



Cystic Fibrosis Screen Positive Inconclusive Diagnosis

What is CFSPID?

CFSPID stands for Cystic Fibrosis Screen Positive Inconclusive Diagnosis. This means your baby has a positive newborn screening result but does not meet all the criteria for a cystic fibrosis (CF) diagnosis.

Children with CFSPID are expected to lead long healthy lives, however, it is not known if these children are at risk of developing a CF-like disease later in life.

How do you diagnose CFSPID?

In Australia, most babies have a CF newborn screening test in the first few days of life. CF newborn screening involves measuring a marker (Immuno Reactive Trypsinogen: IRT) in a blood sample taken from the baby's heel.

Those babies with a high IRT will go on to have a genetic test that looks for the genes responsible for causing CF. The babies will then undergo a sweat test.

There are three possible outcomes from the sweat test:

- A positive result (chloride >60mmol/L)
- A negative result (chloride <30mmol/L)
- A borderline result (30-60mmol/L)



Your child may be diagnosed with CFSPID for the following reasons:

- Two CF genes are found and at least one of those genes has unknown or unclear consequences. The sweat test is normal or borderline.
- One or no CF genes are found, and the sweat test results are borderline on more than one occasion.

How frequent is CFSPID?

The true frequency of CFSPID is still unknown. It is estimated that for every three babies diagnosed with CF through newborn screening, there is one CFSPID diagnosis.

How can CFSPID affect the body?

With widely available genetic testing, more is being discovered about the genes that are found with CFSPID. Although most people with CFSPID will remain healthy, some may experience milder symptoms of CF disease. It is possible that some children may develop more significant evidence of CF disease later in life. Currently, there is not enough information available to predict who will remain healthy and who will develop CF disease. If your child was to develop symptoms of CF disease, the following areas of the body can be affected: lungs, digestive, and reproductive systems.

All children are at risk of developing chest infections, but for children with CFSPID, these may take longer to clear and require treatment with antibiotics. Additionally, children with CFSPID may develop inflammation of the pancreas (a digestive gland) or, if they are male, may be diagnosed with infertility as an adult. These are the main body areas that can be affected in CFSPID, but your CF team can provide you with more information.

The precise onset of symptoms is unknown. We assume that this, if at all, may occur later in life, however, in some individuals, this may be earlier. The CF team will use several tools to monitor the health of your child. This is important, as early treatment of potential symptoms will help to keep your child healthy.

How are children with CFSPID monitored?

It is important to regularly monitor children with CFSPID in a CF clinic, this may be 6-12 monthly depending on your child's age. One in 10 babies with CFSPID may meet criteria for CF as they get older. It is essential for the CF team to monitor your child so early treatment and interventions can begin if necessary. Your CF clinic will arrange regular appointments and assessments for your child. Your CF clinic can provide you with more information.

Please report to your CF clinic if your child:

- Stops gaining weight
- Has loose stools, stools with abundant mucus or, very foul-smelling stools
- Develops acute severe stomach pain and vomiting
- Has a wet or long-lasting cough
- Has frequent lung or sinus infections.

How do you feel?

Many parents say they feel shocked, scared, and confused when they are told their child has CFSPID. These are normal feelings. If you are feeling overwhelmed or have questions, you are encouraged to talk with your CF team or your GP.

Your CF clinic

There is a large multi-disciplinary team at Perth Children's Hospital, all of whom are involved in providing you and your child with the best possible care.

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Acknowledgment to Cystic Fibrosis Clinics at British Columbia Children's Hospital Vancouver and Hospital for Sick Children Toronto and CFQ.