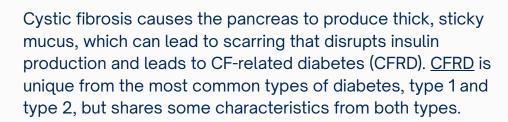
# Cystic Fibrosis-Related Diabetes (CFRD)



# What is cystic-fibrosis related diabetes (CFRD)?

<u>Diabetes</u> is caused by high blood sugar, or glucose, a source of energy that comes from carbohydrates and needs a hormone called insulin to become fuel for energy. Insulin is produced in the <u>pancreas</u>. When the body doesn't produce enough insulin, glucose builds up.







## How is CFRD diagnosed?

About <u>25%</u> of adults living with CF have CFRD. CFF recommends a yearly oral glucose test for all people with CF ages 10 and over. An oral glucose tolerance test is when a patient drinks a sugary liquid and their blood sugar is measured over specific periods of time to measure insulin resistance.

Symptoms of CFRD include:

- Feeling very thirsty
- · Urinating more than usual
- Feeling overly tired
- Losing weight
- Declining lung function
- Increasing exacerbations and infections

### How is CFRD managed?

The goal of managing CFRD is to maintain a good nutritional status and an optimized blood glucose level. CFRD is managed by **taking insulin, monitoring blood sugar, eating a balanced diet,** and **exercising.** An energy-dense, high-fat, and balanced (40% fat, 40% carbohydrates, and 20% protein) diet is recommended to support CFRD management.

People may use insulin pumps or take injections to manage blood sugar. Some people monitor their blood sugar by wearing a device called a continuous glucose monitor, which checks blood sugar levels every few minutes. Others may test their blood sugar by pricking their finger using something called a "finger stick."





#### Other Factors Impacting CFRD

Pregnant women living with CF may experience unique challenges related to CFRD and pregnancy, such as a high risk for hypoglycemia, altered digestion, fluctuating insulin resistance levels, and nutrition needs.

People who have had solid organ transplants may develop diabetes, which presents in <u>35-50%</u> of adults with CF who have had lung transplants.

#### **Mental Health Effects of CFRD**

Living with CF and CFRD can be isolating and may affect social functioning, body image, emotional responses, chest symptoms and interpersonal relationships. CFRD may also increase depression risk.

Person-centered care, routines, improvements in CF symptoms and lung function, and living connected to values may help to promote CFRD management.

