



EOSINOPHILIC GASTROINTESTINAL DISEASE (EGID)

Working Together to Make a Better Tomorrow



The CEGIR Mission

To improve the lives of individuals with eosinophilic gastrointestinal disorders through innovative research, clinical expertise and education via collaborations between scientists, health care providers, patients, and professional organizations.



Initiative of the National Center for Advancing Translational Sciences (NCATS)

Have you been searching for the latest research studies for Eosinophilic Gastrointestinal Disease (EGID)?

Join the Rare Diseases Clinical Research Network (RDCRN) Contact Registry today!

Have you or your child been diagnosed with EGID?

- ◆ Eosinophilic Esophagitis (EoE)
- ◆ Eosinophilic Gastritis (EG)
- ◆ Eosinophilic Colitis (EC)

If so, you are invited to sign up for the Rare Diseases Clinical Research Network (RDCRN) Contact Registry.

What is the RDCRN Contact Registry?

The RDCRN Contact Registry is a way that patients with rare disorders, like EGID, can receive information about new research studies, eligibility for participation in clinical studies/trials, and updates on results of such studies.

Why Join the RDCRN Contact Registry?

Patients who participate in research make it possible for researchers to find new treatments, create new studies, and work for the improvement of all EGID patients. It can also help connect you with Patient Advocacy Groups (PAGs).

To find out more about the RDCRN Contact Registry, or to join, please go to the following website (or scan the QR code with your smartphone):

rdcrn.org/CEGIR



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