



**Congenital Hyperinsulinism International
Support and Awareness Program**

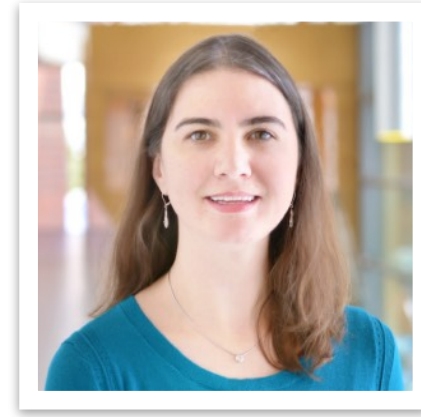
**Jennifer Schmitt, COO
Congenital Hyperinsulinism International
Brisbane Marriott Hotel, Brisbane, Australia
August 5, 2023**



Julie Raskin
Founder and CEO



Jennifer Schmitt
Chief Operating Officer



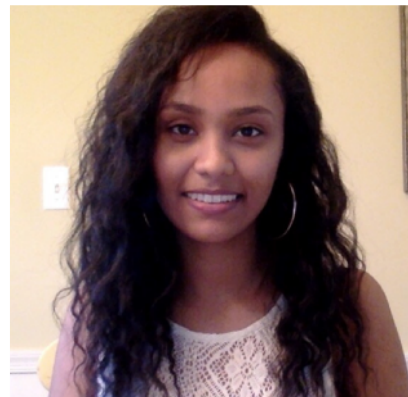
Tai Pasquini, PhD
Chief Research Officer



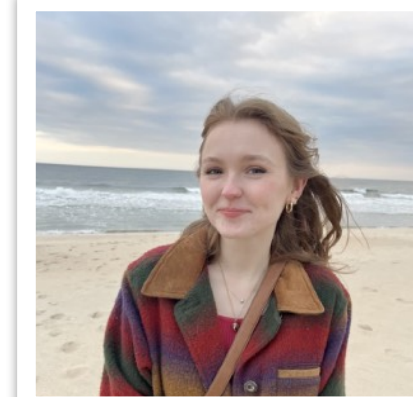
Diane Esty
Development Director



Lauren Lopez, PhD
Registry Director



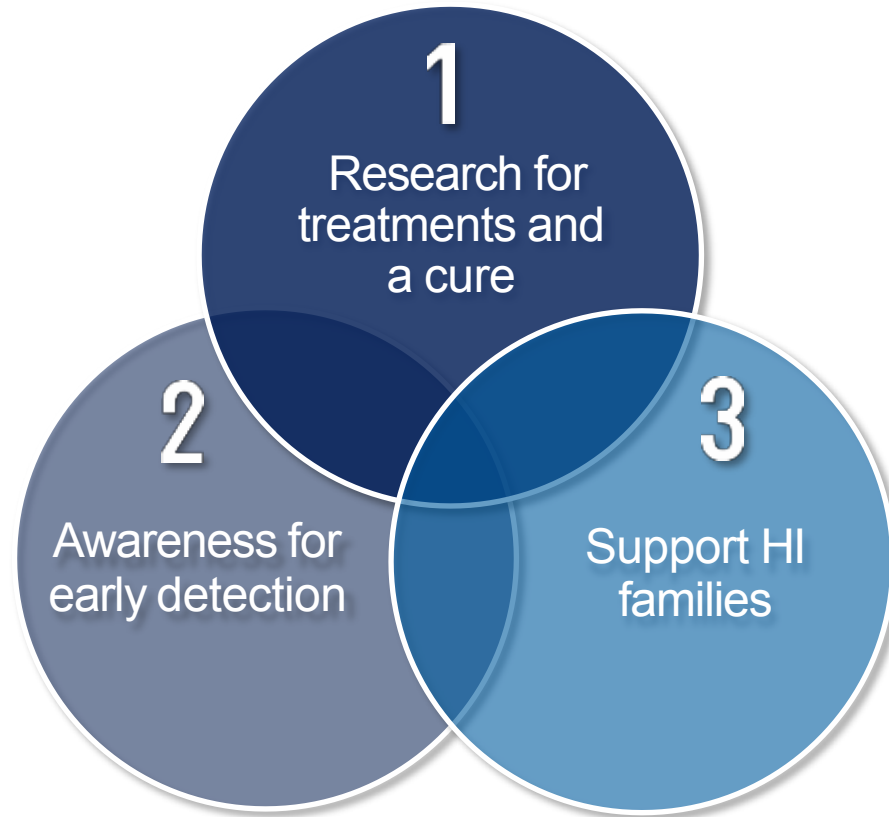
Mahlet Mesfin
Research and Policy
Associate



Lily Barnett
Communications
Associate



Lora Van Arsdell, PsyD
Research and Support
Specialist



Since its founding in 2005 CHI:

- Helps patients around the world receive life-saving medications, treatment, and medical supplies.
- Serves as the global source of HI patient and family support available to thousands of people in the HI community around the world with an online support group which is active 24 hours a day.
- Ensures the patient voice is included in drug development with the hope that new treatments will be developed to help thousands of patients.



