

Tools Enabling Metabolic Parents LEarning

ADAPTED BY THE DIETITIANS GROUP

BIMDG

British Inherited Metabolic Diseases Group

CACT Deficiency

BASED ON THE ORIGINAL TEMPLE WRITTEN BY BURGARD AND WENDEL VERSION 4, MARCH 2025



TEMPLE foreword

TEMPLE (Tools Enabling Metabolic Parents LEarning) are a set of teaching slides and booklets that provide essential information about different inherited metabolic disorders that require special diets as part of their management. These teaching tools are aimed at parents who may have an infant or child that has been recently diagnosed with a disorder. They are also useful when teaching children, extended family members, child minders, nursery workers and a school team.

They have been developed by a team of experienced clinical and research metabolic dietitians from the UK who are members of the British Inherited Metabolic Disease Group (BIMDG).

The team are Rachel Skeath, Karen van Wyk, Pat Portnoi and Anita MacDonald. The group is facilitated by Heidi Chan from Nutricia.

Each module produced is reviewed by a consultant clinician who is a member of the BIMDG.

This teaching tool is not designed to replace dietary information that may be given by a dietitian in clinic.

Carnitine Acylcarnitine Translocase Deficiency

Information for families following a new diagnosis



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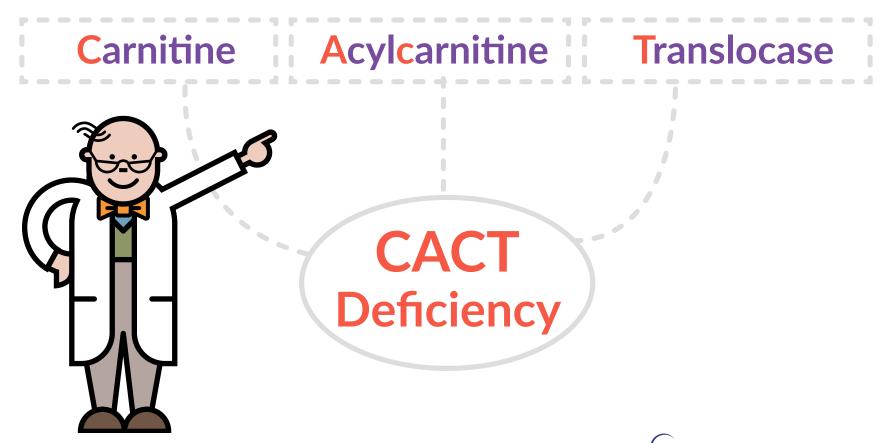




What is CACT deficiency?

CACT stands for Carnitine Acylcarnitine Translocase deficiency

It is an inherited metabolic condition



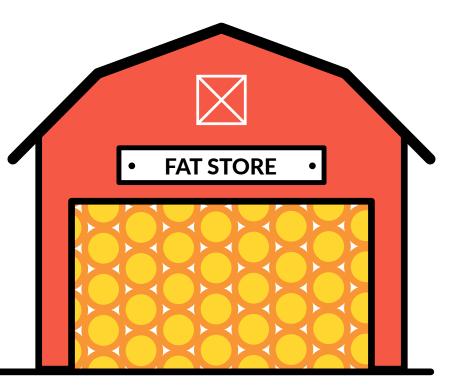
Which foods supply the body with energy?

There are two main food groups that supply the body with energy:

 Carbohydrates (starches and sugars) provide a readily available energy source

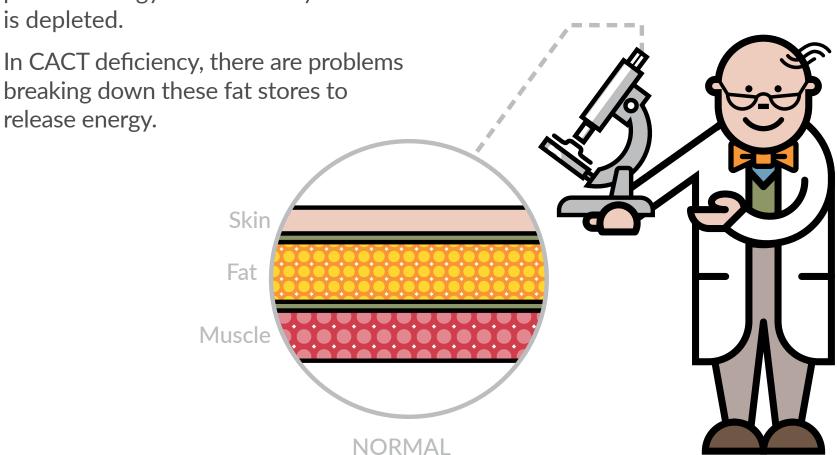
 Fats also provide energy. Fat is stored in the body so it can be used as an energy reserve





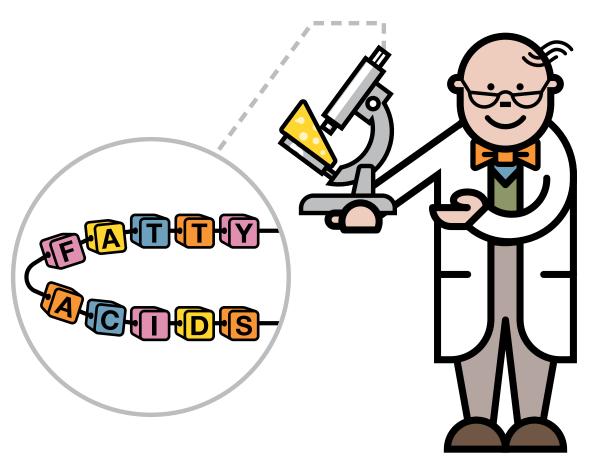
CACT deficiency and fat

The body uses its own fat stores to provide energy when carbohydrate is depleted.



