

CONGENITAL DIAPHRAGMATIC HERNIA PATIENT SUPPORT

Sponsorship Proposal



CDH International is the
world's oldest, largest, and
leading Congenital
Diaphragmatic Hernia charity



www.cdhi.org



sponsor@cdhi.org



+1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



CDH International supports 1000's of patient families in 84 countries, however 52,000 babies are born with Congenital Diaphragmatic Hernia every year. We have work to do.

CDH International was founded in 1995 with just one goal; that no family would have to face Congenital Diaphragmatic Hernia alone.

It was an admiral goal, if not naive. There is much more work to be done than just holding the hand of a diagnosed family. These families need accurate, unbiased information in easy-to-understand terms in their native languages.

They also need emotional support, mental support, trauma therapy, financial support and family support. They need to gain skills to help their children or manage their grief. Or both.

At CDH International, we take a whole family approach to support these patients. They must not fight CDH alone - but they must fight it empowered with knowledge and support.

Every 10 minutes a baby is born with Congenital Diaphragmatic Hernia. At least 3 will die each hour.

CDH International is dedicated to finding the cause, prevention, and best treatments for Congenital Diaphragmatic Hernia while also supporting affected patients and families.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



About Us

Founded in 1995 as CHERUBS, CDH International is a global non-profit working to help over 6500 patient families in 84 countries through support services, raising awareness and funding, and conducting CDH research.

CDHi is a registered non-profit in the United States, Canada, United Kingdom, Switzerland, Hong Kong, Singapore, and the Netherlands.

Headquartered in Wake Forest, North Carolina, CDH International was founded by 2 CDH mothers and is run by patient parents, grandparents, and adult CDH survivors.

Our Mission:

CDH International was created to help families of babies born with Congenital Diaphragmatic Hernia by providing support services, promoting research and raising awareness.



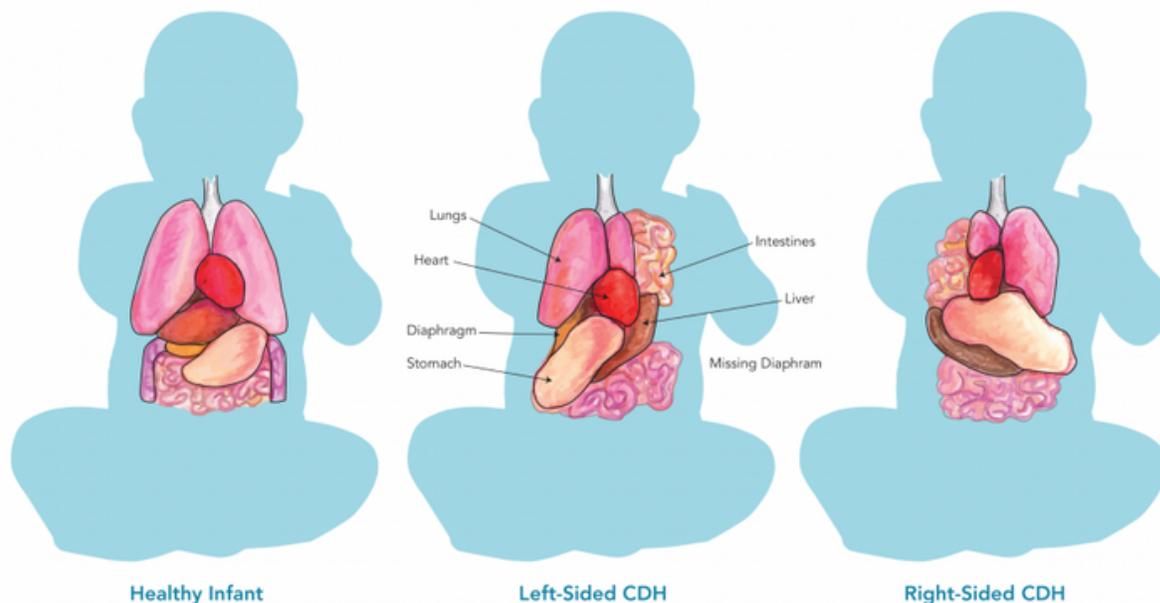
CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Congenital Diaphragmatic Hernia (CDH) is a birth defect that occurs when a baby's diaphragm (a thin sheet of muscle that separates the abdomen from the chest) fails to fully form, allowing abdominal organs to enter the chest cavity and prevent lung growth.

CDH occurs in approximately 1 in 2500 births, with over 1600 babies diagnosed each year in the United States alone. There is currently no known cause, but studies have suggested that it is likely due to a number of genetic and environmental factors.

Treatments:

Surgery to repair the diaphragm and place organs into their proper places is the most common treatment of Congenital Diaphragmatic Hernia. Timing and types of repairs vary on the size of the hole, organs involved, and other health issues. Babies diagnosed in utero can sometimes be treated with fetal intervention to help the lungs to grow.

