



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: Carnitine acylcarnitine translocase deficiency

Acronym: CAT

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This fact sheet has general information about CAT deficiency. Every child is different and some of this information may not apply to your child specifically. Not all is known about CAT deficiency and, at present, there is no standard treatment plan. There may be treatments recommended for some children but not others. Children with CAT deficiency should be followed by a metabolic doctor in addition to their primary doctor.

What is CAT deficiency?

CAT deficiency stands for “carnitine acylcarnitine translocase deficiency”. It is one type of fatty acid oxidation disorder. People with CAT deficiency 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.

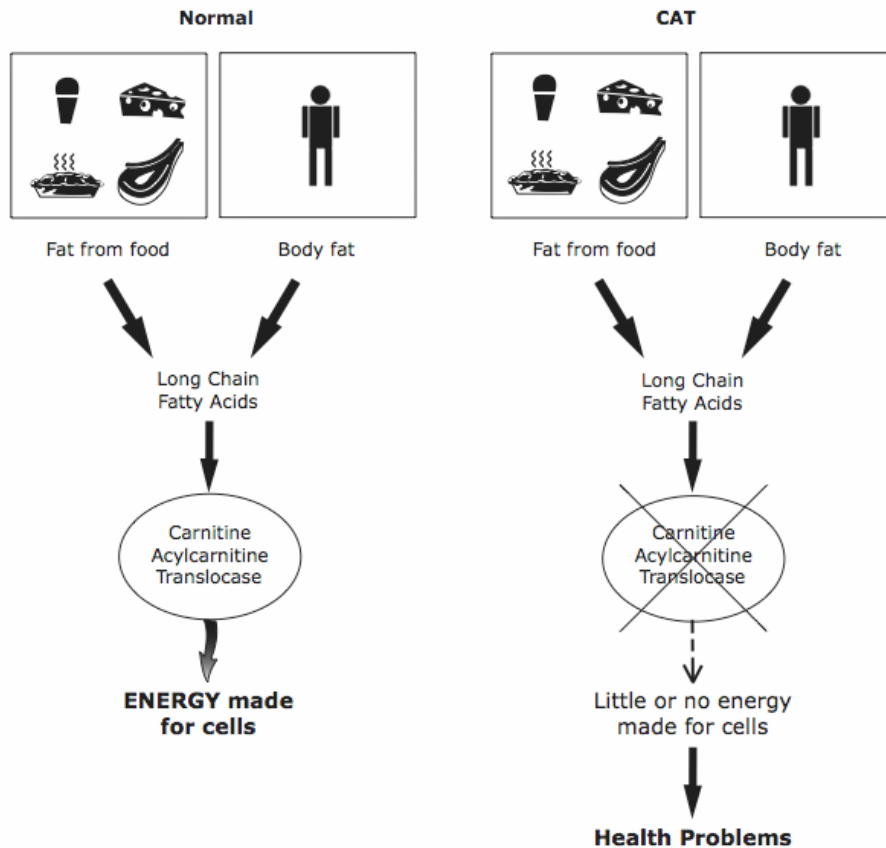
The symptoms and treatment vary between different FAODs. They can also vary from person to person with the same FAOD. See the fact sheets for each specific FAOD.

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

What causes CAT deficiency?

CAT deficiency occurs when an enzyme, called “carnitine acylcarnitine translocase” (CAT), is either missing or not working properly. This enzyme’s job is to help change certain fats in the food we eat into energy. It also helps to break down fat already stored in the body.

Carnitine Acylcarnitine Translocase Deficiency CAT



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 for energy when we don't eat for a stretch of time – like when we miss a meal or when we sleep.

When the CAT enzyme is missing or not working, the body cannot use 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 CAT deficiency is not treated, what problems occur?

CAT deficiency causes episodes of illness called metabolic crises. Some of the first symptoms of a metabolic crisis are:

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

Other symptoms then follow:

- fever
- diarrhea
- vomiting
- hypoglycemia

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

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

There are two forms of CAT deficiency. The most common type happens in newborns. A milder, less common type happens in older infants and children.

