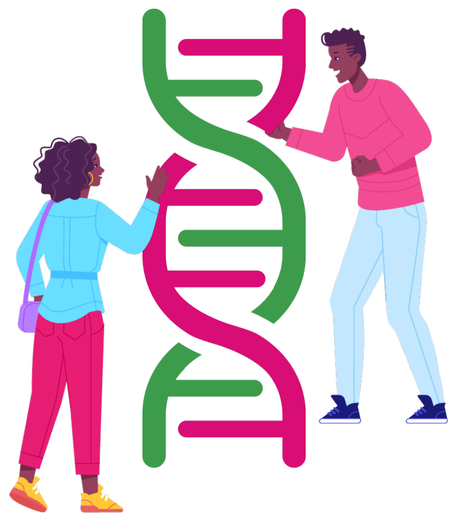
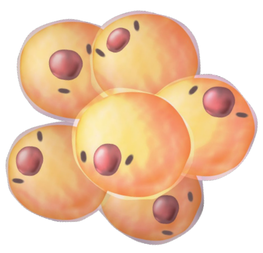
What Is Familial Chylomicronemia Syndrome (FCS)?

FCS OCCURS IN ABOUT ONE IN 1 MILLION PEOPLE



A GENETIC DISORDER

CAUSES CERTAIN FATS TO NOT BREAK DOWN CORRECTLY



SYMPTOMS OF FCS

- Lipemia retinalis
- Neurological symptoms
- Pancreatitis
- Hepatosplenomegaly
- Xanthomas
- Vomiting or diarrhea
- Frequent abdominal & back pain
- Numbness in feet or legs
- Forgetfulness
- Fatigue
- Triglyceride levels exceeding 1,000 mg/dL

2018 June 11th — **2020** — **2021** — **2022** Dec 23rd

FCS Foundation founded by Melissa Termine-Goetz & Lindsey Sutton

Joined Haystack Project to advocate in the US Congress

Continued to be advocators for rare disease provisions

HEART Act* (Helping Experts Accelerate Rare Treatments) PASSED!

*The HEART Act strengthens the U.S. Food and Drug Administration review process for drugs that treat rare and ultra-rare diseases by involving scientists and doctors with expertise along with patients and families .

FCS FOUNDATION

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