

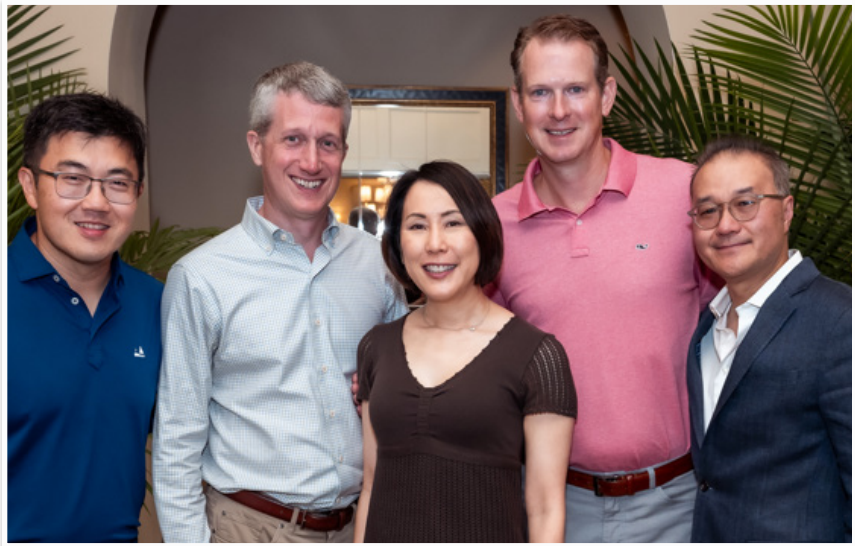
CONNECTIONS



2023 ANNUAL REPORT



CONTENTS



CCF Board Members
(L-R): Won Choi,
Brian Nold, Lisa Yue,
Carney Hawks, Eddie
Yu. Carolyn Kong and
Raymond Yue not shown.

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LETTER FROM THE FOUNDER & EXECUTIVE DIRECTOR



Dear Friends,

As the founding board president and executive director, we're proud to highlight our work in 2023 focused on fostering **connections and collaboration**. This year we strengthened existing ties and forged new relationships, all in support of our mission to assist families and children affected by cardiomyopathy.

In 2023, we built on our collaboration with the **American Heart Association** (AHA) to address the need for better medical management of children with cardiomyopathy. Led by CCF's Chief Medical Officer, Steven Lipshultz, MD, with the involvement of other CCF medical advisors, Steven D. Colan, MD, and Shelley D. Miyamoto, MD, a scientific statement on pediatric cardiomyopathy was published in AHA's peer-reviewed medical journal, *Circulation*. CCF has been a long-standing member of the AHA's **Young Hearts Pediatric Heart Failure & Transplantation Committee** and supported this collective effort to advance knowledge in this very specialized field.

We had a dynamic year connecting with several strategic partners, elevating CCF's presence nationally and globally. This included interviews with Dr. John Whyte, **WebMD's** chief medical officer and the **Transplant Families** monthly podcast and presenting at the **Global Cardiovascular Clinical Trialists** forum. CCF also participated in the inaugural **Global Heart Hub** Cardiomyopathy Patient Council meeting in Italy.

Beyond research and education, we've extended our patient support services with the launch of a "Heart to Heart" chat series to give newly diagnosed families a safe and supportive environment to discuss the challenges of living with a chronic heart disease. We remained focused on disease awareness with our social media campaigns and partnerships with heart-related organizations during **National Heart Month** and **Children's Cardiomyopathy Awareness Month**.

Through our many connections and collaborations, we are working towards our mission of earlier diagnoses, enhanced treatments, and eventual cures. We know that our work is making an impact based on the many family reviews that we receive each year. For 13 years, CCF has been recognized as a top-rated health organization by **Great NonProfits**.

Looking ahead to 2024, plans are already underway to host the 6th International Conference on Cardiomyopathy in Children with the Cincinnati Children's Heart Institute and Pediatric Cardiomyopathy Registry. We're looking forward to bringing the best minds in the field together and offering more engagement opportunities for patients and families.