Breaking down fat stores for energy

Body fat stores are broken down into fatty acids.



What are fatty acids?

Fatty acids are made up of carbon atoms joined together to form chains of many different lengths.



Short chain



Medium chain



Long chain

Fatty acid transport

Fatty acid chains need to be transported into the cells of the body. This enables the body to produce energy in a form which it can use.



What happens in CACT deficiency?

In CACT deficiency, the body lacks a chemical (enzyme) that helps convert fat stores into energy.

This **stops** the long chain fats from being used for energy.

This is a problem when it is necessary to break down fats quickly.

This causes a shortage of energy supply.

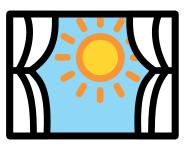


What can go wrong in CACT deficiency in infants?

There can be a shortage of energy supply and a build up of harmful chemicals with illness or lack of food.

Symptoms include:

- low blood sugar
- floppiness
- seizures
- heart problems
- irregular heartbeat
- muscle weakness
- liver problems

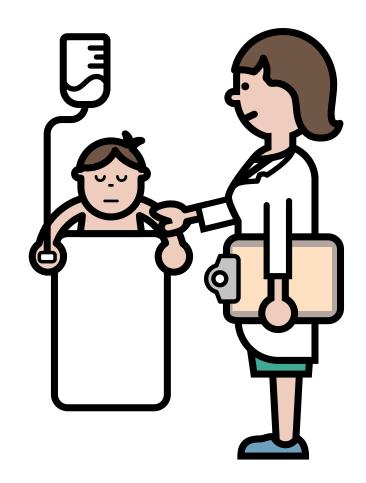




What can go wrong in CACT deficiency?

If there is a shortage of energy and this is not corrected, it can lead to coma and brain damage and it may be life threatening.

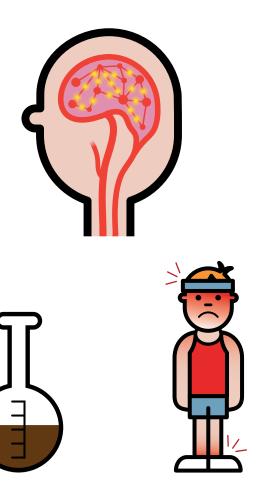
However, please remember, this can all be prevented with timely and correct management.



Some children may be diagnosed in early or later childhood

Signs and symptoms:

- Low blood sugars
- It may affect development and learning
- Exercise may cause muscle pain which may lead to reddish brown coloured urine
- Reddish-brown coloured urine



Metabolic crisis

- A metabolic crisis triggers the CACT deficiency symptoms
- This leads to a lack of energy and build up of toxic chemicals
- It is usually triggered by childhood infections causing high temperatures, vomiting, and diarrhoea
- It can also be caused by going a very long time without food
- Avoidance of a metabolic crisis is essential



How is CACT deficiency diagnosed?

CACT deficiency is suspected because of the pattern of chemicals (acylcarnitines and organic acids) found in the blood and urine.

The diagnosis is confirmed by finding mutations in the CACT gene.





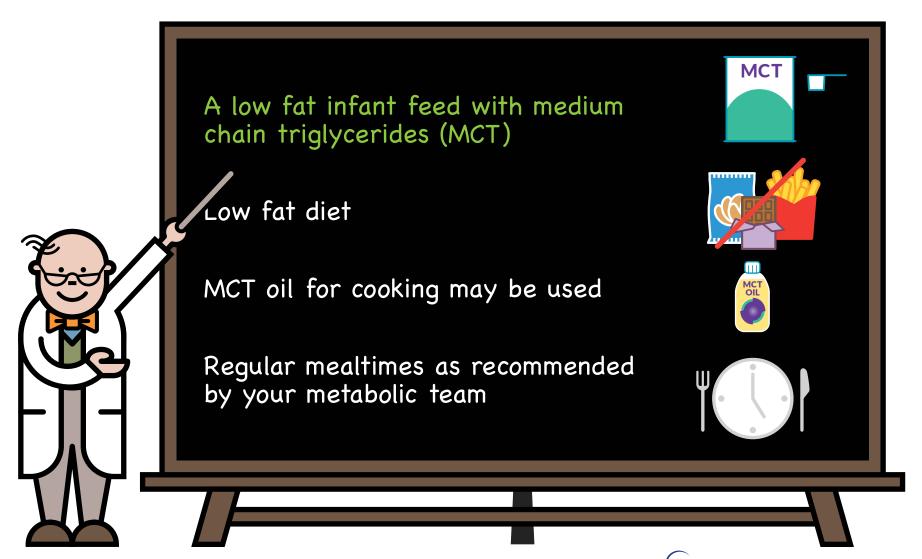
How is CACT deficiency managed?

