The HI Global Registry
Congenital Hyperinsulinism Patient-Powered Research for a Brighter Future

Congenital Hyperinsulinism International has developed the HI Global Registry to advance treatments and find cures. The HI Global Registry is a resource to better understand how the whole community of HI patients is affected by the condition.

The HI Global Registry is a patient-driven study made up of surveys about the patient experience and health outcomes. Patients, their parents, or legally authorized representatives can participate by entering information from anywhere in the world. The data is made anonymous and stored securely in an online database. Upon receiving consent from participants, registry data will be shared with other participants and those conducting research and clinical trials, as approved by the HI Global Registry Steering Committee that includes, doctors, scientists, and patient advocates.

Join us to help advance HI research and find a cure!
www.congenitalhi.org/higlobalregistry
Your Participation is Critical!

Join the HI community to advance research and cures!

Congenital Hyperinsulinism International (CHI)
P.O. Box 135
Glen Ridge, NJ 07028

(973)-842-7559
info@higlobalregistry.org

To participate visit:
www.congenitalhi.org/higlobalregistry