Sincerely,

Lisa Yue
Founder & Board President

Gina Peattie
Executive Director

FOUNDATION OVERVIEW

The mission of the **Children's Cardiomyopathy Foundation (CCF)** is to accelerate the search for causes and cures for pediatric cardiomyopathy through increased **research, education, awareness and advocacy**, as well as to provide **support** to affected families.



Platinum
Transparency
2023

Candid.

DISEASE FOCUS

- Dilated Cardiomyopathy (DCM)
- Hypertrophic Cardiomyopathy (HCM)
- Restrictive Cardiomyopathy (RCM)
- Arrhythmogenic Right Ventricular Cardiomyopathy (ARVC)
- Left Ventricular Noncompaction Cardiomyopathy (LVNC)

BOARD OF DIRECTORS

Lisa Yue
President & Founder

Carney Hawks
Treasurer

Raymond Yue
Secretary

Won Choi

Carolyn Kong

Brian Nold

Eddie Yu

MEDICAL ADVISORS

Wendy Chung, MD, PHD
Boston Children's Hospital

Steve Colan, MD
Boston Children's Hospital

Daphne Hsu, MD
Children's Hospital at Montefiore

Steve Lipshultz, MD
*University at Buffalo School of
Medicine and Biomedical Sciences*

Shelley Miyamoto, MD
Children's Hospital Colorado

Joseph W. Rossano, MD, MS
Children's Hospital of Philadelphia

Jeff Towbin, MD
Le Bonheur Children's Hospital

CCF STAFF

Gina Peattie
Executive Director

Sheila Gibbons
Senior Director of Marketing

Liz Newsome
Project & Events Manager

Grace Weng
Operations Coordinator

CONNECTIONS



RESEARCH

RESEARCH

It was a year of progress as CCF extended its research partnership with the North America Pediatric Cardiomyopathy Registry (PCMR) and pursued new initiatives to advance understanding of the disease and enhance clinical care for children with cardiomyopathy.

AHA SCIENTIFIC STATEMENT ON PEDIATRIC CARDIOMYOPATHY

As a member of the American Heart Association (AHA) **Young Hearts Pediatric Heart Failure & Transplantation Committee** (PCMR), CCF supported the development and publication of a scientific statement on treatment strategies on pediatric cardiomyopathy in *Circulation*, AHA's peer-reviewed medical journal. This effort was led by CCF Medical Advisor, Steven Lipshultz, MD with CCF Medical Advisors, Steven D. Colan, MD, and Shelley D. Miyamoto, MD on the writing committee.

Treatment Strategies for Cardiomyopathy in Children: A Scientific Statement From the American Heart Association

BOGLE C, COLAN SD, MIYAMOTO SD, CHOUDHRY S, BAEZ-HERNANDEZ N,
BRICKLER MM, FEINGOLD B, LAL AK, LEE TM, CANTER CE, LIPSHULTZ SE

Circulation, June 8, 2023



“The PCMR has helped us advance and refine future cause-specific therapy options for pediatric cardiomyopathy.”

STEVEN E. LIPSHULTZ, MD
CCF CHIEF MEDICAL OFFICER & PROFESSOR OF PEDIATRICS
UNIVERSITY AT BUFFALO JACOBS SCHOOL OF
MEDICINE & BIOMEDICAL SCIENCES

RESEARCH

CCF FUNDED RESEARCH FINDINGS

In 2023, CCF-supported research resulted in the following medical presentations and publications.