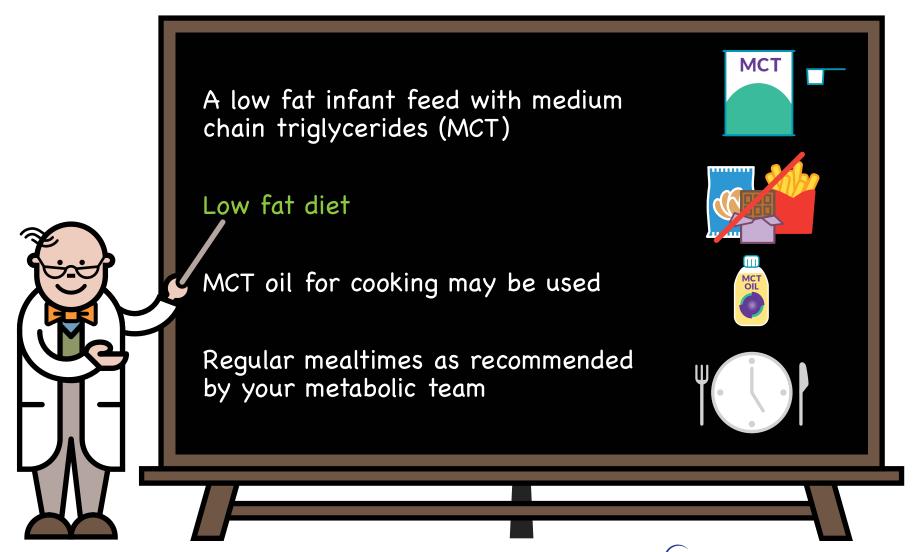
In babies, it is managed day to day by avoiding long periods without feeding, even when well.

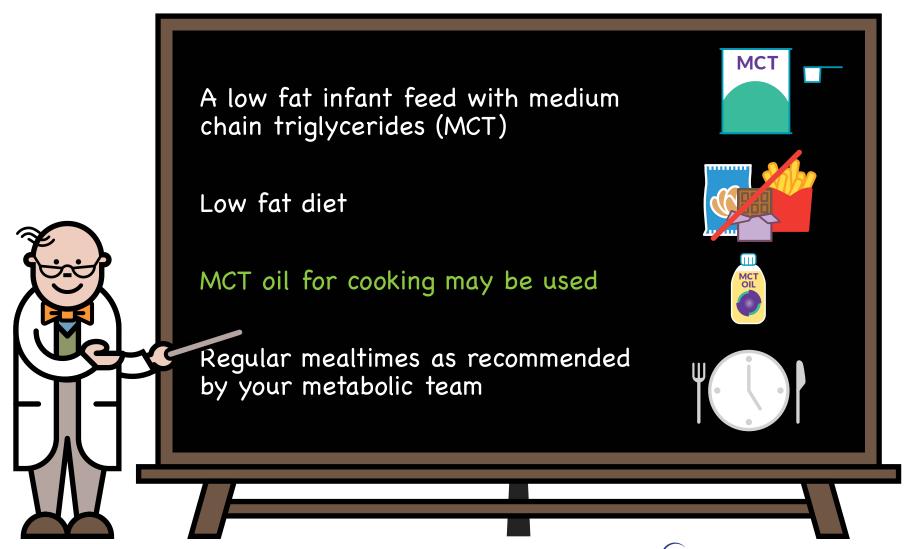
The length of time babies can go without feeds is known as **the safe fasting time**.

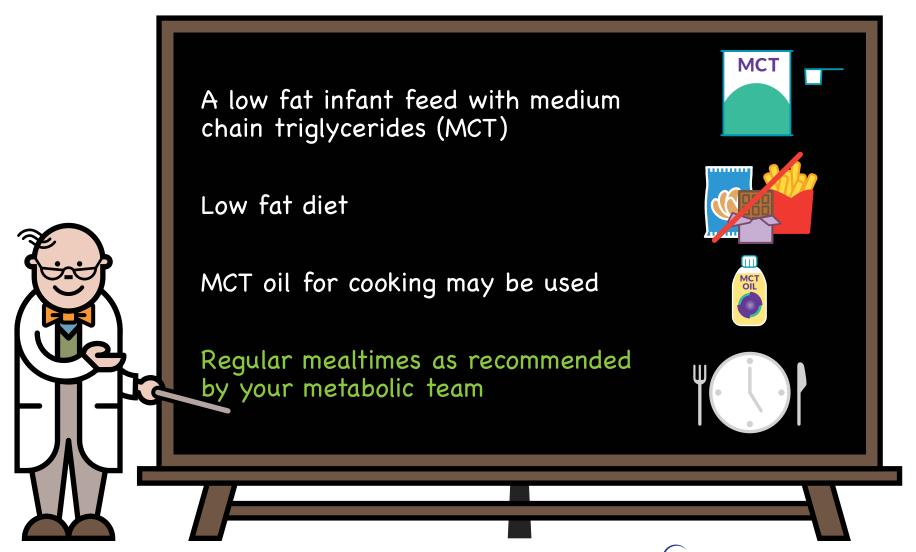
SAFE FASTING TIMES

- The safe fasting time varies for each baby with CACT deficiency. Your metabolic team will advise.
- It is important they receive regular feeds during the day and at night.
- They should not miss scheduled feeds.









CACT deficiency and fat

The diet needs to be low in fat.

Foods high in fat are avoided.

