



What are clinical research studies?

Clinical research studies help determine if an investigational medicine is safe and effective. Before a doctor prescribes a new medicine, it must go through several phases of clinical research. The rules and ethics that doctors must follow to practice medicine also apply to clinical research studies.

Participation in clinical research studies is a choice. Volunteers may stop participating at any time.



CONQUEST

Scleroderma Research Foundation
Platform Clinical Trial

**Thank you for considering
this quest.**

For more information, contact:



**Join the quest to
advance systemic
sclerosis research**

Consider enrolling in the CONQUEST study for interstitial lung disease associated with systemic sclerosis.



What is the CONQUEST study?

Over time, the CONQUEST study will research several investigational medicines in volunteers with systemic sclerosis.

- Only research studies can use investigational medicines because health and regulatory authorities have not approved them for public use

Currently, the CONQUEST study is evaluating 2 investigational medicines for interstitial lung disease associated with systemic sclerosis (SSc-ILD) using 1 study plan, called the master study protocol.

Each investigational medicine has its own “subprotocol” with details that only apply to the study group.

- Subprotocols are extensions of the master study protocol

By sharing study resources, these investigational medicines may become available to patients faster.



Will I receive an investigational medicine?

The CONQUEST study is researching 2 investigational medicines, each a potential SSc-ILD therapy.

If you are eligible and decide to join, you will be assigned at random to 1 of the available investigational medicines or a placebo. Placebos look like the investigational medicine, but they do not have therapeutic effects. Study staff will give you more information about the investigational medicines and dosing instructions.

Who can join the study?

Candidates with the following criteria may be able to join the study:

- Adults ages 18 or older
- SSc-ILD diagnosis
- SSc symptoms that began within the past 5 years
 - This study permits earlier onset of Raynaud’s disease

Other criteria apply.

What can study participants expect?

CONQUEST study participation is about 1 year. You will travel to a study center periodically to take study tests.

The following timeline is an estimate. Lengths may vary for some study groups.

- Screening: 1 month (4 weeks)
- Receive investigational medicine or placebo: about 12 months (52 weeks)
- Attend all follow-up visits: frequency of visits depends on the investigational medicine

