

HOPE **for** HIE
awareness • education • support

IMPACT REPORT 2021



A MESSAGE FROM OUR LEADERSHIP

2021 has been an incredible year. The word "unprecedented" has been used quite a bit in many different contexts over the last two years.

For Hope for HIE, unprecedented has been associated with positive growth in moving forward our organizational mission. We embraced the theme of "Team Hope" this year, and truly it is a team effort.

We have collaborated in ways that were not possible before with the investments we have made into resources, both human and program-related, that we are excited to share more in this report.

We could not do this work without you and your support. We want to thank you for ensuring we have the resources that we need to continue on the path to develop more programs to support our families, and ways to move forward key research in the myriad of ways HIE can impact someone and their family.

Here's to an exciting 2022!

With Hope,

David Ford & Betsy Pilon



David Ford

President, Board of Directors



Betsy Pilon

Executive Director



ORGANIZATIONAL PROFILE

Hope for HIE is the only global nonprofit organization that serves children and families with neonatal and pediatric-acquired Hypoxic Ischemic Encephalopathy.

HIE is a leading cause of cerebral palsy, epilepsy, learning and attention differences and disabilities, and other neurodevelopmental challenges. It has a range of causes, a spectrum of outcomes, and very few therapeutic interventions available to decrease death and disability...

...but that is beginning to change.

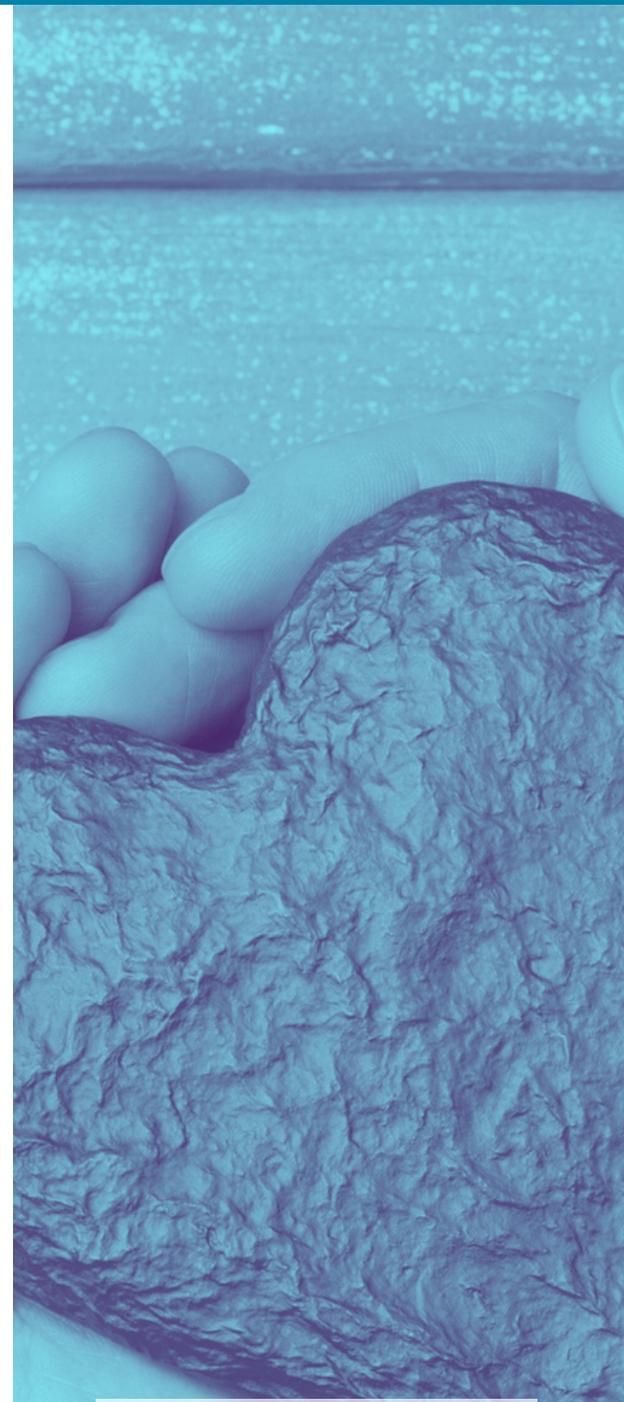
Through our accelerated organizational growth and development, we've been able to not only have a seat at the table to bring the unmet needs of our community forward, but create tables for discussion and action.

Our network represents families and providers from over **30 countries, across six continents**, around the world. We are able to facilitate projects and conversations with researchers and clinicians that other groups are not, due to our unique approach and lens as a patient and family organization.