Prognosis:

The overall survival rate is 50% in Higher-Income Countries and less than 1% in Lower-to-Middle-Income Countries. CDH is a spectrum disease, with each survivor's journey and outcome very different.

Costs:

Babies born with Congenital Diaphragmatic Hernia are the sickest and most expensive NICU patients. They spend weeks or months in intensive care, often requiring heart and lung bypass, fetal treatments, and other costly interventions. Medical bills can be in the millions of dollars per patient.

Families of CDH patients also lose time at work, and sometimes must relocate out of state, losing jobs, leaving other children behind, doubling their cost of living and leaving behind their support system of family and friends.

CONGENITAL DIAPHRAGMATIC HERNIA



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity



www.cdhi.org



sponsor@cdhi.org



+1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA

Helping Congenital Diaphragmatic Hernia Families

"We found CDH International after our son, Kieran, was born and are incredibly grateful to have a continued network of support through other families and professionals all across the globe.



We've gained a knowledge and insight through the educational resources that we haven't found anywhere else to help us along this journey with our son."

— MELISSA AND JUSTIN JOHNSTONE
Parents to CDH Survivor Kieran

"CDH International has meant the world to our family then and now!



It was like a mental wellness break to talk to families going through not only our, but their CDH journey. 7 years later, we know how lucky we are to have our beautiful and healthy daughter, Lindsey. We continue to raise awareness by sharing our story, lobbying on Capitol Hill, speak with other families in Virginia that were given the horrible diagnosis of CDH, and hold an annual fundraiser at a local restaurant. Giving back is a selfish act that does not take a lot of time, makes you feel good, and also helps others along the way. We are so grateful to CDH International for all the support they provide to all of our families."

— JEFF AND LISA BAXTER
Parents to CDH Survivor Lindsey

Having a child born with Congenital Diaphragmatic Hernia can be emotionally, mentally, physically, and financially devastating to families. CDH International is here to help.

Our Services & Projects:

- Public Education
- Educating Newly Diagnosed Families
- NICU Care Packages
- On-Call Emotional Support
- Financial Assistance Grants
- Funeral Assistance Grants
- Scholarships
- Congenital Diaphragmatic Hernia Patient Registry
- Research Grants & Collaborations
- CDH Family Conferences & Events
- Raising Global CDH Awareness Through Events, the Telethon, April 19th Light Ups, Proclamations, Legislation, Social Media and More
- CDH Radio Show
- CDH Phone App for Doctors and Patients
- Weekly CDH News

"Thanks to CDH International my daughter has met other CDH children and we have made many new friends.



CDH International has helped us find families with similar experiences for support. Most importantly CDH-ii has given me hope that my daughter will always have a place to turn as she navigates CDH as an adult."

— AMANDA FLAKHOLM
Mom to CDH Survivor Elizabeth

"After my family experienced a life changing event when my grandson was born in October with a birth defect called Congenital Diaphragmatic Hernia or CDH our lives have been altered forever.



Never having heard of it, I started researching and this led me to information that we all needed. So much was frightening and odds were not on our side. Our baby, just one of the 50% that has survived his major surgery and ongoing side effects. The greater miracles surrounded him. We only prayed for miracles. He has been a good boy for quite some time, but he is doing so well. When we were in his hospital the first month, this was the only place I had to go to for information and support. Thank you."

— BILLIE WHEELER
Grandma to CDH Survivor Tesh

"Seeing a prayer request shared in the midst of complete fear, knowing total strangers were pulling for our son was sometimes the only thing that helped carry me through our ECMO days.



I am forever grateful to CDH International for being there with us not only in the good times, but beside us through the hard times when we didn't know if we could make it through."

— SARAH TANNER
Mom to CDH Survivor Austin

"CDH International has been there for us since 2004 when my son Brandon was born with a congenital diaphragmatic hernia.



We found out about his condition two days prior to his birth. CDH International was the only resource we could find that explained to us his condition and what to expect (good or bad) moving forward. Fifteen years later we still are so grateful for all the support and information they continue to provide to so many families."

— CHERYL SANDOVAL
Mom to CDH Survivor Brandon



"While I was pregnant with my first son Hunter, I was connected with CDH International shortly after his LCDH diagnosis.

I had never heard of CDH before and felt very alone. However on this site, I was able to connect with other families going through the same situation as our family. I was able to read stories and post my own, even asking questions and for advice while Hunter was in the NICU. I received so much love and support from the other families that I cherish greatly. The charity helped our family tremendously with Hunter's diagnosis and gave us hope when we had none. When we first received his diagnosis and learned that he may not survive, you can only imagine the emotions we felt in knowing that our baby may never come home. CDH brought light to a very dark situation and we are thankful for them everday!"

— LIBBY MATTESON
Mom to CDH Survivor Hunter

"CDH International has provided our family with an extended CDH family that has shared the same experiences, ups and downs as we have.



It has also given us a world of information that we didn't even know existed!

— RYAN AND CHRISTINA BRAY
PARENTS TO CDH SURVIVOR CONNOR

"CDH International has helped me so much from the beginning.



