

## **Your antenatal blood test result: you are a sickle cell carrier**

This information is for you if you are pregnant and have had a blood test which shows that you are a sickle cell carrier (often referred to as 'having a trait'). This leaflet gives you information about being a carrier, and what this means for you and your family.

### **How your test result is written**

Your result is written as 'results consistent with sickle cell carrier (Hb AS)'. The letter A represents your usual haemoglobin A gene. The letter S represents your sickle cell haemoglobin gene.

### **What your blood test result means**

Haemoglobin is the substance in red blood cells that carries oxygen around your body. Being a sickle cell carrier means you have inherited one usual haemoglobin gene called haemoglobin A from one biological parent and one sickle cell gene from your other biological parent. Because you have inherited usual haemoglobin A from one parent, you are healthy. Being a carrier does not make you ill.

Knowing that you are a sickle cell carrier means that you could pass the sickle cell gene on to your baby. If your baby's biological father is also a carrier, this means together you could have a baby with an inherited haemoglobin condition. If one biological parent is a sickle cell carrier and the other biological parent has an unusual haemoglobin gene, such as haemoglobin S (sickle cell) this could cause sickle cell disorder.

Please let your healthcare professional know if you:

- became pregnant following fertility treatment with donor sperm or a donor egg;
- have had a bone marrow or stem cell transplant; or
- are pregnant as a surrogate.

### **Inherited haemoglobin conditions**

Haemoglobin conditions are blood disorders inherited from both biological parents. There are many haemoglobin conditions. The type of condition your baby could inherit will depend on what type of haemoglobin genes both biological parents have. Some conditions are more serious than others. The most serious conditions are called sickle cell disorders or transfusion dependent thalassaemia (thalassaemia major). There are different types of sickle cell disorders, but the most serious type is sickle cell anaemia (Hb SS), which happens when both biological parents are sickle cell carriers.

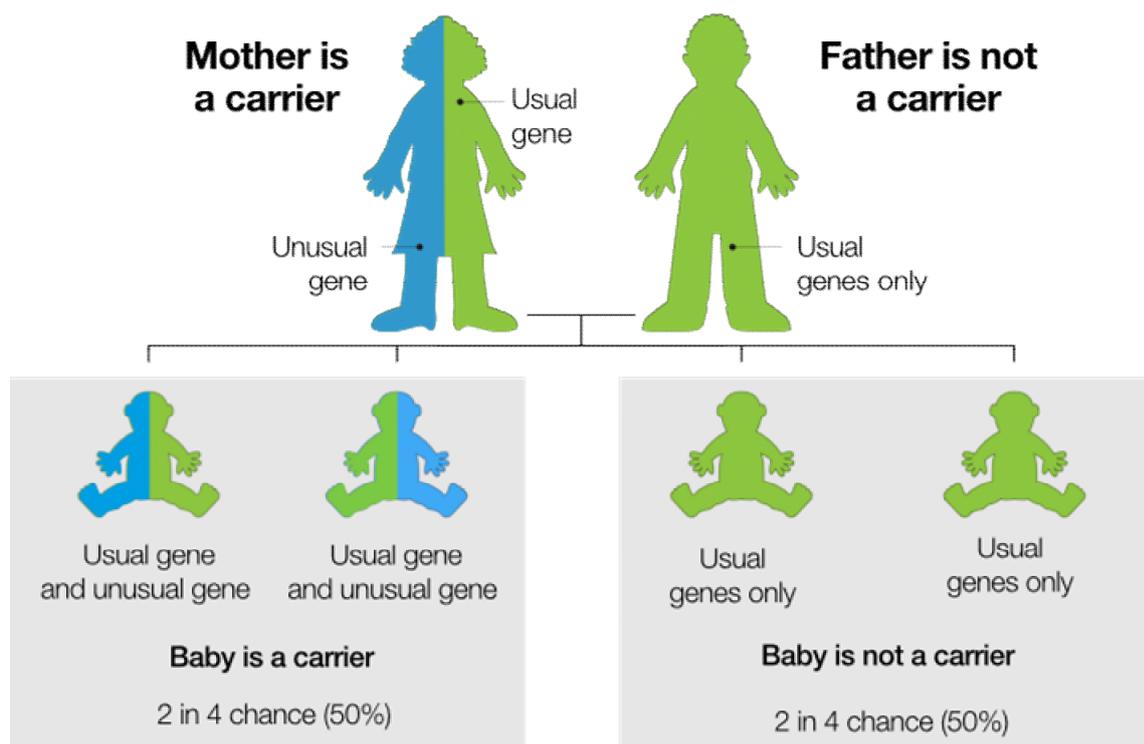
Your healthcare professional can discuss this with you, so that you understand what condition your baby could inherit and how this could affect the health of your baby.

### **How would my baby inherit a sickle cell disorder?**

#### **One biological parent is a carrier and the other biological parent is not**

If your baby's biological father has two usual haemoglobin genes (AA), there is a 2 in 4 (50%) chance your baby will be a sickle cell carrier (like you) and a 2 in 4 (50%) chance they will have two usual haemoglobin genes (AA). See diagram below.

These chances are the same in every pregnancy for this couple.