Through our initiatives, we've been able to:

- Connect more than **7,000 families** to research-backed peer-to-peer support, host online and in-person connections, and distribute over 1,200 care packages.
- Generate awareness for HIE, the range of outcomes and impacts children and families face, advocating for improvement in care, communication and connection.
- Develop comprehensive, accessible, and vetted educational information, addressing health information literacy gaps.

Our work is only made possible by the generous support of our community, our Partners in Hope, community giving, and you.



Hope for HIE is committed to the highest integrity for financial and organizational accountability and reporting.

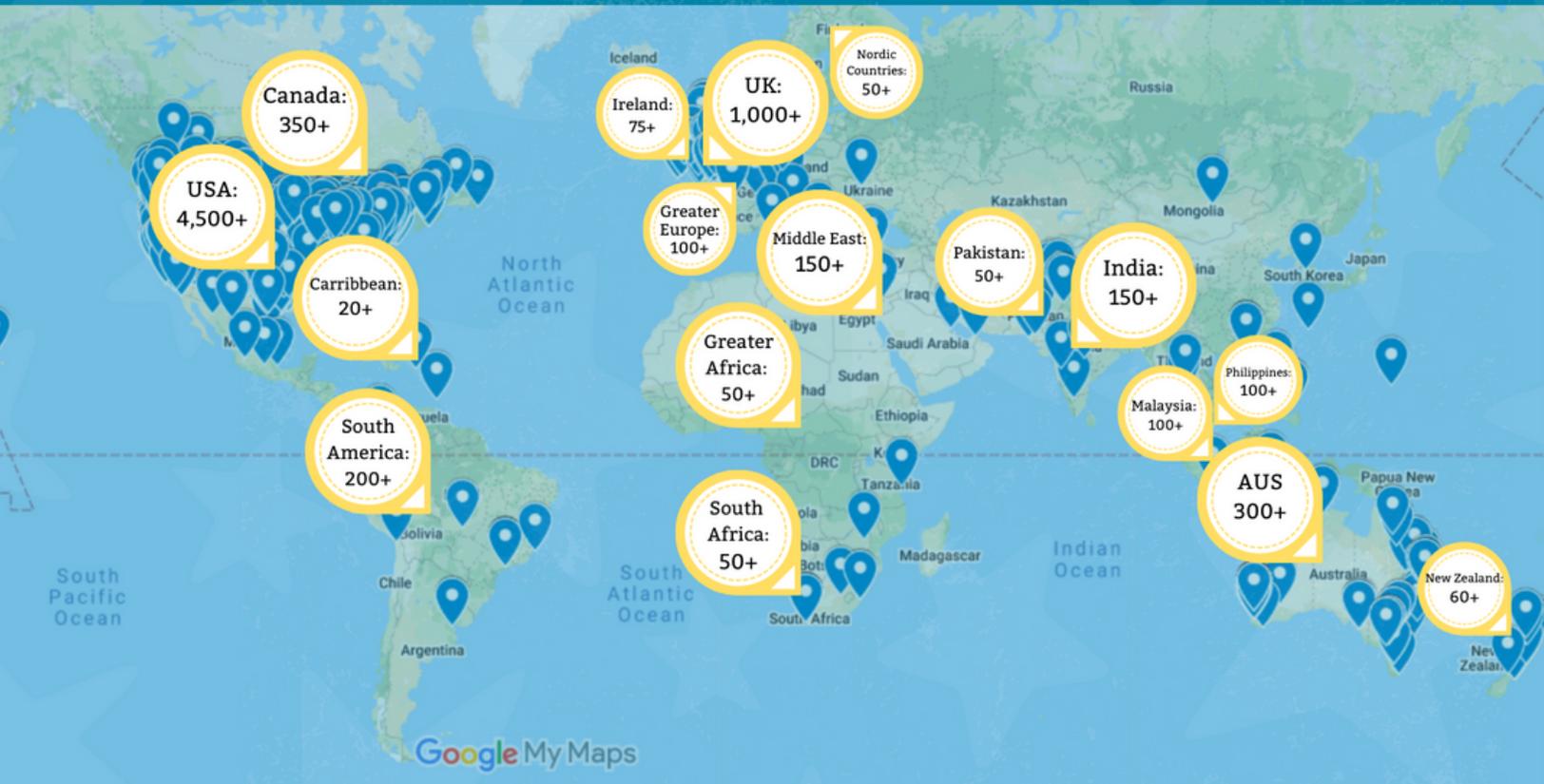
IRS EIN: 46-4038344

[Latest 990 filing](#)

2021 WORLDWIDE IMPACT

HOPE for **HIE**
awareness • education • support

Connecting families, researchers and clinicians, worldwide,
since 2010 - connecting 7,000+ today!