I was so scared when I found out Rosey had CDH and literally thought she was going to die but through your help I gave me the hope I needed. Scarie scars just remind us that we survived."

— RACHEL BARR
Mom to CDH Survivor Rosey

"CDH International was and continues to be a tremendous network and resource for my family.



Our daughter's diagnosis was incredibly difficult to comprehend, but CDH International provided support and a network of families dealing with the same challenges. From the day of Harper's diagnosis through her time in the hospital and even now, years later after her passing, we are comforted knowing we will forever be connected through this charity and families that belong to CDH International."

— LYNN GRAY
Mom to CDH Angel Harper

"CDH International has helped us to know that we were never alone throughout our daughter's life and we are not alone now after her death.



Because of so many we've met through CDH International we know that our Grace is not forgotten."

— MIRANDA HALL
Mom to CDH Angel Grace

"CDH has helped me connect with parents that have gone through something similar with their baby.



It is nice being able to talk with parents that have gone through the same things my daughter and I have. We all have different stories and journeys, but have been affected by the same condition."

— MADISON CARPENTER
Mom to CDH Survivor Adalynn

"Because of our experience with CDH International, our hope and vision for CDH has expanded beyond just our personal journey.



We believe that so many more families and children can be supported as we travel, and keep in touch with each other online and in person. We couldn't be more grateful for the dedication that CDH International has brought to our families over the last 25 years, and are looking forward to the next 25 years."

— HEIDI AND HER SON SEAN FORNEY
Seen is an adult CDH survivor

"CDH International is more than just an organization. It is a family, a support system, and an unbreakable bond between people that have never met.



CDH gives hope to families who feel like they have no where to turn. My family is one of those families and we couldn't be more thankful for coming into contact with CDH."

— THE FLOYD FAMILY
Family of CDH Survivor Bryer



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

www.cdhi.org

sponsor@cdhi.org

+1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA

Meet Our Boards



Exec. Board of Directors

- Dawn Ireland, President
- Tracy Meats, CHERUBS President
- Warren Sumner, Director
- Rhonda Montague, Director
- Dr. Brooke Newman, Director
- David Holt, Director
- Bill Opal, Director



Medical Advisory Board

- N. Scott Adzick, MD, MMM, FACS, FAAP - Children's Hospital of Philadelphia
- Kristin Aigner, RN, BSN - OSF St. Francis Medical Center
- Charles Carapinha, MBBCh, MRCS, FC Paed Surg, MMED, MPH - University of the Witwatersrand Johannesburg, South Africa
- Badr Chaban, MD - Imperial College Healthcare NHS Trust
- Priscilla Chiu, MD - Sick Kids, University of Toronto
- Wendy Chung, MD, PhD - Columbia Presbyterian / DHREAMS
- Jan Deprest, MD, PhD - University Hospital Gasthuisberg | Leuven, Belgium
- Patricia Donahoe, MD - Massachusetts General Hospital
- Mahmoud El Fiky, MD - Cairo University, Harvard University | Cairo, Egypt
- Matthew T. Harting, MD - University of Texas Medical School at Houston, CDH Study Group
- Timothy Jancelewicz, MD - Le Bonheur Children's Hospital Memphis, TN
- Gabrielle Kardon, MS, PhD - University of Utah CDH Lab
- Richard Keitzer, MD, PhD, MS - University of Manitoba
- Mauro Longoni, MD - Massachusetts General Hospital
- Steadman McPeters, CPNP - Pediatric Surgery Nurse Clinician | Huntsville, Alabama
- Henry Rice, MD - Duke University | Durham, NC
- Giovanna Riccipettoni, MD - Ospedaledei Bambini Buzzi, Milan
- Dick Tibboel, MD, PhD - Erasmus University MC, Sophia Children's Hospital | Netherlands
- Jay Wilson, MD - University of Texas Medical School at Houston, CDH Study Group
- Naomi Wright, MBChB (Hons) BSc (Hons) MRCS DCH MSc - Global Initiative in Pediatric Surgery, Oxford, UK
- Edmund Yang, MD - Peace Health Sacred Heart Medical Center at River Bend
- Augusto Zani, MD - Toronto Sick Kids Hospital



Patient Advocacy Board

- Tracy Meats, Board Vice-President
- Tina Ingham, Patient Parent
- Renci Scurlock, Patient Parent
- Sean Forney, Patient
- Lauren Dietz, Patient
- Morgan Nuchols, Patient
- Danae Perkins, Patient Parent
- Taylor Steffensmeier, Patient
- Fani Dragati Stavrou, Patient Parent
- Melissa Johnstone, Patient Parent
- Robbie Fletcher, Patient Parent
- Shelly Moore, Patient Grandparent





Our CDH Support Projects

As the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity, it is our duty to support as many affected families as possible on a global level.

♥ CDH HOPE Totebags

New families are sent a totebag full of CDH information, CDH Awareness items and gifts.

♥ CDH Phone App

CDH information available at your fingertips, 24/7, available in Google Play and the Apple Store.