MEDICAL PUBLICATIONS

Circulating MicroRNAs Identify Early Phenotypic Changes in Sarcomeric Hypertrophic Cardiomyopathy

SUCHAROV, CC, NELTNER B, PIETRA AE, KARIMPOUR-FARD A, PATEL J, HO CY, AND MIYAMOTO SD

Circulation: Heart Failure, March 2023

Circulating and Cardiac Tissue miRNAs in Children with Dilated Cardiomyopathy

HAILU FT, KARIMPOUR-FARD A, NELTNER B, STAUFFER BL, LIPSHULTZ S, MIYAMOTO SD, SUCHAROV CC

Journal of Cardiovascular Development, September 2023

Cardiac imaging and biomarkers for assessing myocardial fibrosis in children with hypertrophic cardiomyopathy

AKIRMANI S, WOODARD PK, SHI L, HAMZA TH, CANTER CE, COLAN SD, PAHL E, TOWBIN JA, WEBBER SA, ROSSANO JW, EVERITT MD, MOLINA KM, KANTOR PF, JEFFERIES JL, FEINGOLD B, ADDONIZIO LJ, WARE SM, CHUNG WK, BALLWEG JA, LEE TM, BANSAL N, RAZOKY H, CZACHOR J, LUNZE FI, MARCUS E, COMMEAN P, WILKINSON JD, LIPSHULTZ SE.

Journal of the American Heart Association, October 2023

MEDICAL PRESENTATIONS

Pre-emptive Treatment of Cardiac Dysfunction in Pediatric Oncology Patients

INTERNATIONAL PEDIATRIC VAD AND HEART FAILURE SUMMIT

St. Louis, MO, September 2023

Pediatric Cardio Oncology: The Past, The Present, The Future

INTERNATIONAL CARDIOONCOLOGY SOCIETY MEETING

Virtual, September 2023

The Genetic Architecture of Pediatric Cardiomyopathy

UNIVERSITY AT BUFFALO CLINICAL AND TRANSLATIONAL SCIENCE INSTITUTE (CTSI) ANNUAL FORUM

Buffalo, NY, March 2023

PHACS Cardiac Toxicity Study: Preliminary Results

PEDIATRIC HIV/AIDS COHORT STUDY MEETING
Bethesda, MD, May 2023

Anthracycline Induced Cardiotoxicity and Effective Ways of Cardioprotection

THE INTERNATIONAL CARDIO-ONCOLOGY SOCIETY IN COLLABORATION WITH THE GLOBAL CANCER CONSORTIUM (GCC)

Virtual Course for IC-OS Certification
Exam Eligibility

RESEARCH

PCMR SCHOLARS OF DISTINCTION

CCF believes in the importance of investing in the next generation of pediatric cardiomyopathy researchers and thought leaders. In 2023, Sonya Kirmani, MD and Justin Godown, MD were named Pediatric Cardiomyopathy Registry (PCMR) Scholars of Distinction for their research efforts on multi-center studies funded by CCF.

Dr. Sonya Kirmani is the Medical Director of the Pediatric Heart Failure, Ventricular Assist Device (VAD), and Transplant Program at the University of Wisconsin School of Medicine and Public Health.



Dr. Justin Godown is the Associate Director of the Pediatric Heart Transplantation Program and Medical Director of the Pediatric Cardiomyopathy and Pediatric Cardio-Oncology Program at Monroe Carell Jr. Children's Hospital at Vanderbilt.



RESEARCH

PEDIATRIC REPRESENTATION AT GLOBAL CARDIOVASCULAR CONFERENCE

CCF Executive Director, Gina Peattie, presented “Pediatric Cardiology Trials” and “Pediatric Cell and Gene Therapy Trials” at the Global Cardiovascular Clinical Trialists Forum (CVCT) December 1–2. Held in Washington DC, CVCT is the largest global gathering of physicians involved in cardiovascular clinical trials.



