

Carrying Beta Thalassaemia

(Also known as having Beta Thalassaemia Trait or having Beta Thalassaemia Minor)

- A carrier of Beta Thalassaemia is a healthy person.
- Carrying Beta Thalassaemia does not weaken them physically or mentally.
- They do not need any medical treatment because they carry Beta Thalassaemia.

What does it mean to carry Beta Thalassaemia?

Beta Thalassaemia is one of many possible variations in the blood called haemoglobin gene variants or haemoglobin variants.

Haemoglobin is what makes blood red. It is packed into red blood cells. Carriers of thalassaemia have smaller red blood cells, but more of them than other people.

A carrier will always be a carrier and no-one can catch it from them. They inherited Beta Thalassaemia from one of their parents and could pass it on to their children.

Anyone can carry Beta Thalassaemia. It is common in the Mediterranean area, the Middle East, the Indian subcontinent and South and South-East Asia. It also occurs among West Africans and African-Caribbeans. It occurs occasionally among North Europeans.

Can carrying Beta Thalassaemia cause any health problems?

Carrying Beta Thalassaemia is not an illness and will never turn into an illness. In fact, carriers are less likely than other people to catch malaria or suffer from heart attacks. Carriers can eat what they want and do any kind of work they choose.

Some carriers are slightly anaemic (have a lower haemoglobin level than usual). This anaemia has no effect on health or length of life.

Occasionally a doctor thinks a carrier must be short of iron because they have small red blood cells. If the doctor prescribes iron medicine, in the long run this could do more harm than good. A carrier should take iron medicine only if a special blood test (serum iron or serum ferritin) shows that they are short of iron.

Carriers can give blood provided that they are not anaemic.

Could a carrier of Beta Thalassaemia have children with a serious haemoglobin disorder?

Only if their partner also carries a haemoglobin variant. With medical help, such a couple can have healthy children.

What should a carrier do if they are thinking of having children?

They should tell their partner that they carry Beta Thalassaemia and ask him or her to have a blood test "for haemoglobin disorders". This test should be done before they start a pregnancy, or as soon as possible once a pregnancy has started. Their GP can arrange it.

If their partner is not a carrier, there is nothing to worry about.

What should they do if their partner is also a carrier?

They should ask their GP for an immediate appointment with a specialist counsellor. This is particularly important if they have already started a pregnancy. They can also contact the counselling service directly.

Is there anything else that a carrier should do?

If a carrier has brothers or sisters or already has children, they need to know that they may also carry Beta Thalassaemia. They should ask their GP or practice nurse for a blood test “for haemoglobin disorders”.

Further useful information

The NHS Screening Programmes website publishes a number of leaflets for patients about haemoglobin disorders, one of which is about being a Beta Thalassaemia carrier. They are available at: [Screening in pregnancy: information leaflets - GOV.UK \(www.gov.uk\)](https://www.gov.uk/government/publications/screening-in-pregnancy-information-leaflets)

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