♥ Emotional Support

Volunteers are available for one-on-one support to ask questions or to be there to listen.

♥ Financial Assistance

CDHi offers financial assistance to families currently in the NICU or undergoing in utero procedures.

♥ Funeral Assistance

Funds available to assist families with a funeral and burial expenses.

♥ Parent Reference Guides

The handbook to help understand Congenital Diaphragmatic Hernia.

♥ CDH Conference

A Conference that brings families, researchers and medical professionals together for research, support and awareness.

♥ CDH Grieving Retreats

A relaxing, supportive, and respectful retreat for our grieving families.

♥ Local Support

CDHi Ambassadors are on-call for local information, an ear or a shoulder.

♥ Scholarship Fund

Scholarship to a recent graduate or a parent of a cherub seeking their first college degree.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



The CDH H.O.P.E. (Helping Other Parents Expecting) Totebag Project assists families expecting babies born with CDH by providing them with much needed free information and support items through a community project in which all CDH families can participate and honor their children while helping new families affected by CDH. Donated items are collected by our members, CDH families, and friends and put together into totebags by volunteers. Totebags are then mailed to families who are expecting babies with CDH or who have newborns who are still hospitalized and currently battling CDH.

Sponsorship (\$76,000)

- 1000 baby onesies (\$3)
- 2000 parent t-shirts (\$5)
- 1000 CDH baby books (\$7)
- 1000 hat and booties sets (\$2)
- 1000 baby blankets (\$4)
- 1000 Parent Reference Guides (\$3)
- Shipping Costs (estimated \$30 per box)
- 1000 info packets (\$10)
- 1000 journals (\$1)
- 1000 puzzle books (\$1)
- 1000 teddy bears (\$1)
- 1000 pacifiers (\$1)
- 1000 chapsticks (\$1)
- 1000 pairs of wings (\$2)



CDH HOPE Totebag Project

The Goal:

Creating and gifting 1000 CDH HOPE Totebags to patient families in the United States, Canada, the United Kingdom and Ireland by December 2024.

Steps:

1. Gathering items for the totebags
2. Putting the totebags together
3. Boxing the totebags
4. Shipping the boxes to newly diagnosed CDH patients

Measurements of Success:

Meeting the goal of gifting 1000 CDH HOPE Totebags to patient families by December 31, 2024.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity



www.cdhi.org

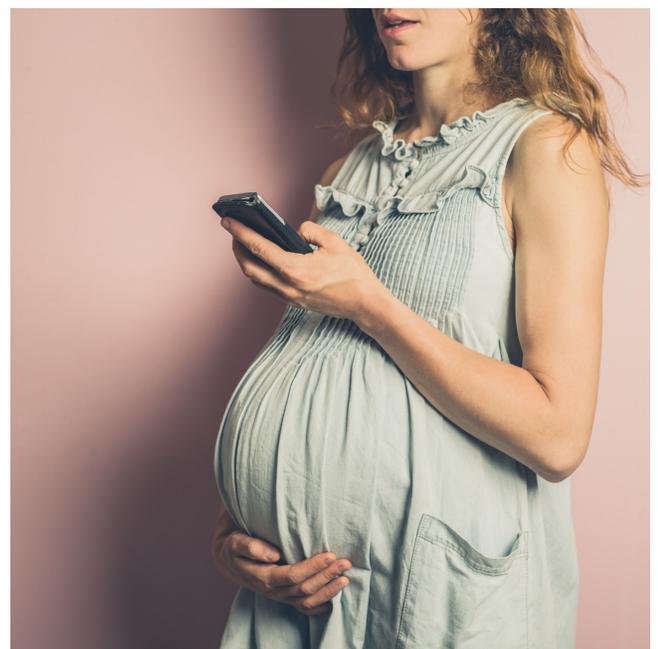
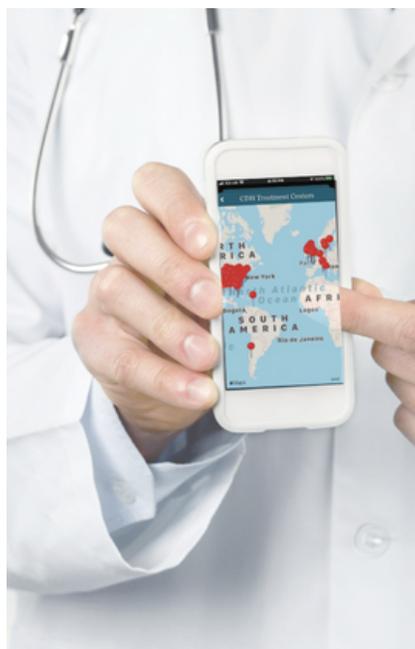
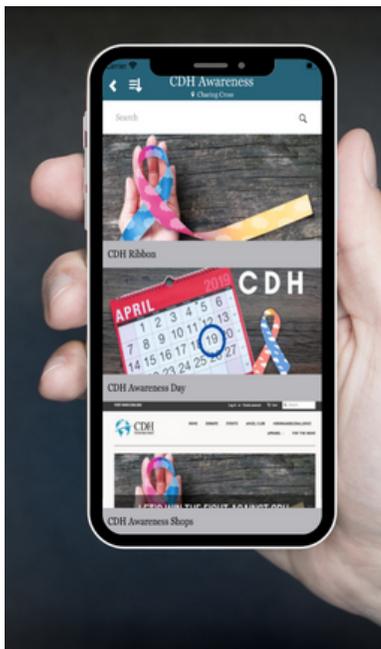


sponsor@cdhi.org



+1 919-610-0129 / +1 919-610-2972 (cell)