Many foods are high in fat e.g. full fat milk, full fat cheese, fatty meat, eggs, ice cream, chips, crisps and chocolate.

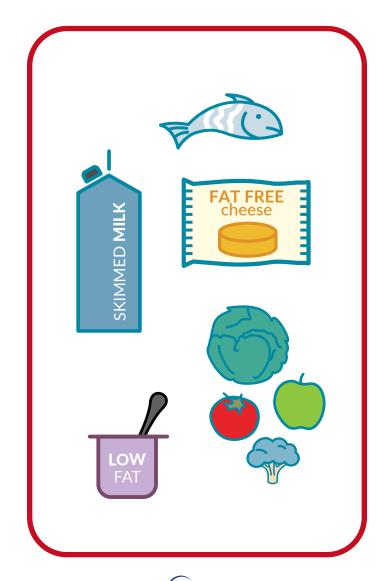


Low fat foods

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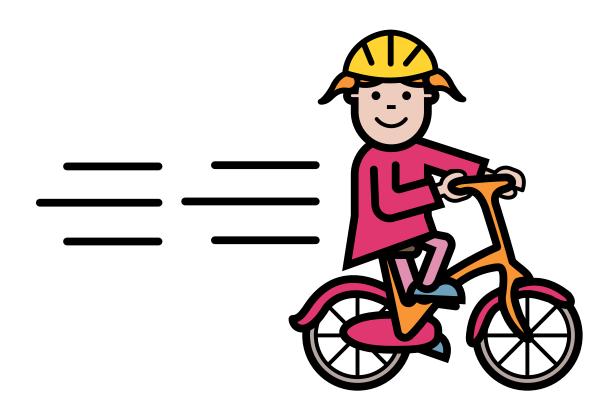
Low fat foods are permitted.

There is a wide range of low fat foods available e.g. skimmed milk, low fat yoghurt/cheese, white fish, fruit and vegetables.



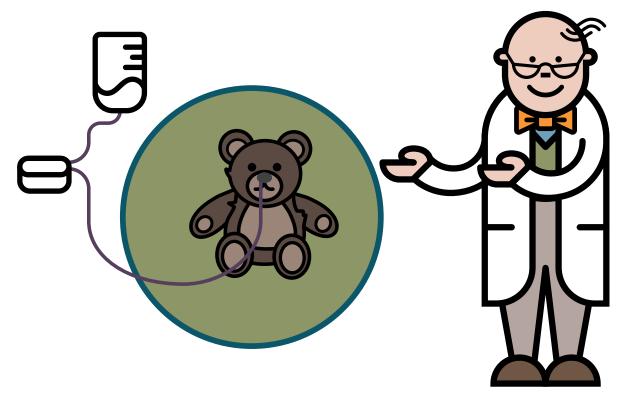
CACT deficiency and exercise

It is advisable to take a high sugar snack or drink before exercise. This will supply an extra source of energy.

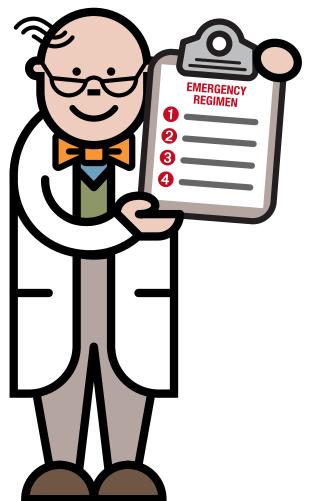


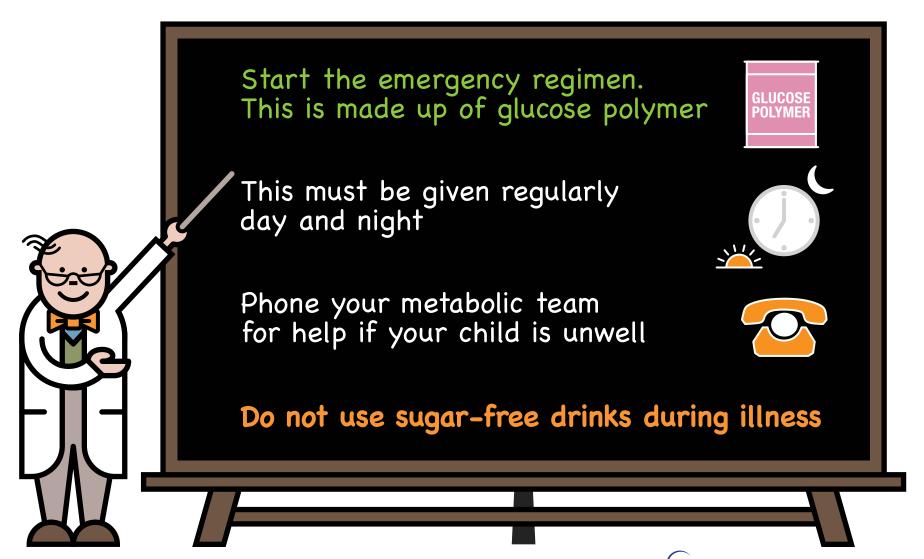
Is tube feeding needed?