ABOVE Elfriede Pahl, MD, Northwestern Feinberg School of Medicine; Gina Peattie, Children's Cardiomyopathy Foundation; Daphne Hsu, MD, Children's Hospital at Montefiore (L-R)



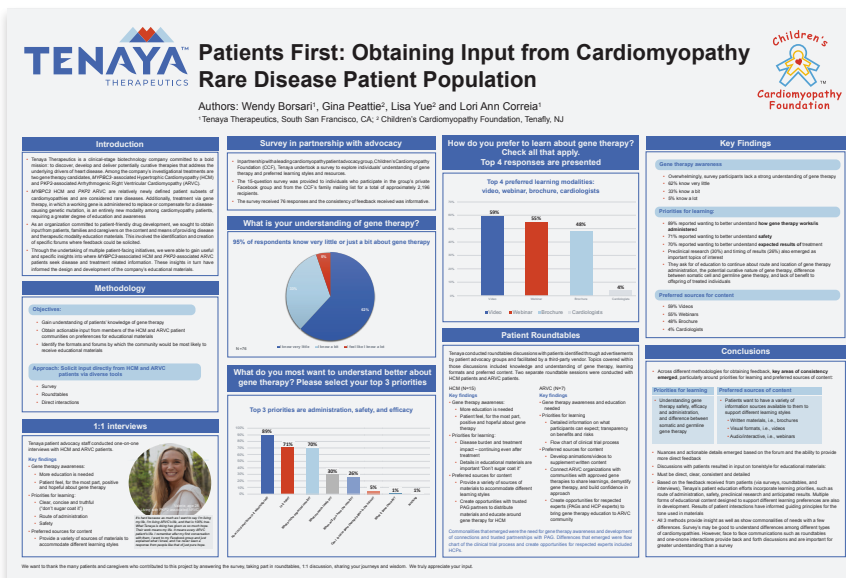
“Connecting with the Children’s Cardiomyopathy Foundation at the CVCT Forum to explore and discuss cardiovascular clinical trials is crucial for advancing cures for rare diseases like pediatric cardiomyopathy. These forums provide a platform for collaboration, enabling us to make significant strides in improving patient outcomes and bringing hope to countless families.”

ELFRIEDE PAHL, MD
PROFESSOR EMERITA OF PEDIATRICS
NORTHWESTERN UNIVERSITY FEINBERG SCHOOL OF MEDICINE

RESEARCH

INSIGHT GATHERING ON GENE THERAPY

CCF collaborated with Tenaya Therapeutics on a caregiver survey, “Patients First: Obtaining Input from Cardiomyopathy Rare Disease Patient Population,” to understand current perceptions on gene therapy and to determine educational needs among CCF family members affected by cardiomyopathy. Survey findings were presented by Tenaya Therapeutics in May at the 2023 World Orphan Drug Congress in Washington, D.C.



“We believe that building a strong partnership with CCF enables us to address patient needs more effectively and improve patient outcomes. Collaborating with CCF on our Patients First study was instrumental in identifying the educational needs of the patient population.”

WENDY BORSARI

ASSOCIATE DIRECTOR OF PATIENT ADVOCACY

TENAYA THERAPEUTICS

CONNECTIONS



EDUCATION

EDUCATION

Throughout the year, the Children's Cardiomyopathy Foundation focused on disease state education on topics of interest from the patient and caregiver community.

EDUCATIONAL WEBINARS

CCF's "Experts on Pediatric Cardiomyopathy" webinar series covered eight educational topics. It featured pediatric cardiomyopathy specialists from various medical centers and generated over 2,400 views.

Peer Support for Patients Transitioning to Adult Care

SAMANTHA J. ANTHONY, PHD, MSW

JIA LIN, MPH

Hospital for Sick Children Research Institute

Exercise, Sports, & Cardiac Rehabilitation Considerations for Children with Cardiomyopathy

JONATHAN EDELSON, MD

Children's Hospital of Philadelphia

Updates in Medical Management for Pediatric Heart Failure Patients

NEHA BANSAL, MD

Mount Sinai Kravis Children's Hospital

Pediatric Cardiomyopathy: What Newly Diagnosed Families Need to Know

BETSY BLUME, MD

Boston Children's Hospital

ICD Considerations for Pediatric Cardiomyopathy Patients

HEATHER GIAcone, MD

LAUREN SCHNEIDER, PSYD

Lucile Packard Children's Hospital

Sudden Cardiac Arrest: Education & Prevention for Pediatric Cardiomyopathy Families