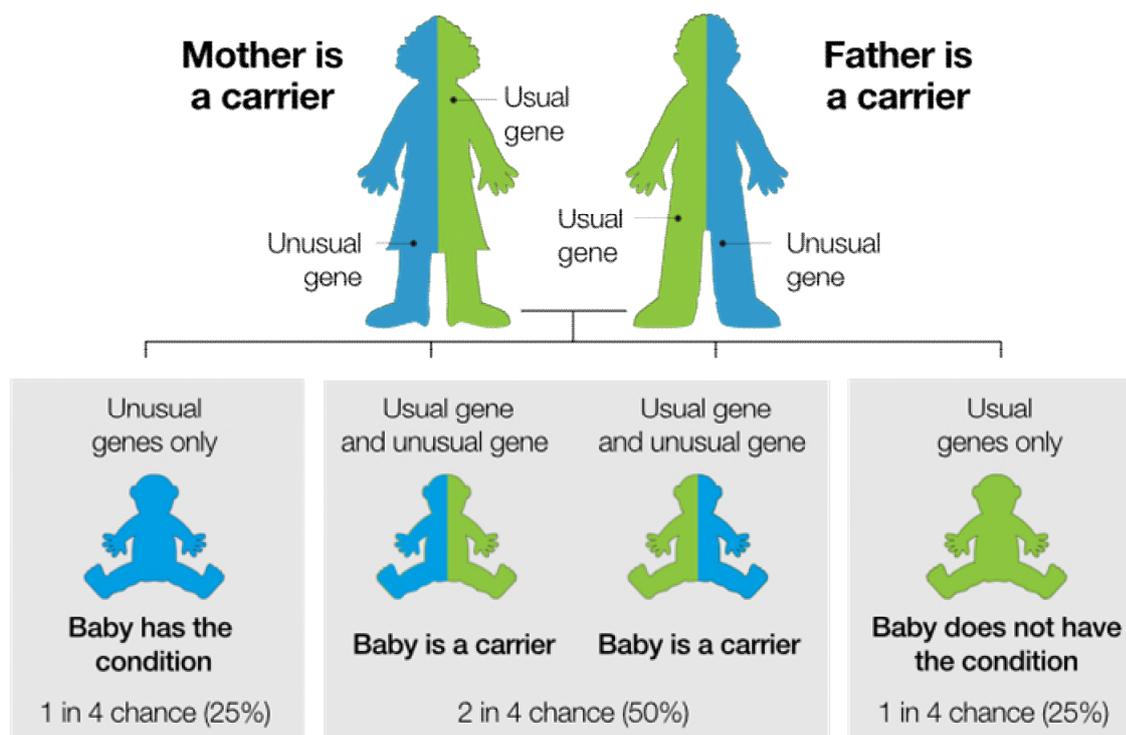


### What if both biological parents are carriers?

If your baby's biological father is a carrier, there is a 1 in 4 (25%) chance your baby will inherit a haemoglobin condition. The seriousness of the condition will depend on what haemoglobin genes are inherited.

There is also a 2 in 4 (50%) chance your baby will be a carrier (like you), and a 1 in 4 (25%) chance your baby will have usual haemoglobin genes (AA). See diagram below.

These chances are the same in every pregnancy for this couple.



### What else is it important to know as a sickle cell carrier?

There are very rare occasions when you should take extra care because of your carrier status. If you need a general anaesthetic for an operation or for dental treatment, make sure healthcare professionals know that you are a sickle cell carrier.

If you take part in any form of vigorous physical activity, you should:

- make sure you drink plenty of fluids;
- avoid extreme exhaustion; and
- make sure you are never short of oxygen.

### Should my baby's biological father be offered testing?

If your baby's biological father has an unusual haemoglobin gene it is important to identify the type of gene and the chance of your baby inheriting a haemoglobin condition. For this reason, we will also invite the baby's biological father for screening. He will only know if he carries a gene for unusual haemoglobin if he has a blood test to check his haemoglobin type.

[phw.nhs.wales/services-and-teams/screening/antenatal-screening-wales/information-resources/leaflets/sickle-cell-and-thalassaemia/sickle-cell-and-thalassaemia/information-for-fathers-invited-for-a-screening-test-for-sickle-cell-disorder-and-thal/](http://phw.nhs.wales/services-and-teams/screening/antenatal-screening-wales/information-resources/leaflets/sickle-cell-and-thalassaemia/sickle-cell-and-thalassaemia/information-for-fathers-invited-for-a-screening-test-for-sickle-cell-disorder-and-thal/)

If the test shows your baby's biological father is a carrier of an unusual haemoglobin gene you will be offered specialist counselling and prenatal diagnosis.

### What happens when my baby is born?

All babies are offered a newborn bloodspot screening test around four days after their birth. This test will screen for sickle cell disorder.

### **Should I tell my family the result?**

Other family members, such as brothers and sisters, aunts, uncles and cousins may want to consider getting tested as they could be carriers too. Sharing this information with them may be helpful.

### **Next steps and choices**

You can also:

- let your GP or healthcare professional know you are a sickle cell carrier; and
- contact All Wales Medical Genomics Service (AWMGS) where you can get free information and advice to help you understand the implications of being a sickle cell carrier.

### **More information**

You can get more information from the following.

[111.wales.nhs.uk/Encyclopaedia/s/article/sicklecelldisorder](http://111.wales.nhs.uk/Encyclopaedia/s/article/sicklecelldisorder)



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