2021 WORLDWIDE IMPACT BY THE NUMBERS:

HIE impacts **2-3 per 1,000 live births** in developed countries, and even more in low and middle income countries.

Hope for HIE grew in 2021 to connect over **7,000 families, researchers and clinicians worldwide** from over 30 countries on six continents through our comprehensive support network, Medical Advisory Board and outreach.

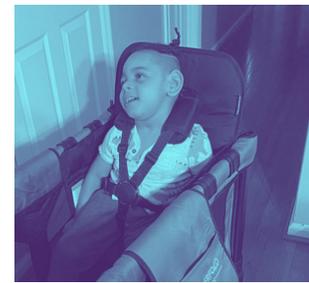
Connected to over **300 NICUs**, distributing educational materials translated into **seven** different languages.

Over **25,000 individuals** viewed Hope for HIE educational videos.

Over **100,000 individuals** from all over the world accessed Hope for HIE's website, which was relaunched in April 2021.

Over **1,000,000 impressions** were generated from HIE Awareness Month in April, worldwide.

2021 IMPACT: SUPPORT PROGRAM HIGHLIGHTS



Hope for HIE's **SUPPORT** programs are focused connecting families and empowering parents through discussion-based forums, across the age and outcome continuum, and connecting to local resources for additional support. The heart of who we are is the spirit of connection and community. What started as three moms coming together to support one another, has transformed into a comprehensive worldwide community of support with special emphasis on equity, accessibility and cultural competence, with a goal of integrating the principles of trauma-informed care.

| Program | Details | Impact |
|--|--|---|
| Comprehensive Peer-to-Peer Support Network | Support membership grew by over 15% to connect over 7,000 families, worldwide | <ul style="list-style-type: none"> Started dedicated groups in Indonesia, Malaysia, India, Pakistan to better connect families. |
| Newly Diagnosed Family Support Packages | Kits contain comfort items, connection to peer support, and educational information about HIE | <ul style="list-style-type: none"> Distributed over 1,000 support packages, including new distribution into Israel, Bangladesh and Brazil. |
| Loss Support Packages | Provided support packages for new to loss families, and a commemorative gift for all loss families | <ul style="list-style-type: none"> Supported over 25 new to loss families Sent out over 150 commemorative loss gifts |
| Cup of Hope Support Program | Distribution of Starbucks gift cards to families, NICU nurses, and clinicians | <ul style="list-style-type: none"> Distributed to over 100 nurses, 200 families and 50 other identified clinicians. |
| Peer-to-Peer Support Training Program | Two support administrators audited Child Neurology Foundation's P2P training | <ul style="list-style-type: none"> Will be launching comprehensive peer support mentorship program in 2022, with CNF training secured. |
| Hope for the Holidays Community Support Program | 10 Days of Giveaways of adaptive equipment, family quality of life equipment and experiences | <ul style="list-style-type: none"> Over 150 families benefitted from our sixth annual program, with over \$10,000 worth of items given to families. Second annual dedicated program for loss families |

2021 SUPPORT IMPACT: MEET OUR FAMILIES



The Castanon Family Texas, USA

"Hope for HIE is there to help the lifelong journey of adapting, learning, advocating and loving more than we ever thought possible. It is finding the joy and the good things along the way, even when the road is unknown."

The Burns Family Ireland

"Hope for HIE was the first place we found when Ella was born. Straight away the people welcomed us and made us feel part of this global family. We were gifted a switch toy last Christmas, and it has helped Ella tremendously with therapies and play.

It's the first place we go when we need to talk to someone who gets it."



The Benso Family New Jersey, USA

"Hope for HIE is our lifeline. To know that there is always a community to support our family, validate our feelings & experiences, and celebrate every inchstone is the most comforting. "



SUPPORT

THE PEOPLE BEHIND THE ORGANIZATION IN 2021

STAFF



Betsy Pilon
Executive Director

Hope for HIE easily outgrew the all-volunteer model in 2019, and the Board of Directors made the decision to hire an Executive Director at the end of 2020 to manage organizational operations and programming. Dedicating full time towards Hope for HIE has opened multiple doors for the organization this year, positioning it to be stronger and resourced for this growth curve.



Simran Bansal
Intern

Hope for HIE was connected to Duke University senior, Simran Bansal, through Medical Advisory Board member, Dr. Monica Lemmon. Simran has been instrumental in helping the organization with capturing stories, and is turning them into a published book. She is Hope for HIE's first intern.

BOARD OF DIRECTORS



David Ford
President



Becky Detlef
Secretary &
Loss Coordinator



Matt Kegyes
Treasurer



Crystal Kostick
Director



Amy Cutler
Director &
Medical Advisory Board
Co-Chair



Annie Goeller



Jennifer Kegyes

Our greatest thanks to Annie Goeller & Jennifer Kegyes who finished their Board terms in 2021.

