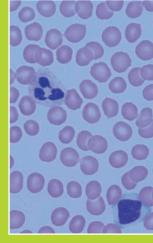
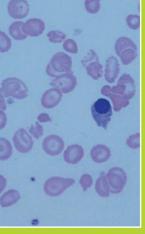
Treatment of Thalassaemic Patients

- Patients need monthly blood transfusions for their entire life.
- Frequent blood transfusions will result in iron accumulating in major organs such as the liver, spleen, heart and endocrine glands. This eventually results in multiple organ malfunction.
- Treatment to remove excess iron in the body has to be carried out 5 times a week.
- Bone marrow transplant is an option if there is a suitable donor in the family.

Thalassaemia Major Can be Prevented!

- Get tested if you:
 - 1. are a teenager
 - 2. have a family history of Thalassaemia
 - 3. are planning to get married
- If you are a thalassaemia carrier, avoid marrying another carrier.





Normal red blood cells

alassaemic blood cells

Your child's future is in your hands. Get tested for Thalassaemia.





For further information:

Please contact the nearest hospital, clinic or Thalassaemia Association



THALASSAEMIA What every Malaysian needs to know!



MINISTRY OF HEALTH MALAYSIA



Do you carry the thalassaemia gene? 1 in 20 Malaysians are carriers!



What is Thalassaemia?

Thalassaemia is a genetic disorder of the red blood cells. Thalassaemia is inherited from either or both parents and can occur in males and females.

There are two types of thalassaemia

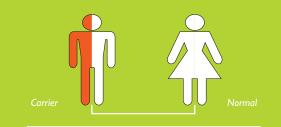
- 1. Thalassaemia carrier (Thalassaemia minor)
- 2. Thalassaemia patient (Thalassaemia major)

Thalassaemia Carrier

A thalassaemia carrier does not have any symptoms or health problems. The condition can only be detected through a special blood test. Carriers can pass on the thalassaemia gene to their children.

IF EITHER PARENT IS A THALASSAEMIA CARRIER, THE FOLLOWING CAN OCCUR:

When EITHER the mother or father is a







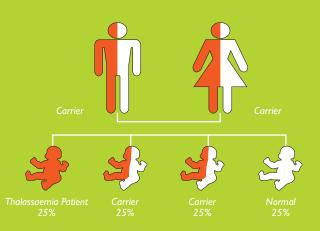


If you are a Thalassaemia Carrier, avoid marrying another Carrier.



IF BOTH PARENTS ARE THALASSAEMIA CARRIERS, THERE IS A HIGH CHANCE THAT IN EVERY PREGNANCY, THE CHILD CAN GET THE THALASAEMIA DISEASE -A LIFE THREATENING CONDITION.

When BOTH parents are thalassaemia carriers



Signs & Symptoms of a Thalassaemic **Patient** (Thalassaemia major)

- Children with Thalassaemia major appear normal at birth. However they start to suffer from serious anaemia between 3 to 18 months of age.
- Increasingly visible paleness, accompanied by constant fatique and lethargy
- Difficulty in breathing due to severe anaemia
- Jaundice
- Distended abdomen due to swelling of the liver and spleen
- Stunted growth
- As the child grows changes occur in the facial bone structures