Since its founding in 2005 CHI:

- Partners with Exeter Clinical Laboratory to provide genetic testing to over **786** patients in **60** countries who are suspected of having HI and who do not have the ability to pay for the testing.
- Designated six CHI Centers of Excellence in the US, UK, and Germany.





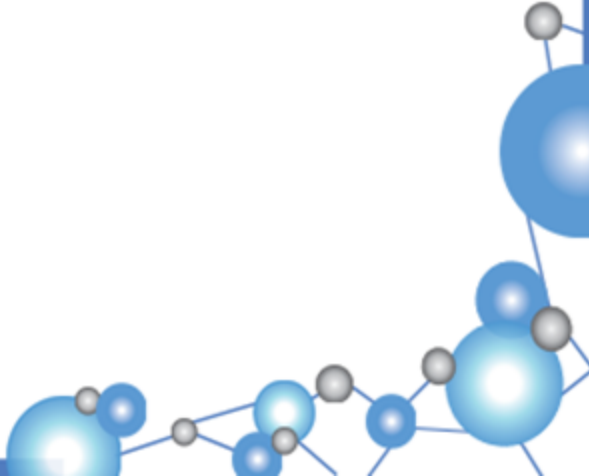
Since its founding in 2005 CHI:

- Has organized 25 HI conferences and meetings around the world bringing together leading global HI specialists to share best practices and research.
- Widely shares and advocates for the adoption of the PES Hypoglycemia Guidelines.



Since its founding in 2005 CHI:


- Created and distributed the global HI and hypoglycemia awareness poster campaigns to save lives and prevent brain damage. The posters are available in 24 languages with a 25th on the way.



Since its founding in 2005 CHI:

- Has created the *Future Without Lows*, *Be My Sugar*, *Stop the Lows*, *HI Never Sleeps* and many other awareness campaigns to save lives and prevent brain damage. Glucose as a Vital Sign coming soon!

Recurrent hypoglycemia in newborns, infants and children is **NOT NORMAL.**



LEARN THE SIGNS & SYMPTOMS OF HYPOGLYCEMIA TO PREVENT BRAIN DAMAGE AND DEATH

CHI Congenital Hyperinsulinism International congenitalhi.org

CHIbras are strong because...



"He went from screaming over every sugar check to telling me, 'It doesn't hurt, mommy,' at just three years old!"

- Kimberly Pfaff

congenitalhi.org

Alexander loves to play soccer, but his mom worries about blood sugar drops when he does.



Learn about HI clinical trials #FutureWithoutLows™

congenitalhi.org

SUGARPARENT QUOTES



"She's our little rock star and fighter - she's been through so much already and always finds a way to keep her smile."

- Janine Grizzle Cook

congenitalhi.org

CELEBRATIONS OF STRENGTH

#SWEETESTCAUSE



Dante is such a special boy who just knows that we must check his blood sugar multiple times a day. He is aware that certain foods are good for his blood sugar and others make his sugar drop. He tells everyone goodbye followed by "I love you with my whole heart." He loves to play with his brother and sister. He is a great student & very analytical. It would not surprise anyone if Dante grows up to be some type of engineer.

SUGAR SOIREE 2019

#STOPTHELOWS!



"With Research, Possibilities Are Limitless."



I support the #SweetestCause because no one should have to fight HI alone!

- Klersten Smith

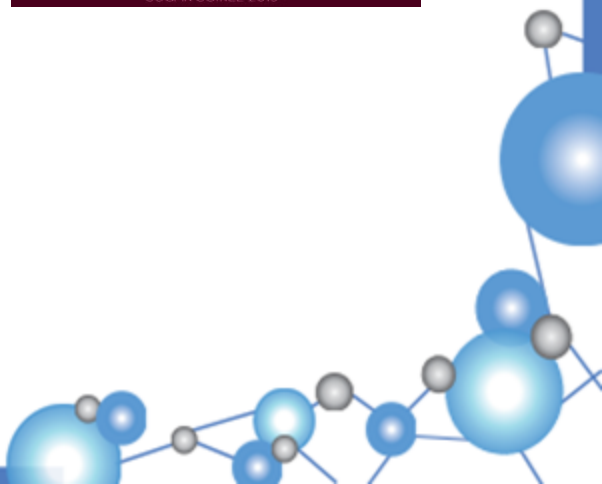
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I DARE TO DREAM...

that Daphne and the rest of the HI kids in the Philippines will have access to medicine and treatment. We dream that we will no longer have to get medicine and treatment from other countries and not wait for months. Just because HI is a rare disease, our medicines and treatment should not be rare. Our goal is for our country to understand this rare disease and its treatment.

- Faith, Philippines

Rare Disease Day is February 28 | #RareandDaretoDream



THANK YOU!