MEDICAL ADVISORY BOARD

NEONATOLOGY



Lina Chalak, MD, MSCS



Mohamed El-Dib, MD



Sue Hall, MD, MSW



Nathalie Maitre, MD PhD

NURSING



Brooke Rakes, PhD, RN, RNC-NIC



Kathi Randall, RN, MSN, CNS, NNP-BC



Jayne Solomon, MSN, RNC-NIC, NNP-BC

PEDIATRIC NEUROLOGY



Monica Arroyo, MD



Bhooma Avaramuthan, MD, DPhil



Danielle Barber, MD PhD



Hannah Glass, MD



Monica Lemmon, MD



Adam Numis, MD



Renee Shellhaas, MD, MS



Janet Soul, MD

PEDIATRIC NEUROLOGY & SLEEP MEDICINE



Chris Carosella, MD



Brittney Pryor Craig, MD

NEUROPSYCHOLOGY



Seth Warschausky, PhD

PEDIATRIC DENTISTRY & ORAL SURGERY



Scott Siegel, MD, DDS, FACS, FICS, FAAP

PHYSICAL MEDICINE & REHABILITATION



Ed Hurvitz, MD

PEDIATRICS



Beth Robinson Swartz, MD

OCCUPATIONAL THERAPY



Amy Cutler, OTR/L Co-Chair



Sue Ludwig, OTR/L



Tamara Smith, OTR

PHYSICAL THERAPY



Ashley Fielding, PT



Sara Morgan, DPT

SOCIAL WORK & COUNSELING



LaTrice L. Dowtin, PhD, LCPC, NCSP, RPT



Evelyn Mascarenas, MSW



Adriana Salcido, MSW Co-Chair

SPEECH & LANGUAGE PATHOLOGY



Nancy Calamusa, MA CCC-SLP, IBCLC



Jennifer Keyges, MA, CCC-SLP



Carole Zangari, PhD, CCC-SLP



Tina Moreno, MA, CCC-SLP

VISION SPECIALISTS



Christine Roman-Lantz, PhD



Deborah Zelinsky, OD, FNORA

SUPPORT

THE PEOPLE BEHIND THE ORGANIZATION IN 2021

2021 SUPPORT IMPACT: PARTNERS IN HOPE

Medtronic

Platinum Supporter

Medtronic plc (www.medtronic.com), headquartered in Dublin, Ireland, is among the world's largest medical technology, services and solutions companies – alleviating pain, restoring health and extending life for millions of people around the world. The company is focused on collaborating with stakeholders around the world to take healthcare Further, Together.



Gold Supporter

Aspect Imaging is a global leader in the design and development of compact, MR imaging systems for Pre-Clinical and Medical applications. Through state-of-the-art technology, Aspect Imaging offers innovative point-of-care MRI solutions, including the Embrace® Neonatal MRI System, which is designed to facilitate enhanced patient-centered diagnostic capabilities inside the NICU.

tréxō robotics

Bronze Supporter

Tréxō Robotics is on a mission to redefine mobility solutions for all people. We want to create devices that are designed with the user in mind, and can be used at home. Fundamentally, we are driven by the belief that people should have access to enabling technologies. Trexo Robotics builds wearable robotics to help children with disabilities and walking impairments experience over-ground walking in clinic and at home.

2021 IMPACT: COMMUNITY CONNECTION



Laura Villanueva Germany

"I ran the race virtually with my brother back home in Germany today, it was great doing an activity together, as we haven't seen each other in almost 2 years. Thanks for the event, we ran for our friend Peter."

The Shah Family Wisconsin, USA

"Hope for HIE was a huge resource for answers and comfort as we were new parents and didn't know what was happening with our baby during those few first weeks,. As we were nearing Kiran's first birthday, we wanted to have an event to celebrate Kiran and support Hope for HIE's efforts. We are so happy to give back."



Margie Jones & Julie Makow North Carolina & Texas, USA

"Having Hope for HIE over the years through these very complex journeys that we find ourselves on, and having this support has meant everything. Meeting each other in person felt like we've been lifelong friends. It was so special to be able to reconnect on the Just Say HIE podcast."



COMMUNITY

CONNECTING THROUGH HOPE

Connecting through our community has definitely been impacted by the global pandemic. Our families have found themselves more isolated than usual, and restrictions to safely hold in-person events have been a hardship. After postponing our retreats set to be held in 2020, we had hoped to be able to host in 2021, but that was not to be.

Instead, we celebrated the opportunities when two families met up at a beach, or a park, or anywhere else to safely connect, and celebrated the two events we were able to support to connect our global community offline.