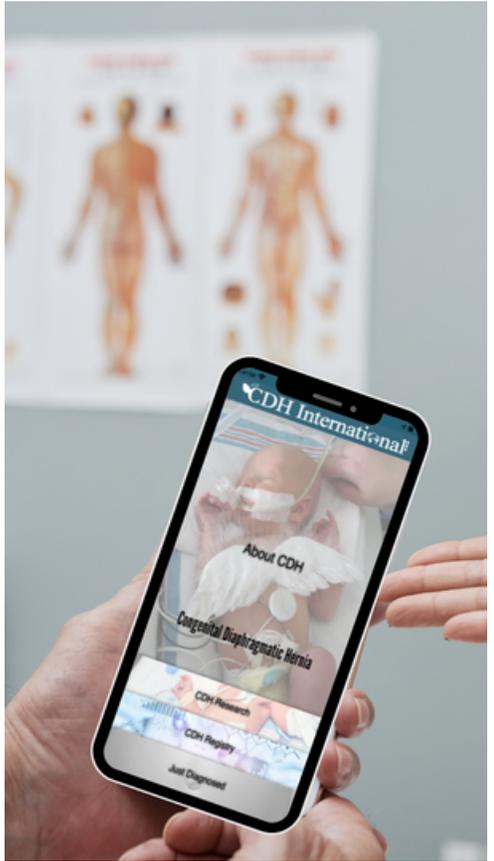
CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



The CDH App is available for free on Apple and Google Play

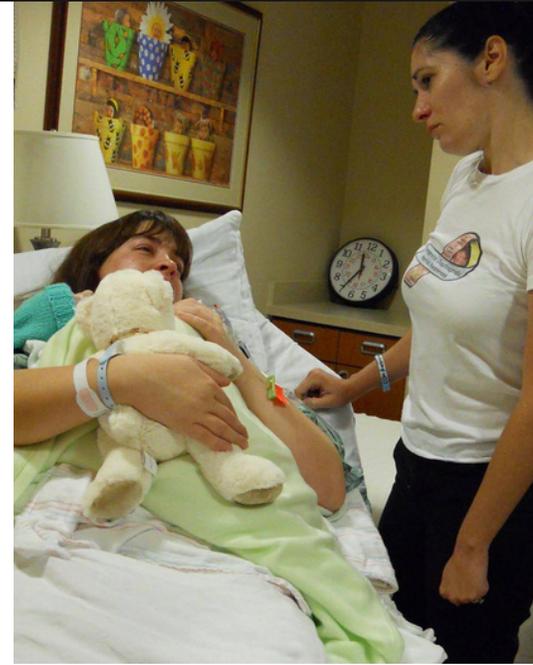


- The CDH App Contains:**
- Information for Newly Diagnosed Families
 - The CDH Parent Reference Guide
 - CDH Hospital Map
 - CDH Research Registry
 - Digital Research Library
 - Social Media Platform Just for CDH Families
 - CDH Awareness Shop
 - Event Calendar
 - Links to other CDH Apps



Phone App (\$10,000)

It's imperative that newly diagnosed patient families gain access to accurate, unbiased information on Congenital Diaphragmatic Hernia as soon as possible. \$10,000 will be spent on Google and social media ads.



Emotionally Supporting Families Affected by Congenital Diaphragmatic Hernia

Raising a child with CDH or living with Congenital Diaphragmatic Hernia is a life-long journey. Families look to CDH International for emotional support through all stages of their journey. Sadly, some babies diagnosed with CDH will not make it past their first few hours, days, or weeks of life. CDH International offers grieving parents a safe place of support where we understand what a broken heart feels like.