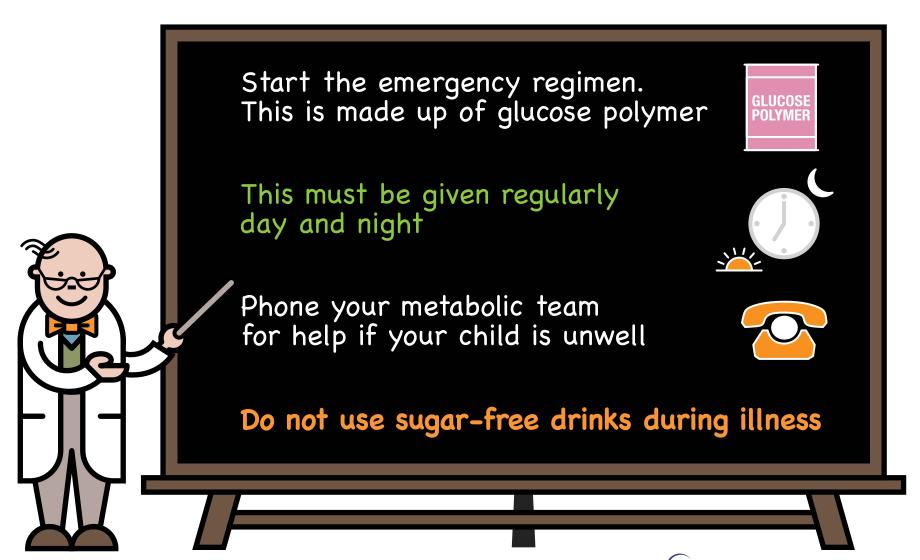
In babies with the most severe forms of CACT deficiency, tube feeding may be necessary. This will ensure energy, nutrient and fluid needs are met. In children not on tube feeds, a late night snack containing carbohydrate may be necessary. This is to limit fasting time.

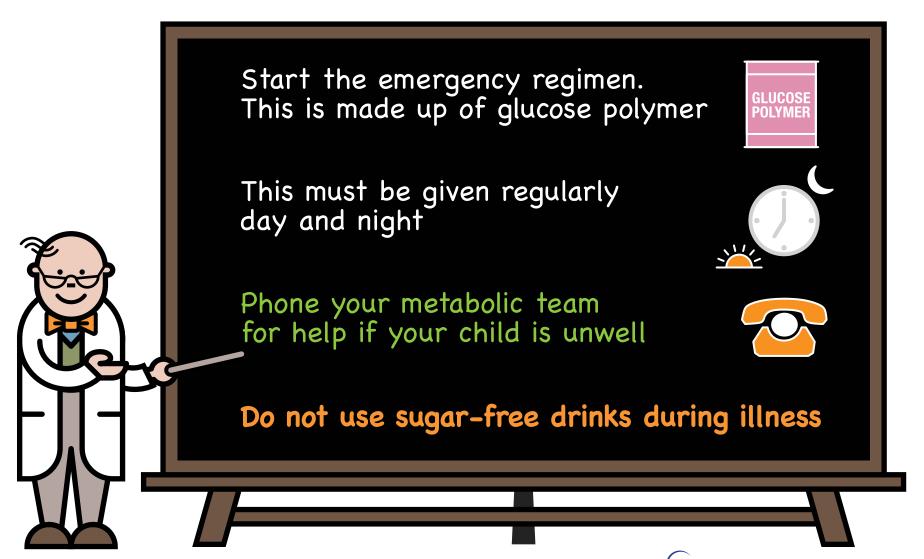


- During any childhood illness, an emergency regimen is given
- This provides energy and prevents build up of harmful chemicals that cause a metabolic crisis