ROBERT WHITEHILL, MD

Children's Healthcare of Atlanta

Genetic Testing and Gene Therapy: Updates and Future Directions in Pediatric Cardiomyopathy

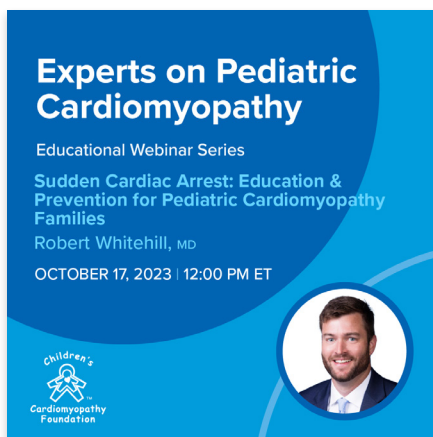
ANDREW LANDSTROM, MD, PHD

Duke Children's Hospital

Overview of Clinical Trials

JENNIFER SU, MD

Children's Hospital Los Angeles



EDUCATION

WEBMD FEATURE ON PEDIATRIC CARDIOMYOPATHY

CCF partnered with WebMD to produce an episode, “A Medical History Helps Unravel a Mystery,” for their “Champions of Change” heart disease video series, which was featured during National Heart Month and Children’s Cardiomyopathy Awareness Month.

Gina Peattie, CCF’s executive director, spoke with Dr. John Whyte, chief medical officer of WebMD about cardiomyopathy in children. The episode was viewed more than 30,000 times on social media by the public.

Gina Peattie, Advocate



Video

A Medical History Helps Unravel a Mystery

Early diagnosis is key for treating HOCM. Family medical history, highlighting heart-related conditions, and genetic testing can help pinpoint a mutation.



EDUCATION

OUTREACH TO THE PEDIATRIC HEART TRANSPLANT COMMUNITY

Since cardiomyopathy is a leading cause for heart transplants in children, CCF partnered with Transplant Families during National Pediatric Transplant Week in April. At the 2023 Pediatric Transplant Conference Executive Director, Gina Peattie, moderated a virtual session on “Genetic Testing Before & After Transplant.” More than 450 families attended with a third comprised of heart transplant families.



During National Heart Month in February, Gina Peattie was a guest on Transplant Families monthly podcast, “Growing the Gift.”



CONNECTIONS



ADVOCACY

ADVOCACY

In federal advocacy, CCF weighed in on public policies related to increasing research, education, and access to specialized care.

CHRONIC DISEASE COALITION

CCF broadened its partnerships with health advocacy organizations by working with the Chronic Disease Coalition. CCF elevated the profile of cardiomyopathy on their website and provided a new engagement platform as well as advocacy resources and training to caregivers.

ORGAN PROCUREMENT & TRANSPLANTATION NETWORK

CCF submitted a public comment to support the OPTN request to modify the heart transplant policy to allow pediatric candidates access to incompatible blood type heart, heart-lung, or lung organs. With increased transplant options, this policy modification would potentially improve waitlist mortality.

CONGRESSIONALLY DIRECTED MEDICAL RESEARCH PROGRAMS

Every year, CCF works with the Defense Health Research Consortium, consisting of 126 organizations, to request congressional support of the Defense Appropriations Act. This bill funds the Congressionally Directed Medical Research Programs and includes cardiomyopathy-related research studies.



“Cardiomyopathy is often a silent disease and may be difficult to detect without symptoms. Providing educational materials on cardiomyopathy to more families has the potential to save the lives of children at risk of sudden cardiac arrest.”