Some of our Resources:

- CDH Forums
- CDH Facebook Support Group for All Families
- CDH Facebook Support Group for CDH Survivors
- CDH Facebook Support Group for Grieving Families
- CDH Facebook Support Group for Grandparents
- CDHi Support Facebook Page
- CDHi Facebook Pages for all NGO's
- CDH International Website
- CDH Phone App. Social Platform
- Support Through Instagram and TikTok
- On-Call Support
- Kind Thoughts and Prayers (posted within social media)
- Virtual Chats with CDH International
- Local Get-Togethers
- Free Customized CDH Awareness Ribbons
- ACDHO Charity Resources
- Resource List for CDH Families



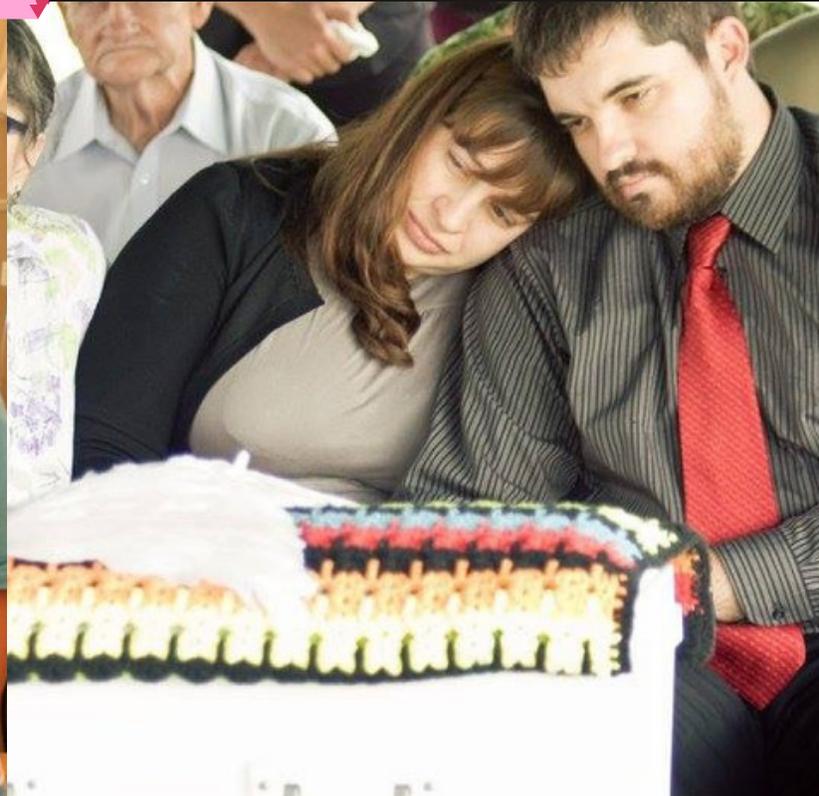
CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Financial Assistance Fund (\$50,000)

50 grants of \$1000 each to CDH families in the United States or Canada

Funeral Assistance Fund (\$30,000)

10 grants of \$3000 each to help CDH families bury / cremate their children

CDH is financially devastating to families, requiring parents to be out of work, often relocate and yet continue to pay regular bills and now medical bills. Our organization started giving small Financial Assistance Grants of \$100 in 2012. That grew to \$500 when funding was available. We would like to offer \$1000 grants per family if they meet qualifications (note from their hospital social worker proving they are in need and in the hospital) so that they can pay bills, get to the hospital, and cover any expenses they may have without having to meet any other criteria or go to any specific hospital.

As of 2020, our non-profit has not been able to establish a Funeral Assistance Fund to assist the many families who struggle with burial costs after they have lost their child unexpectedly. Often, parents are young and not financially stable; already burdened by astronomical healthcare bills, finding the money to bury their child makes the unfathomable even harder. Our organization has helped where we can over the years to reach out to funeral homes and their local resources for assistance but a fund to help families pay last respects to their child with dignity is something that is so very needed.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity



www.cdhi.org



sponsor@cdhi.org



+1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



The CDH Parent Reference Guide is a purse-size booklet that explains Congenital Diaphragmatic Hernia in layman's terms. With a glossary of medical terms, diagrams of CDH and ELSO, advice from other parents, and much more, it is the go-to guide for all newly diagnosed CDH patient families.

1. Arabic
2. Italian
3. Portuguese
4. Dutch
5. Hebrew
6. Mandarin
7. Cantonese
8. Hindi
9. Bengalese
10. Urdu
11. Swahili
12. Amharic
13. Japanese
14. Afrikaans
15. Indonesian
16. Korean

Sponsorship (\$68,800)

Translation - \$15,000 for 20 languages

Printing - \$45,750 for 25,000 copies at \$1.83 per copy

Shipping - \$8,050 at \$16.10 per 25 guides to send to 500 hospitals in the United States.

If we hire the service to translate the CDH Patient Registry, they will include the Parent Reference guides.



Parent Reference Guides

The Goal:

The CDH Guide is currently available in 6 languages, and we would like to translate it into 15 more languages to help newly affected families on a more global scale.

Steps:

1. Hire a translation company.
2. Send them the English version of the guide.
3. Tell them which 20 languages we want.
4. Wait for them to translate.
5. Make them available online.
6. Print newly translated guides and sent to hospitals.

Measurements of Success:

1. Doctors requesting more guides.
2. The percentage of new CDH Families registering on our website from countries who speak these languages.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Every year, the CHERUBS division of CDH International holds a family support conference. The location of the conferences alternates annually between the United States and Europe, with European conferences a joint effort between CDHi and fellow ACDHO charities.