Checklist for illness



Checklist for illness

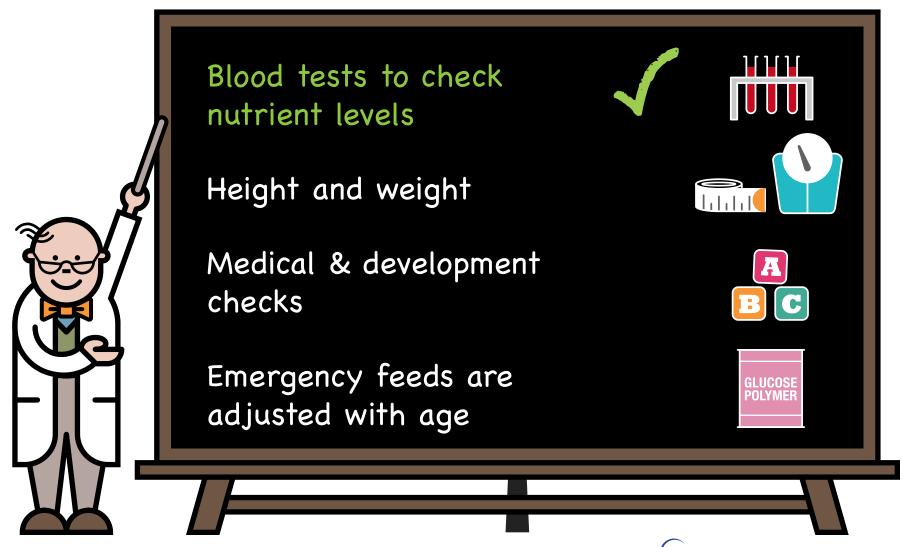


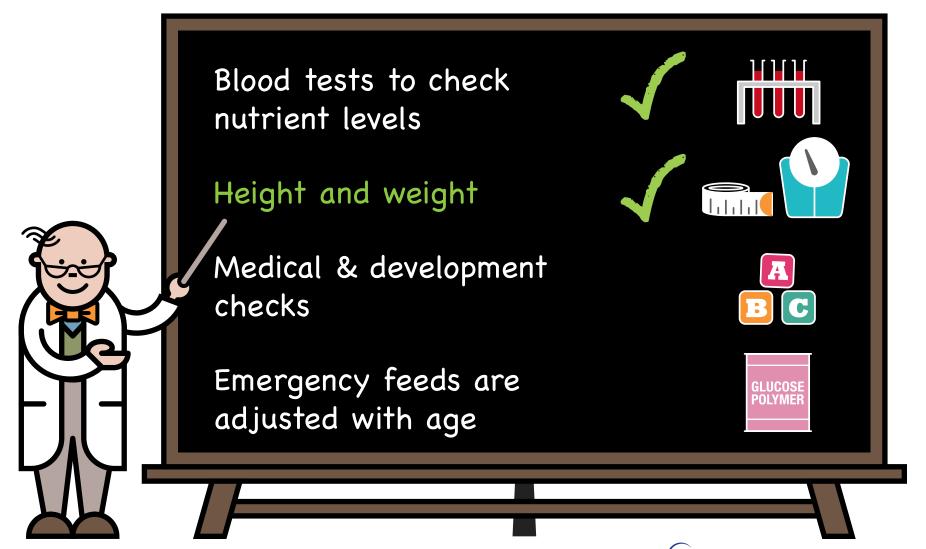
Checklist for illness

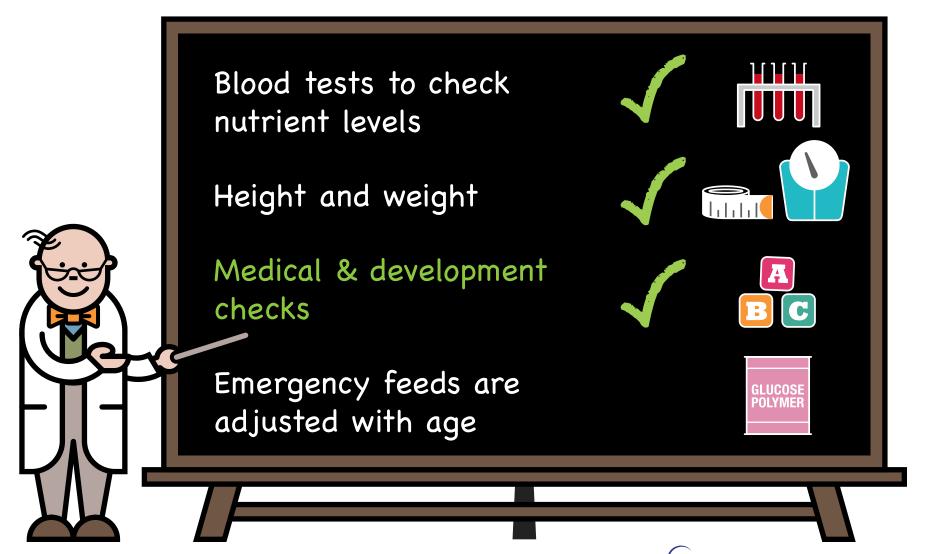


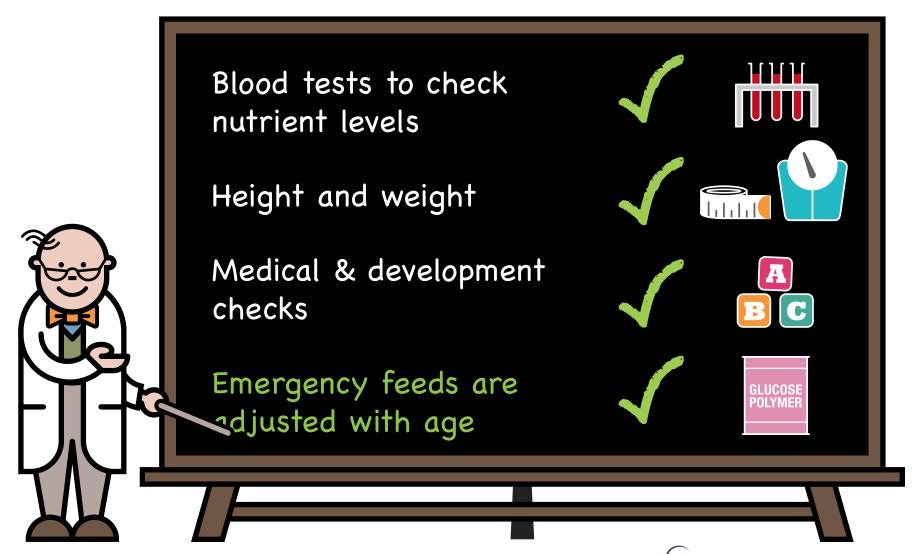
Key message

It is imperative that emergency feeds are started promptly and there are no delays in management.