GINA PEATTIE
EXECUTIVE DIRECTOR
CHILDREN'S CARDIOMYOPATHY FOUNDATION

ADVOCACY

CARDIOMYOPATHY HEARTS ACT

Over the years, CCF has worked closely with Congressman Frank Pallone (NJ-06) on the Cardiomyopathy Health, Education, Awareness, Research and Training in Schools (HEARTS) to reduce sudden cardiac death among young people. CCF supported Rep. Pallone's reintroduction of the Cardiomyopathy HEARTS Act in December. The bill proposes that the Centers for Disease Control (CDC) develop educational materials and resources on cardiomyopathy for public awareness and distribute materials to schools, teachers, and parents.



PHOTO CREDIT
Office of Congressman
Frank Pallone, Jr. (NJ-06)

“It’s critical that we raise awareness about the causes of sudden cardiac arrest and ensure schools are more prepared to deal with cardiac emergencies so we can prevent these tragic deaths. I’m proud to introduce the HEARTS Act to take these much-needed measures to combat this devastating condition in young people.”

CONGRESSMAN FRANK PALLONE, JR. (NJ-06)

CONNECTIONS



AWARENESS

AWARENESS

National Heart Month in February and Children's Cardiomyopathy Awareness Month in September are CCF's two disease awareness initiatives.



NATIONAL HEART MONTH

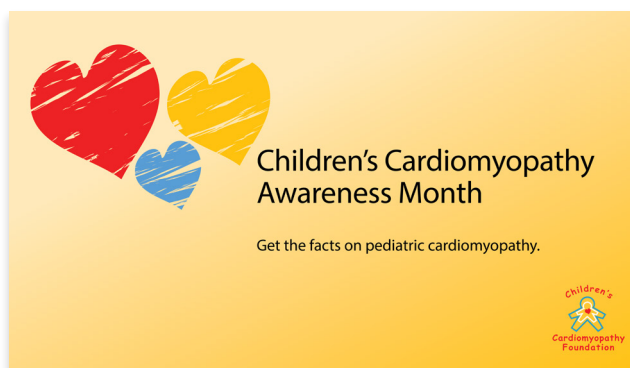
- WebMD launched their "Champions of Change" heart disease video series in February with a dedicated episode on pediatric cardiomyopathy.
- Pediatric cardiomyopathy and CCF's family programs and services were featured on Transplant Families podcast "Growing the Gift."
- For CCF's Meet the Experts educational webinar, Neha Bansal, MD of Mount Sinai Kravis Children's Hospital presented "Updates in Medical Management for Pediatric Heart Failure."
- Bake for a Cure events were organized in communities across the U.S. and shared on CCF's social media channels.



AWARENESS

CHILDREN'S CARDIOMYOPATHY AWARENESS MONTH

- Enlisted 20 national partners to highlight Children's Cardiomyopathy Awareness Month on their website and social media channels.
- Launched #GoTheDistance campaign on CCF's social media platform to showcase the efforts of CCF families and supporters in raising awareness of pediatric cardiomyopathy.
- Distributed disease facts and resources through email blasts and social media to promote public education and awareness.
- Held a two virtual Heart-to-Heart coffee chat to facilitate discussions about awareness month activities and provide support to newly diagnosed families.
- Raised funds for CCF's family support programs through the 9th Annual Walk for a Cure. Event sponsor, Merck, and dedicated walk teams from New York, Ohio, California, Nevada, Pennsylvania, North Carolina, Maryland, New Jersey supported the event.
- WebMD highlighted their video on pediatric cardiomyopathy, "Medical History Helps Unravel a Mystery," to support Children's Cardiomyopathy Awareness Month