Conferences include scientific speakers, round-table support sessions, the latest research and awareness efforts, as well as give the opportunity for patients and families to meet other patients and families.

All CDH Families are encouraged to attend and all researchers are welcome.



Conference Sponsorship (\$50,000)

- \$10,000 for Venue and Food
- \$40,000 for 40 grants of \$1000 for conference scholarships for families.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Grieving Parents Retreat

A 3-day weekend at an all-inclusive adult only venue where grieving parents can meet other grieving CDH parents, attend group and couples counseling sessions, find positive ways to remember their children and rest in a beautiful setting.

- Meet other grieving parents who understand how they feel
- Grief counseling
- Couples counseling
- Memorial service
- Yoga
- Night Out with other grieving parents to dance, have fun and feel "normal" again for a little while

Sponsorship - \$30,000

This will cover the costs 50 people at an all-inclusive resort in the United States with a budget of \$200 per person, per night.

CDH Camp

A week-long camp for CDH patients and their siblings, supervised by CDHi staff, nurses and a pediatric surgeon on call. The event will allow the kids to have fun, meet other children who understand how they feel, attend camp activities and do things that "normal" kids get to do - all while giving their caregivers a much-needed break.

- Meet other CDH patients and families
- Outdoor games
- Swimming
- Equine therapy
- Arts and crafts
- Group discussions
- Campfire signing and s'mores

Sponsorship - \$52,500

This will cover the costs of 90 children and 10 adults at an all-inclusive camp in the United States with a budget of \$75 per person, per day.

Volunteer Retreat

Volunteering while taking care of or grieving a child born with CDH is extremely hard and emotionally exhausting. Burn out is a very common issue among volunteers at medical nonprofits. These wonderful people give so much of themselves, and they need more rest and appreciation.

The CDHi Volunteer Retreat is 4-day weekend that allows CDH International Boards, Leaders and Volunteers to come together and work 2 days on charity goals, planning and issues, but also have 2 days to have fun together and bond to build a stronger charity team and CDH community.

Sponsorship - \$30,000

This will cover the costs 50 people at an all-inclusive resort in the United States with a budget of \$150 per person, per day.





Local CDH Support

Ways that CDH International supports patient families on a local level:

- Hospital visits (when requested)
- Funeral attendance (when requested)
- On-call emotional support
- Member picnics and get-togethers
- Local fundraisers

CHERUBS has been tirelessly working to connect families since 1998 when with a ListServ (email mailing list) and online chat. In 2001, CHERUBS created private forums on the charity website which provided families a safe, secure, and private place to discuss any issues they may be facing.

With the advancement of social media, CHERUBS created private Facebook groups for families to have a place to openly discuss their lives with CDH. In addition to online support, we encourage local family gatherings and promote local fundraising events families can participate in to support each other.

Local Support Sponsorship (\$10,000)

The venue costs of 50 events at a \$200 budget each



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Our first CDH Scholarship was established as the Oz Kidd-Ward Scholarship in memory of a child with money raised by his parents. Since then, we have awarded a dozen scholarships up to \$2500 in honor / memory of many patients and even in other family members.

It is our goal to offer \$25,000 in scholarships by 2025 to CDH survivors, siblings, or parents who are returning to school and have an interest in studying medicine to help further the fight against CDH.

Sponsorships (\$25,000)

- 3 scholarships of \$5000 each to CDH survivors
- 2 scholarships of \$5000 each to CDH siblings or parents going into the medial field



CDH Scholarships

The Goal:
To offer \$25,000 in scholarships to CDH Survivors, CDH siblings or parents of a CDH child.

Steps:

1. Get families to create a scholarship in honor or in memory of their loved one born with CDH.
2. Donate / raise a minimum of \$1000 and we will create a scholarship in their name for the upcoming new school year.
3. Families can review applications and choose the recipient.
4. Legacy scholarships or one-time scholarships can be created.

Measurements of Success:
Meeting the goal of gifting 5 CDH Scholarships to patient families by December 31, 2025.



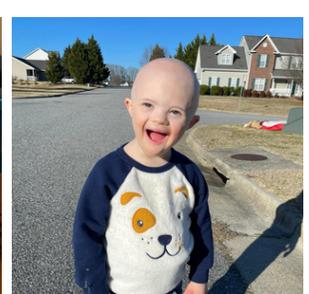
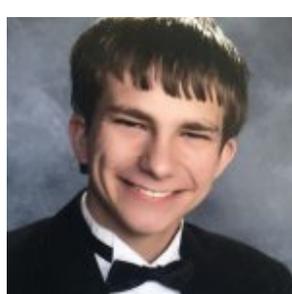
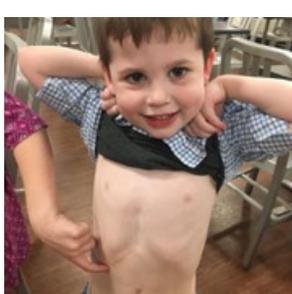
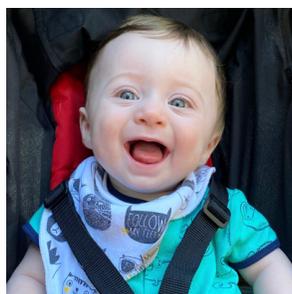
CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