CAT deficiency in newborns

Newborns with CAT deficiency often show symptoms within the first week of life. Episodes of metabolic crisis are common. High levels of ammonia in the blood can occur. This can cause serious brain damage.

Newborns with CAT may also have:

- low muscle tone (floppy muscles and joints) and muscle weakness
- enlarged liver
- heart problems and enlarged heart
- breathing problems

Babies who are not treated usually die of heart problems, breathing problems, liver failure or high levels of ammonia in the blood.

CAT deficiency in children

Children with the mild type of CAT deficiency usually start having symptoms before age 3. Children are at risk to have episodes of metabolic crisis, but usually do not have heart problems.

In both types of CAT deficiency, symptoms often happen after having nothing to eat for more than a few hours. Symptoms are also more likely when a person with CAT deficiency gets sick or has an infection.

What is the treatment for CAT deficiency?

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 CAT deficiency.

Certain treatments may be helpful for some children but not others. When necessary, treatment is usually needed throughout life. The following are treatments sometimes recommended for children with CAT deficiency:

1. Avoid going a long time without food

Babies and young children with CAT deficiency need to eat often to avoid having 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 CAT deficiency should 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 CAT deficiency 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 advised. 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 food). Any diet changes should be made under the guidance of a dietician.

People with CAT deficiency cannot use particular building blocks of fat called "long chain fatty acids". A dietician can help you create a food plan low in these fats.

Ask your doctor whether your child needs to have any changes in his or her diet.

3. L-carnitine and MCT oil

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 L-carnitine without checking with your doctor.

Medium Chain Triglyceride oil (MCT oil) is sometimes used as part of the food plan for people with CAT deficiency. This special oil has medium chain fatty acids that people with CAT deficiency can use in small amounts for energy. Your

metabolic doctor or dietician can guide you in how to use this supplement. You will need to get a prescription from your doctor to get MCT oil.

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
- persistent muscle pain or weakness

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

Unless you are advised otherwise, use only L-carnitine prescribed by your doctor.

What happens when CAT deficiency is treated?

Prompt and careful treatment may help prevent or control symptoms in children with CAT deficiency. However, some children continue to have metabolic crises and other health problems despite treatment. Even with treatment, there is a risk of death, especially in newborns with symptoms.

What causes the CAT enzyme to be absent or not working correctly?

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

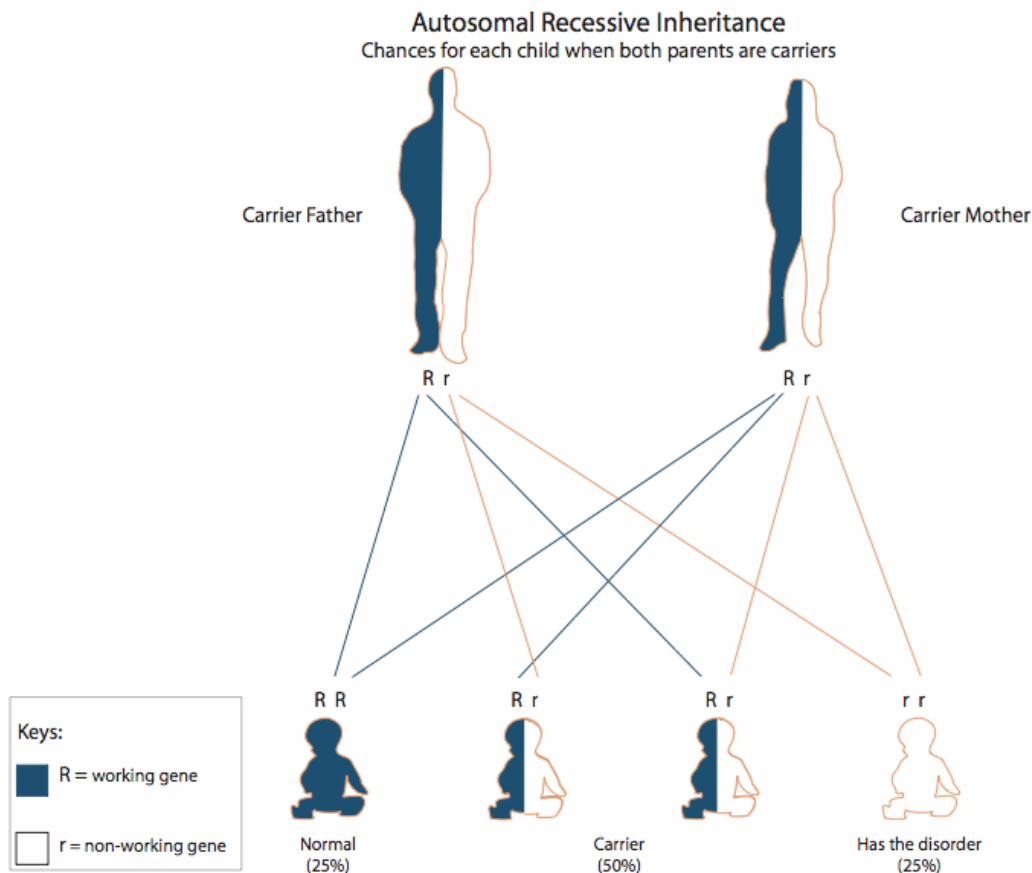
How is CAT deficiency inherited?