Humans have chromosomes composed of DNA



Genes are pieces of DNA that carry the genetic instructions. Each chromosome may have several thousand genes

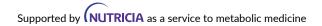


The word mutation means a change or error in a genetic instruction



We inherit particular chromosomes from the egg of the mother and sperm of the father







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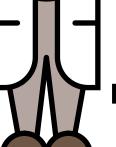




CACT deficiency is an inherited condition. There is nothing that could have been done to prevent your baby from having CACT deficiency

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

Parents of children with CACT deficiency are carriers of the condition





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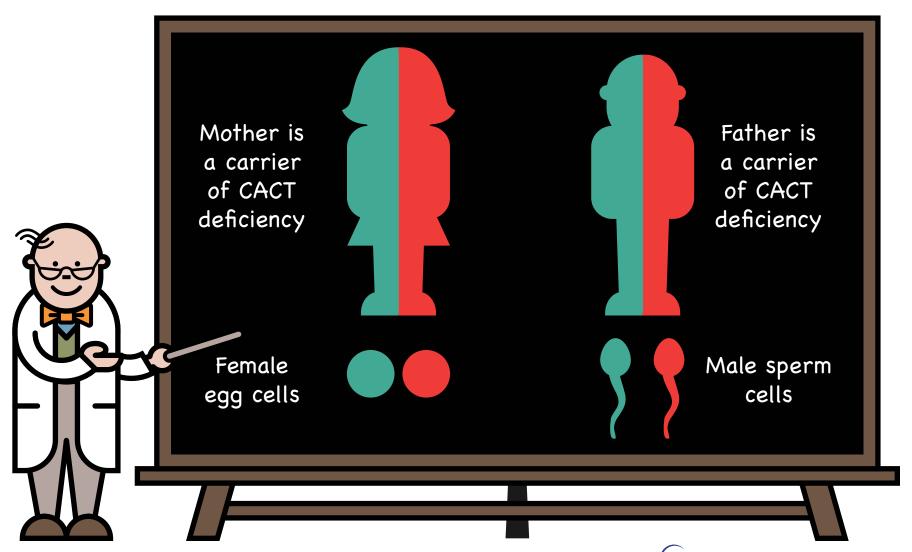
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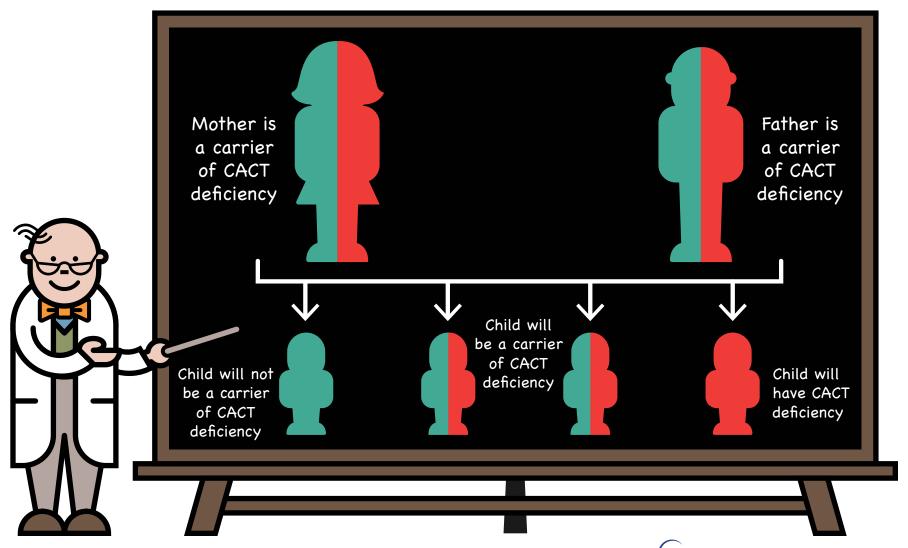
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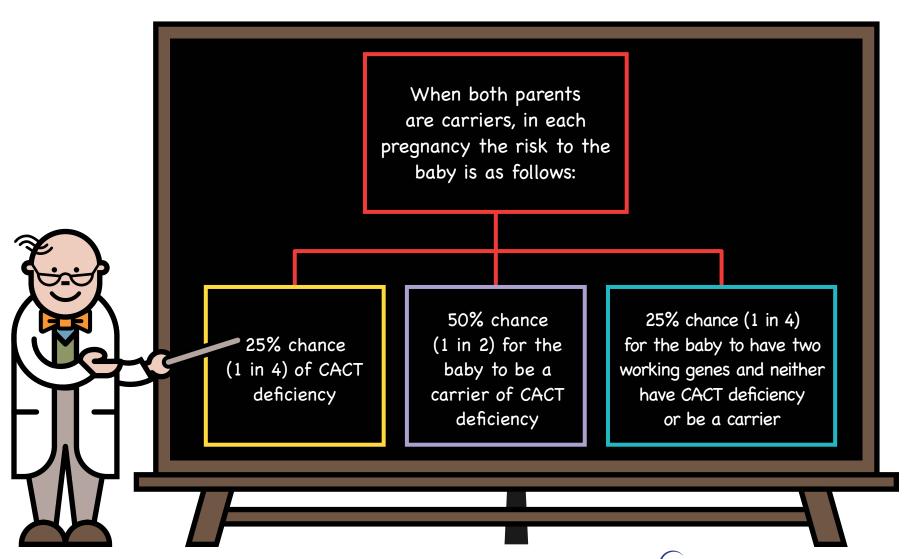
Inheritance – Autosomal recessive (carriers of CACT deficiency)