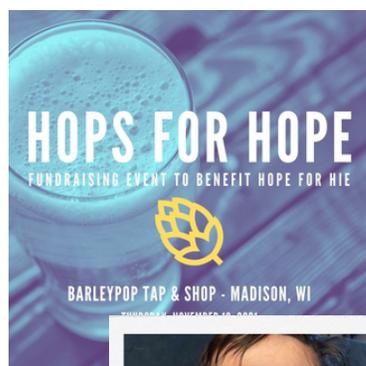
HUSTLE FOR HOPE 5K - RUN, WALK, ROLL

Our second annual Hustle for Hope 5K grew exponentially in 2021. We saw over 400 registrants, and people gathered across the globe to safely participate, either in small groups outdoors, or individually.

Some families even took this opportunity to train and participate with family members who live in different countries. We were able to raise over \$20,000 through this, and above all, families felt connected to each other once again.

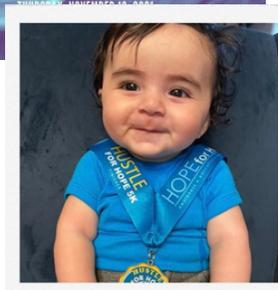
We had terrific sponsors like Trexo Robotics, Bonfire, and the Ford family.

We hope to host regional low-key gatherings for the 2022 Hustle for Hope.



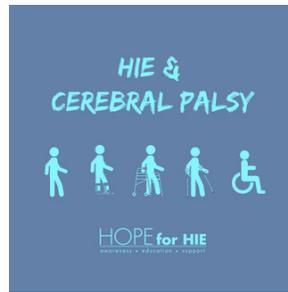
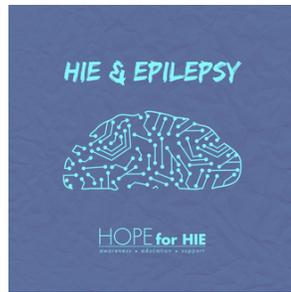
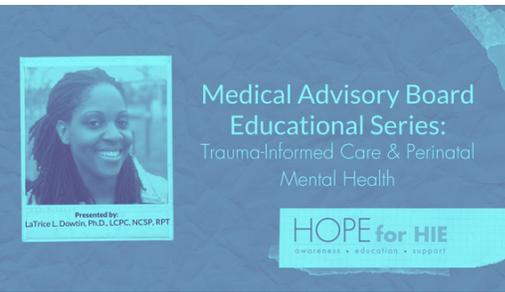
HOPS FOR HOPE MAKES ITS WAY TO MADISON, WISCONSIN

With their son Kiran's first birthday, Sarah & Ravi Shah wanted to find a way to support Hope for HIE, and celebrate this milestone birthday. Partnering with BarleyPop Tap & Shop, guests had an opportunity to bid on some exciting baskets, get some Hope for HIE swag, and enjoy Kiran's birthday celebration for a cause raising nearly \$2,500!



We are so grateful for the Shah family for their incredible work to make this event happen, for BarleyPop Tap & Shop's support, and for the community of support that rallied around this event and their family during the first year on the HIE journey.

2021 IMPACT: EDUCATIONAL PROGRAMS



Hope for HIE works diligently to provide **EDUCATION** for its global community. Families of children with HIE face many obstacles accessing educational materials that we are working to remove. Health information literacy is one of the most significant issues impacting families, in addition to the lack of connecting families early, naming the diagnosis, and providing educational materials that are written free of medical jargon and available in first use language for our diverse population.

| Program | Details | Impact |
|---|--|---|
| Medical Advisory Board video educational series | MAB members lent their expertise to develop Q&A resources for HIE families | Over 20,000 individuals viewed 15 produced videos with experts addressing the biggest challenges in the HIE community. |
| HIE NICU Educational Timeline Materials | Trifold handout developed for patient education at time of diagnosis in the NICU | <ul style="list-style-type: none"> Over 3,000 copies printed in seven languages and distributed to over 150 NICUs, worldwide Also available for download via website |
| Educational Materials: HIE | Developed new website with additional educational toolkits distributed for HIE Awareness Month and beyond | <ul style="list-style-type: none"> Over 1,000,000 impressions recorded of the toolkit across communication channels |
| Educational Materials: Epilepsy, Cerebral Palsy, Learning & Attention, Sleep | Developed several educational blog posts and social media educational toolkits relating to the biggest identified educational gaps in subsequent diagnoses from HIE. | <ul style="list-style-type: none"> Seizure First Aid resources viewed over 20,000 times Newly diagnosed resources for Infantile Spasms and Epilepsy in Childhood viewed over 15,000 times All About Tone blog post viewed over 10,000 times |
| Educational Materials: Parent Mental Health | Developed several educational blog posts, videos, and social media educational toolkits relating for parents AND providers. | <ul style="list-style-type: none"> Published article in conjunction with the Newborn Brain Society with practice points regarding parent mental health in Society of Fetal & Neonatal Medicine journal. Toolkits viewed over 12,000 times. |

2021 EDUCATIONAL IMPACT: FAMILY STORIES



Hope Riley
South Carolina, USA

"I had never heard of Infantile Spasms before learning about them through Hope for HIE. We were originally misdiagnosed, which caused me to dig deeper and advocate strongly for my son, fully supported and empowered by the information I had learned through the organization."