Our Families



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA

Sponsorship



Service:	Cost:	Includes:
CDH HOPE Totebags Sponsorship	\$76,000	<ul style="list-style-type: none"> • Logo on the actual totebags • Logo on information packets • Logo on the t-shirts • Social Media and Blog posts
CDH Phone App Sponsorship	\$10,000	<ul style="list-style-type: none"> • Logo on the Phone App • Social Media and Blog posts
Financial Assistance Sponsorship	\$50,000	<ul style="list-style-type: none"> • Social Media and Blog posts
Funeral Assistance Sponsorship	\$30,000	<ul style="list-style-type: none"> • Social Media and Blog posts
Parent Reference Guide Sponsorship	\$68,800	<ul style="list-style-type: none"> • Logo on the Parent Reference Guides • Social Media and Blog posts
Annual CDH Conference Sponsorship*	\$50,000	<ul style="list-style-type: none"> • Logo on all event signage and programs • Logo on event t-shirts • Logo on the event webpage • Social Media and Blog posts
Grieving Parents Retreat Sponsorship	\$30,000	<ul style="list-style-type: none"> • Logo on all event signage and programs • Logo on event t-shirts • Logo on the event webpage • Social Media and Blog posts
Kids Camp Sponsorship	\$52,500	<ul style="list-style-type: none"> • Logo on all event signage and programs • Logo on event t-shirts • Logo on the event webpage • Social Media and Blog posts
Volunteer Retreat Sponsorship	\$30,000	<ul style="list-style-type: none"> • Logo on all event signage and programs • Logo on event t-shirts • Logo on the event webpage • Social Media and Blog posts
Local Support Sponsorship	\$10,000	<ul style="list-style-type: none"> • Logo on all event signage and programs • Logo on event t-shirts • Logo on the event webpage • Social Media and Blog posts
Scholarship Fund Sponsorship	\$25,000	<ul style="list-style-type: none"> • Social Media and Blog posts • Logo on Scholarship Certificate

* Also listed in the CDH International Research Sponsorship Proposal



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA

Become a Sponsor



Please complete the online registration form at www.cdhi.org/support2023. You may also print and complete the form below. You will receive a confirmation once payment is received.

Please return the completed form and your logo via email to sponsor@cdhi.org

CONTACT NAME

COMPANY/ORGANIZATION

MAILING ADDRESS

CITY

STATE/PROVINCE

ZIP/POSTAL CODE

TELEPHONE

EMAIL

WEBSITE

SIGNATURE

- CDH HOPE Totebag Sponsorship (\$76,000)
- Parent Reference Guide Sponsorship (\$68,800)
- Kids Camp Sponsorship (\$52,500)
- CDH Conference Sponsorship (\$50,000)
- Financial Assistance Sponsorship (\$50,000)
- Funeral Assistance Sponsorship (\$30,000)
- Grieving Parents Retreat Sponsorship (\$30,000)
- Volunteer Retreat Sponsorship (\$30,000)
- Scholarship Fund Sponsorship (\$25,000)
- Local Support Sponsorship (\$10,000)
- CDH Phone App Sponsorship (\$10,000)

PAYING BY CHECK?

Make payable to "CDH International", and reference "Support Sponsorship" in the memo/reference line. Please mail your check and registration form to:

CDH International
3650 Rogers Road #290
Wake Forest, NC 27587



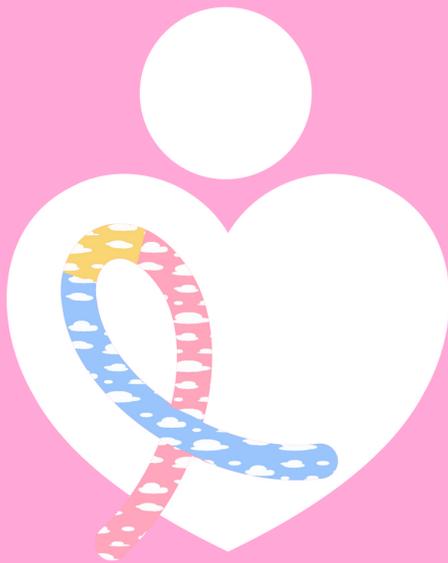
CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity

 www.cdhi.org

 sponsor@cdhi.org

 +1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA



End Of Proposal

THANK YOU



We appreciate your time and consideration and look forward to working with you to help save the lives of children born with Congenital Diaphragmatic Hernia.



CDH International is the world's oldest, largest, and leading Congenital Diaphragmatic Hernia charity



www.cdhi.org



sponsor@cdhi.org



+1 919-610-0129 / +1 919-610-2972 (cell)

CDH International
3650 Rogers Rd #290
Wake Forest, NC 27587
USA