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

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

Parents of children with CAT deficiency rarely have the disorder. Instead, each parent has a single non-working gene for CAT deficiency. They are called carriers. Carriers do not have the condition 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 CAT deficiency. 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 CAT deficiency. Genetic counselors can answer your questions about how CAT deficiency is inherited, options 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 CAT deficiency can be done on a blood sample. Genetic testing, also called DNA testing, looks for changes in the pair of genes that cause CAT deficiency. In some affected children, both gene changes can be found. However, in other children, neither or only one 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?

CAT deficiency can also be confirmed by a special enzyme test on a skin sample. Talk to your doctor or genetic counselor if you have questions about testing for CAT deficiency.

Can you test during pregnancy?

If both gene changes have been found in the child with CAT deficiency, 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, testing during pregnancy can be attempted by performing special tests on fetal cells. Again, the sample needed for these tests is obtained by either CVS or amniocentesis.

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

Can other members of the family have CAT deficiency or be carriers?

CAT deficiency

The brothers and sisters of an affected baby have a chance of having CAT deficiency, even if they haven't had symptoms. Finding out whether other children in the family have CAT deficiency is important because early treatment may prevent serious health problems. Talk to your doctor or genetic counselor about testing your other children.

CAT deficiency carriers

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

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

Some states do not offer newborn screening for CAT deficiency. However, expanded newborn screening is available through private labs 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 CAT deficiency 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 for CAT deficiency

Brothers and sisters of an affected child can be tested for CAT deficiency using either DNA testing or a special enzyme test.

Carrier testing

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

If DNA testing would not be helpful, other methods of carrier testing may be available. Your metabolic doctor or genetic counselor can answer your questions about carrier testing.

How many people have CAT deficiency?

CAT deficiency is very rare. The actual incidence is unknown.

Does CAT deficiency happen more often in a certain ethnic group?

No, CAT deficiency does not happen more often in any specific race, ethnic group, geographical area or country.

Does CAT deficiency go by any other names?

CAT deficiency is also known as:

- carnitine acylcarnitine carrier (CAC)
- CACT deficiency

Where can I find more information?

Fatty Oxidation Disorders (FOD) Family Support Group

www.fodsupport.org

Organic Acidemia Association

www.oaanews.org

United Mitochondrial Disease Foundation

www.umdf.org

Children Living with Inherited Metabolic Disorders (CLIMB)

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

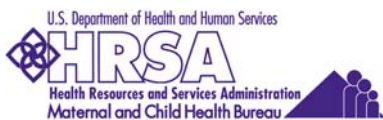
Genetic Alliance

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