Catie Flores
Illinois, USA

"I can honestly say that through all of these educational resources, I am continuously learning new things, especially about disabilities and ableism. I love that things that are posted are also shared in a way that it reaches so many people. I have shared so many postings on my social media so that I can continue to educate family and friends."



Tracey Johnson
England, United Kingdom

"When my son experienced his first seizure since NICU at age 7, I reconnected with Hope for HIE and the educational resources created for families whose children develop epilepsy later in childhood. I think we will need this group and resources more than ever as we navigate a new normal for our family."



2021 IMPACT: AWARENESS & ADVOCACY



Hope for HIE's **AWARENESS** programs are focused on bringing light to and improving the unmet needs of the HIE community. We not only create collaborations, advocacy programs and awareness, but are regularly invited to collaborate as a strong patient advocacy partner by clinicians, researchers, biotech and pharma in the newborn and pediatric brain community.

| Program | Details | Impact |
|---|--|--|
| HIE Awareness Month 2021 Campaign | Developed multi-faceted community awareness tools to educate and inform | <ul style="list-style-type: none"> Over 1,000,000 impressions recorded of the toolkit across communication channels Over 300 new families found support |
| Name the Diagnosis Social Media Campaign | Targeted social media influencers in neonatal neurology to understand the impact of not naming the diagnosis to families | <ul style="list-style-type: none"> Significant engagement from those targeted to understand impact of this specific trauma. Included HIE family voices in published article in SFNM |
| HIE Remembrance Day | April 20th is dedicated to HIE loss, educational and family stories are shared, Wave of Light candlelight service held. | <ul style="list-style-type: none"> Recognition of more than 400 loss families. Wave of Light service included over 500 families worldwide. |
| Rare Epilepsy Advocacy | Identify and participate in advocacy relating to HIE equity and inclusion as a leading cause of several rare epilepsies such as Infantile Spasms, Lennox-Gastaut Syndrome & ESES | <ul style="list-style-type: none"> Member of the Rare Epilepsy Network since 2019, participated in the American Epilepsy Conference in December 2021 Outreach to connect with ESES leading researchers for more equity of inclusion in research and collaborations. Enrolling families into new trial. |
| Just Say HIE Podcast | A new communication channel for families to share their unique HIE experiences & connections. | <ul style="list-style-type: none"> Published ten episodes from April to December 2021 with over 1,000 listeners from all over the world across six channels. |
| Advocacy for Equity in the NICU | Messaging with NICU traditionally leaves HIE families and other full term experiences out. Equity is essential to reduce trauma. | <ul style="list-style-type: none"> Worked with the NICU Parent Network and Project Sweet Peas on inclusive messaging for NICU Awareness events and campaigns. Speaking opportunities listed separately. |

ADVOCACY

BRINGING HIE TO THE FOREFRONT OF RESEARCH & CLINICAL CARE



One of the biggest advantages of bringing on a full time Executive Director at the end of 2020, was the ability to have more dedicated time to build and participate in collaborations that bring awareness to the unmet needs of the HIE community, and those that would benefit more broadly in the child neurology space. We've connected with researchers, clinicians, organizations and more.

We were so grateful to have the following opportunities to have our community needs heard, and to collaborate in 2021:

SPEAKING OPPORTUNITIES

- **Supporting HIE Families**, Cleveland Clinic Neonatal Brain Club Grand Rounds, January 2021
- **Supporting Families Facing HIE**, Graven's Conference on the Care for High Risk Newborns, March 2021
- **Do You See What I See? MRI Day from the Parent Perspective**, NICU Leadership Forum, April 2021
- **Hope for HIE: A Parent Perspective**, NICU Grad Podcast, April 2021
- **Living Your Best Life with HIE**, Empowering NICU Parents Podcast, April 2021
- **Family Care in the NICU**, Newborn Brain Society Webinar, April 2021
- **Supporting HIE Families**, ICON Webinar, May 2021
- **Sleep Issues in Children with Neurological Conditions**, Child Neurology Foundation Webinar, June 2021
- **HIE, Hope & The Art of Communicating in the NICU**, The Incubator Podcast, June 2021
- **Do You See What I See? MRI Day from the Parent Perspective**, Embrace Webinar, July 2021
- **Hope, Healing & the Powerful Impact of NICU Experiences**, Qeepsake Instagram LIVE event, September 2021
- **Improving Critical Transitions**, Vermont-Oxford Network Annual Quality Conference, October 2021

