

# How can CoRDS benefit you?

## THE REGISTRY OPENS THE DOORS TO CLINICAL STUDIES.

Researchers will contact CoRDS if a clinical study may benefit you.\*



CoRDS provides you with this information.



You decide if you want to contact the researcher.



You find a community living with similar experiences, both for you and your family.



Your information can help physicians better diagnose patients and researchers develop treatments for your disease.

Researchers need funding to continue their research. CoRDS data helps them develop their studies and seek funding to continue their research, which could lead to a better treatment or cure for your disease.



*\*CoRDS cannot guarantee participants will be contacted by a researcher for clinical trials and/or research studies.*

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# Join CoRDS

You hold a piece of the puzzle.



You can help researchers see the whole picture. Play a vital role in finding cures and improving treatments by joining the Coordination of Rare Diseases at Sanford (CoRDS).

### Contact us:

Phone: (877) 658-9192

Email: [cords@sanfordhealth.org](mailto:cords@sanfordhealth.org)

Online: [sanfordresearch.org/cords](http://sanfordresearch.org/cords)

### Follow us



Sanford CoRDS



@SanfordCoRDS

CoRDS does not profit financially from sharing your data or personal/contact information, but solely collects the data for the sake of accelerating research into rare diseases.

**SANFORD**  
RESEARCH

**SANFORD**  
RESEARCH

# What is CoRDS?

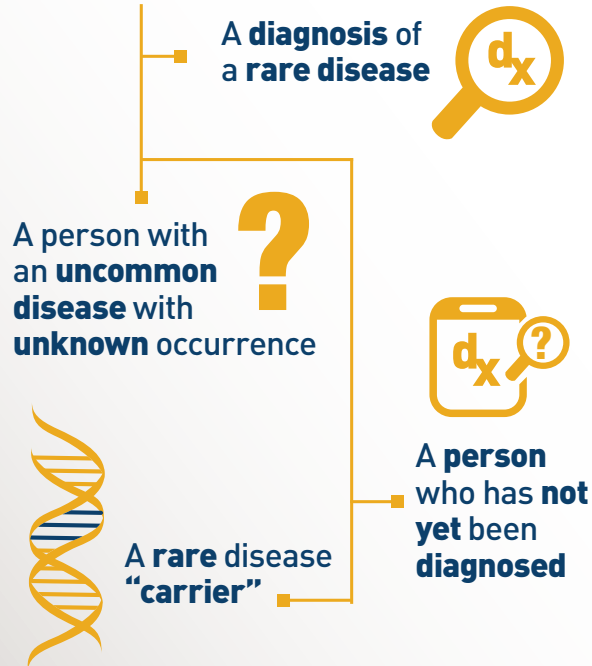
Complete details on CoRDS can be found at [sanfordresearch.org/CoRDS](http://sanfordresearch.org/CoRDS) under the Patients and Families link.

CoRDS is a centralized international patient registry for all rare diseases. Our goal is to connect as many patients and researchers as possible to advance treatments and cures for rare diseases.

The registry is free for participants to enroll. During enrollment, participants provide information on their condition for researchers to access – also free of charge. Because our patients' privacy is our first priority, the database is secure and all information is protected, so the information researchers receive does not identify the participant.

## How to join CoRDS

 **ANY PATIENT WITH THE FOLLOWING CAN ENROLL:**

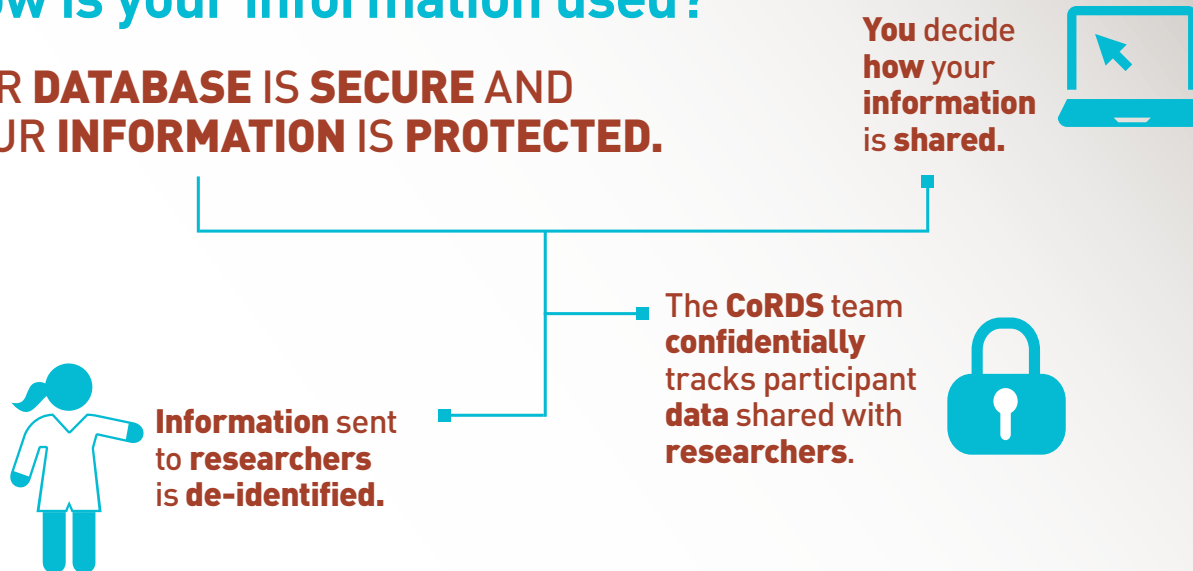


YOU CAN REMOVE YOUR INFORMATION FROM THE REGISTRY AT ANY TIME.

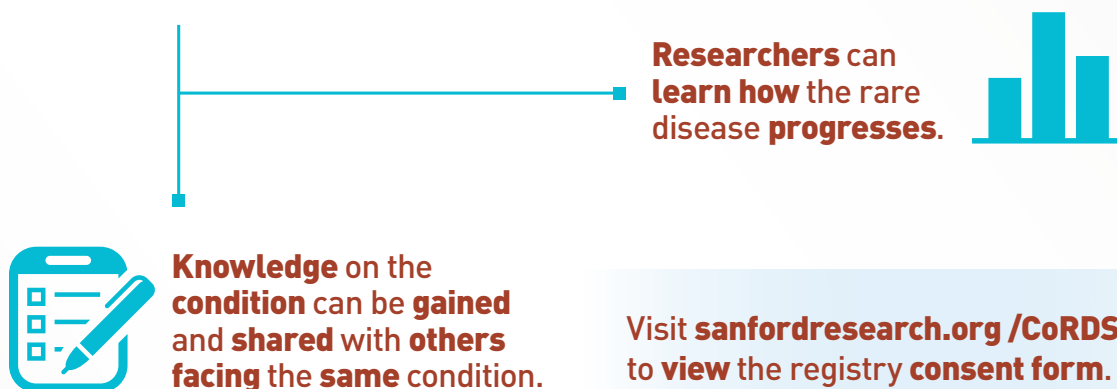
 **Enroll** now at [sanfordresearch.org/CoRDS](http://sanfordresearch.org/CoRDS) by selecting **Enroll Now**.

## How is your information used?

**OUR DATABASE IS SECURE AND YOUR INFORMATION IS PROTECTED.**



**YOUR DATA CAN HELP FUTURE FAMILIES WITH THE SAME DIAGNOSIS.**



Visit [sanfordresearch.org/CoRDS](http://sanfordresearch.org/CoRDS) to view the registry consent form.

