testing your baby after birth.

## **Can other members of the family have MCADD or be carriers?**

### **Having MCADD**

The brothers and sisters of a baby with MCADD have a chance of being affected, even if they have had no symptoms. Finding out whether other children in the family have MCADD is important because early treatment may prevent serious health problems. Talk to your doctor or genetic counselor about testing your other children for MCADD.

### **MCADD Carriers**

Brothers and sisters who do not have MCADD still have a chance to be carriers like their parents. Except in special cases, carrier testing should only be done on people over 18 years of age.

Each of the parents' brothers and sisters has a 50% chance to be an MCADD carrier. It is important for other family members to be told that they could be carriers. There is a small chance they are also at risk to have children with MCADD.

Some states do not offer newborn screening for MCADD. However, expanded newborn screening through private labs is available for babies born in states that do not screen for this condition. To learn more about expanded newborn screening, see [How to obtain MS/MS](#).

When both parents are carriers, newborn screening results are not sufficient to rule out MCADD in a newborn baby. In this case, special diagnostic testing should be done in addition to newborn screening.

## **Can other family members be tested?**

### **Diagnostic testing**

Brothers and sisters can be tested for MCADD using DNA testing or other special tests.

### **Carrier Testing**

If both gene changes have been found in your child with MCADD, other family members can have DNA testing to see if they are carriers.

If DNA testing would not be helpful, carrier testing can also be done by an enzyme test on a skin sample.

## **How many people have MCADD?**

About one in every 15,000 babies in the United States is born with MCADD.

## **Does MCADD happen more often in a certain ethnic group?**

MCADD happens more often in white people from Northern Europe and the United States. About 1 in every 70 Caucasians is a carrier for MCADD.

## **Does MCADD go by any other names?**

MCADD is also sometimes called:

- MCAD deficiency
- ACADM deficiency
- MCADH deficiency

## Where can I find more information?

Fatty Oxidation Disorders (FOD) Family Support Group

<http://www.fodsupport.org>

Organic Acidemia Association

<http://www.oaanews.org>

United Mitochondrial Disease Foundation

<http://www.umdf.org>

Children Living with Inherited Metabolic Diseases (CLIMB)

<http://www.climb.org/uk>

Genetic Alliance

<http://www.geneticalliance.org>

PacNoRGG pamphlet: MCAD

[http://mchneighborhood.ichp.edu/pacnorgg/media/Metabolic/mcad\\_eng.pdf](http://mchneighborhood.ichp.edu/pacnorgg/media/Metabolic/mcad_eng.pdf)

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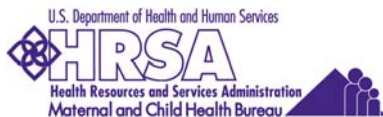
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