COLLABORATIONS & WORKGROUPS

- Child Neurology Foundation Telehealth Workgroup
- Child Neurology Foundation Better Together for Kids Coalition
- American Academy of Neurology Child Neurology Measures Task Force
- Neonatal Seizure Registry
- Mild HIE Research Advocacy
- Rare Epilepsy Network
- Infantile Spasms Action Network
- Newborn Brain Society, Board of Directors
- Songs of Love Foundation Custom Song Program
- NICU Awareness Month Partner Organization
- Epilepsy Foundation & American Brain Coalition Legislative Advocacy



AWARENESS

ENSURING EQUITY & INCLUSION

We know that families find a lot of information on the internet, and that sometimes it is hard to weed through what is factual and what is not. When HIE isn't even listed as a neurological diagnosis, despite it impacting 2-3 per 1,000 live births in neurology-related, and especially pediatric neurology-related directories, we go to work to ensure equity of inclusion and naming the diagnosis. Sometimes, it's a matter of learning the vernacular, like HIE is a structural cause of epilepsy vs. genetic.

Here's where we have expanded our inclusion in 2021:

DISORDER DIRECTORIES & WEBSITES:

- NIH National Institute of Neurological Disorders and Stroke
- Child Neurology Foundation
- Epilepsy Foundation of America
- Lennox-Gastaut Syndrome Foundation (HIE is a top cause of LGS)



NICU Support Organizations and clinicians and researchers in the NICU space that support all NICU families have often left out full-term NICU experiences, and have not connected families to diagnosis-specific resources. Messaging five years ago rarely mentioned anything but the preemie experience, which we know exacerbates the trauma full term NICU families experience. We are grateful for the following organizations and NICU providers who have shifted to more inclusive messaging, and in particular highlighted HIE this year:

NICU SUPPORT EQUITY IN MESSAGING & INCLUSION:

- **Hand to Hold:** Inclusion in referrals, updated messaging, and support of HIE Awareness Month
- **NICU Parent Network:** Updated messaging, support of HIE Awareness Month
- **NICU Social Media Influencers:** Updated messaging, updated hashtag use, education on impact of trauma to HIE families not included.
- **NICU Awareness Month:** Project Sweet Peas updated messaging and specifically developed educational information regarding the impact of full term NICU experiences
- **Empowering NICU Parents podcast:** episode that aired during HIE awareness month



2021 AWARENESS & ADVOCACY IMPACT



Shiru Wanjiru
Nairobi, Kenya, Africa

"Children with disabilities in my community are often associated with taboo. Having Hope for HIE bring awareness helps improve access to treatments, programs and acceptance. Hope for HIE has been that place of acceptance in my journey where I feel normal, and can spread awareness."

Georgie Oliver
Arizona, USA

"I found the HIE diagnosis listed in the extended notes after I requested medical records at 3 months when she was diagnosed with infantile spasms. The worst part is that it was listed as mild and RESOLVED... spoiler alert: it is neither of those things. Families like ours need the diagnosis named in the NICU to ensure we know what our children are at risk for and are connected to resources."



Ashlee Hamlett
North Carolina, USA

"Becoming involved with NSR has allowed me to share the parental side of studies in regards to using easy to understand language and providing recommendations to raise awareness. Advocating for the needs and wants of families, especially HIE families, has always been a passion of mine and this has been a great opportunity."

Lindsey Garrett
Georgia, USA

"This year, we got involved in our local community to spread awareness, through the city of Hahira's "parade of trees". Hudson's Mimi came up with the idea to do one for HIE awareness. Hudson was so happy when he saw his tree. I'm so excited for our little town to learn more about HIE. If one person walks away knowing more, than I am happy."



LOOKING AHEAD

PRIORITIES IN 2022 AND BEYOND

With solid footing under us thanks to the continued and increased support from our global HIE community, we are eager to get started on our upcoming priorities in 2022. These include:

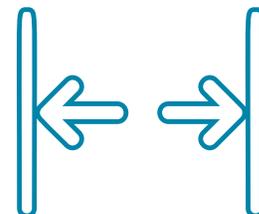


EXPANDED MENTAL HEALTH & PEER SUPPORT FOR OUR FAMILIES

- Contracting a Social Worker to run monthly video support group meetings, connect families to community resources and more.
- Development of formal peer mentorship program for Newly Diagnosed families with comprehensive trauma-informed training provided through the Child Neurology Foundation
- Redevelopment of admin and moderator group programs, investing in paid administrators to be more actively involved in support group management.