Inheritance – Autosomal recessive – possible combinations



Future pregnancies



Take home messages

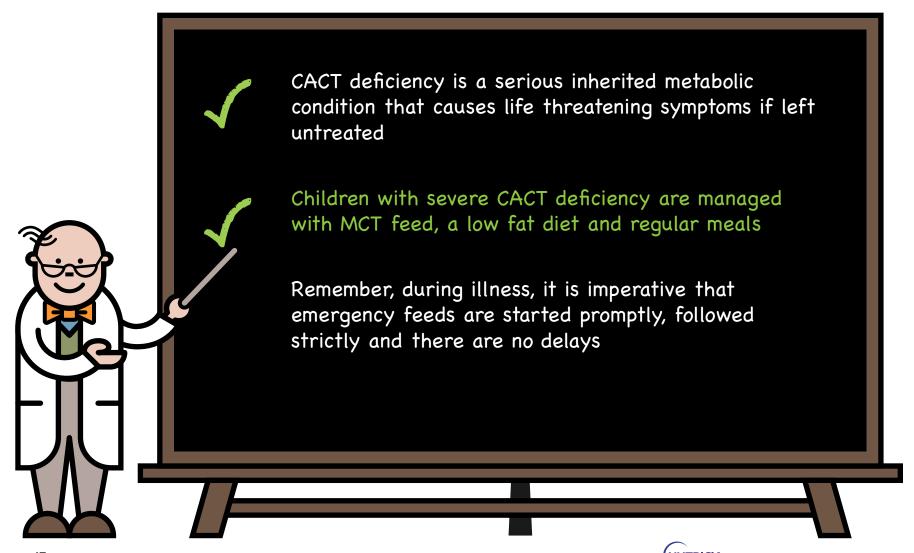


CACT deficiency is a serious inherited metabolic condition that causes life threatening symptoms if left untreated

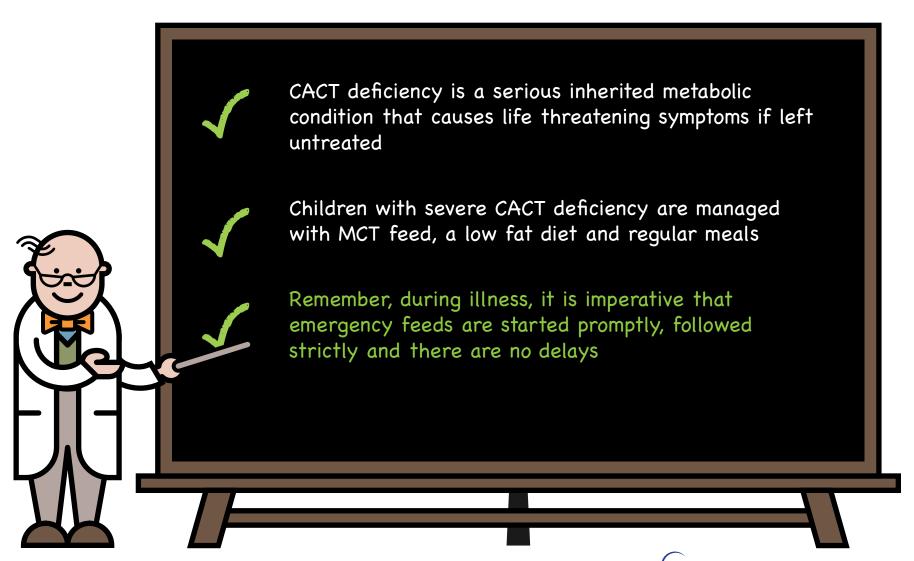
Children with severe CACT deficiency are managed with MCT feed, a low fat diet and regular meals

Remember, during illness, it is imperative that emergency feeds are started promptly, followed strictly and there are no delays

Take home messages



Take home messages



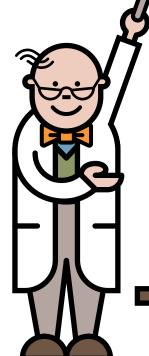


Always ensure you have a good supply of your emergency glucose polymer powder and it is in date

Special feeds and dietary products are prescribed by your GP and you obtain them from your pharmacy or home delivery

Medications to control fever should be given as normally recommended — always keep supplies available

Remember to keep contact numbers of your metabolic team to hand

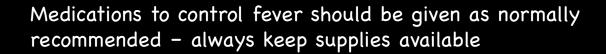




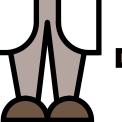
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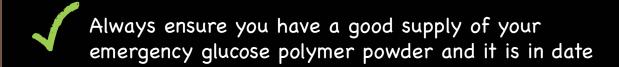


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Who's who

My dietitians

My nurses

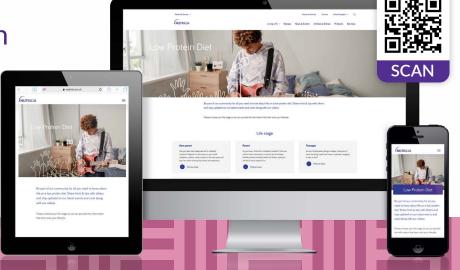
My doctors

- Contact details, address, photos

Visit www.nutricia.co.uk/patients-carers/living-with/low-protein-diet.html
and register to get access to support and practical advice for those living on a low protein diet.

The site also provides information on upcoming events and personal stories from others





BIMDG

British Inherited Metabolic Diseases Group





SUPPORT UK

Our common fight.

on a low protein diet.