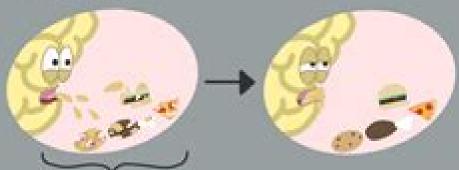
# Cystic Fibrosis Alveoli Bacteria Mucus CFTR protein

CF | K ▲F508

In cystic fibrosis chloride channels become dysfunctional which leads to less chloride transport in certain organs. Less chloride, leads to less fluid transportation and therefore thick mucus production in the lungs. Thick mucus is hard to get rid of and easier for bacteria to grow in





Duodenum

Thicker mucus in the pancreas causes blockage of the pancreatic duct. The pancreatic enzymes can therefore not enter the duodenum to help with digestion. This leads to indigestion and malnutrition

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# **Cystic Fibrosis And Related Human And Animal Diseases**

**Lei Huang** 

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