CLOSING THE GAP FROM DISCHARGE TO FOLLOW UP APPOINTMENTS

- Development of educational modules for HIE families (and staff) in discharge planning from NICU and PICU to educate on seizure risk and identification, Infantile Spasms, building positive partnerships with medical teams, and mental health support.
- Collaborating with existing NICU and PICU discharge resources to ensure interoperability.



BRINGING THE GLOBAL HIE COMMUNITY TOGETHER

- In Fall 2022, we'll bring together the global HIE community for the inaugural **Hope for HIE Global Community Conference**.
- This will be a multidisciplinary event focused on collaboration, problem-solving unmet needs, and learning about the latest research and discoveries from families, clinicians, researchers and industry.
- We hope to have some satellite gatherings, but will make determinations on in-person feasibility later in the year depending on the state of the global COVID pandemic.

EXPAND RESOURCES TO ENSURE QUALITY OF PROGRAMS CONTINUE

- In addition to the contracted social worker and some paid support network administrators, onboard a contracted program manager for large-scale and ongoing programs such as our fulfillment of support packages to NICUs, loss families, and hospitalizations, Hustle for Hope program administration and coordination of our Global Community Conference.
- Continue to engage colleges and universities to further develop our internship program.

FINANCIAL SUSTAINABILITY

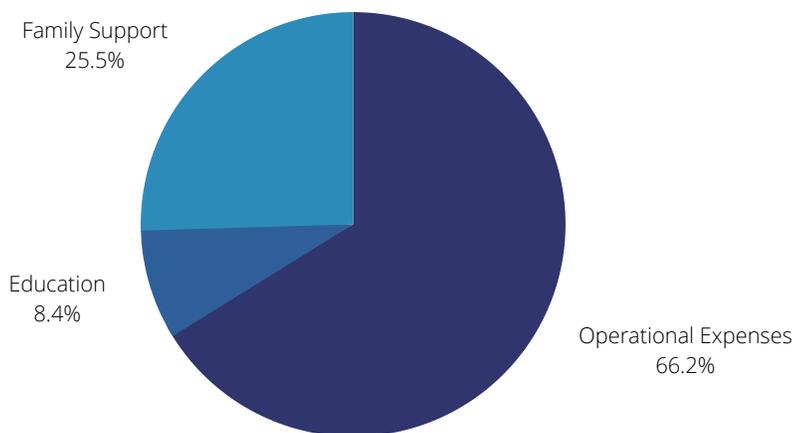
2021 FINANCIAL HIGHLIGHTS

Hope for HIE's Board of Directors made a significant decision in 2020 to invest in contracting a full-time Executive Director for the 2021 year to aide in the organization's growth and opportunities. This allowed the organization to grow significantly in 2021, programmatically and financially, allowing the organization to make additional investments expanding programs and resources into 2022, as the organization continues rapid growth.

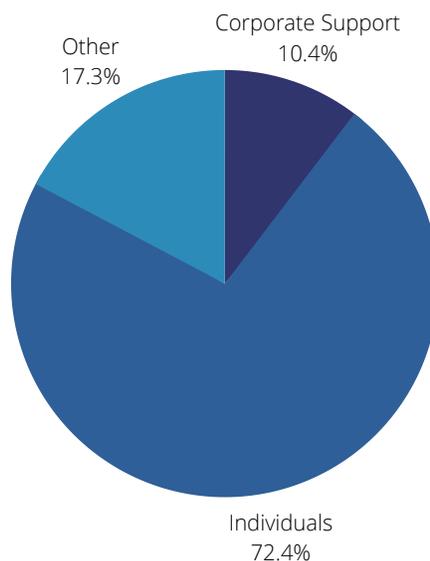
Key highlights:

- Over **\$168,000** raised from donor contributions and fundraising activities
- Nearly **\$50,000** spent on family support, outreach, awareness, and education
- Net assets of **\$278,000** as of December 31, 2021

2021 EXPENSES



2021 REVENUE



YEAR ENDED DECEMBER 31, 2021



HOPE FOR HIE'S ORGANIZATIONAL MISSION:

To improve the quality of life for children and families impacted by neonatal and pediatric-acquired Hypoxic Ischemic Encephalopathy through Awareness, Education & Support.

We want to profusely thank everyone who has contributed in any way to support our organization as we continue on this significant growth curve accelerating our mission through our programs, partnerships, and resources.

HOPE for **HIE**
awareness • education • support

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