AWARENESS MONTH SPONSORS

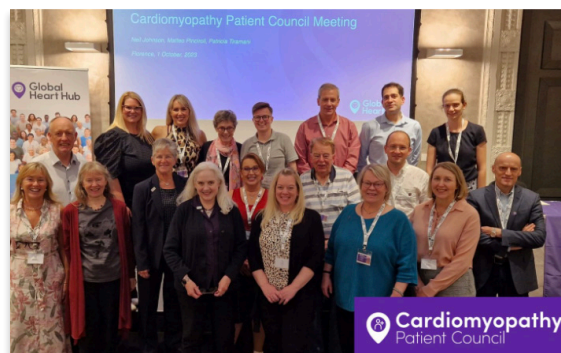


AWARENESS

GLOBAL HEART HUB AWARENESS CAMPAIGNS

As a member of the Heart Failure Patient Council and Cardiomyopathy Patient Council, CCF participated in two multi-country social media campaigns to promote Heart Failure Awareness Month in May and Cardiomyopathy Awareness Week in June.

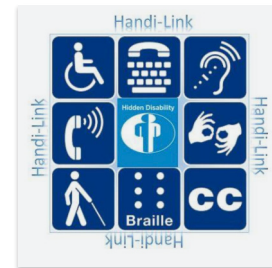
In November CCF attended the first Cardiomyopathy Patient Council meeting in Florence, Italy. 20 patient organizations from 13 countries attended the 3-day meeting to discuss initiatives to improve awareness, diagnosis, and management of cardiomyopathy.



AWARENESS

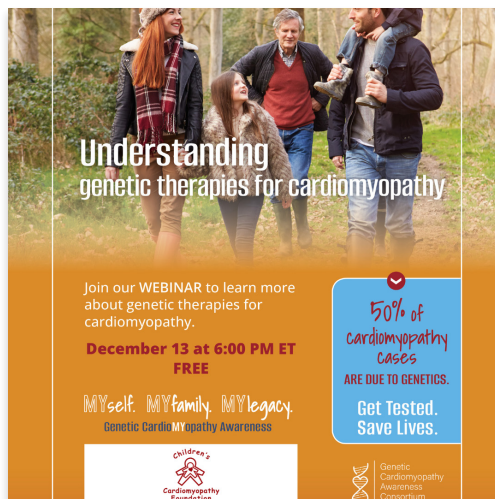
CANADIAN NEWS INTERVIEW

In July, Handi-links, a Canadian radio station focusing on disability issues, spoke to CCF Executive Director Gina Peattie to profile pediatric cardiomyopathy and CCF's services. The station has more than 4,000 followers on X.



GENETIC CARDIOMYOPATHY AWARENESS CONSORTIUM

CCF collaborated with seven cardiomyopathy patient groups to launch the Genetic Cardiomyopathy Awareness Campaign to raise awareness on the need for genetic testing for cardiomyopathy patients and family members. The Genetic Cardiomyopathy Awareness Consortium expanded to 12 patient organizations across 9 countries and launched a “MYself. MY Family. MY Legacy” social media campaign.



Genetic
Cardiomyopathy
Awareness
Consortium

“We stand alongside CCF in the Genetic Cardiomyopathy Awareness Consortium. Through our combined efforts, we aim to address the barriers to genetic testing, ensuring that more patients have access to this crucial diagnostic tool. This collaboration ultimately enhances our ability to connect and advocate for those affected by genetic cardiomyopathy.”

GREG RUFF
EXECUTIVE DIRECTOR
DILATED CARDIOMYOPATHY FOUNDATION

CONNECTIONS



PATIENT &
FAMILY SUPPORT

PATIENT & FAMILY SUPPORT

The Children's Cardiomyopathy Foundation expanded our reach and support of affected families in 92 countries primarily through online connections and virtual meetings.

- CCF has 3,718 active members from 95 countries.
- Grew the CCF Connect Community on Facebook to more than 2,250 members, generating more than 600 posts and 6,500 comments in 2023.
- Assisted more than 7,000 families through CCF's education and support resources.

HEART-TO-HEART FAMILY COFFEE CHATS

Introduced the Heart-to-Heart Coffee Chat as a new platform for caregivers and other family members to have informal discussions about living with pediatric cardiomyopathy. Scheduled after an Experts in Pediatric Cardiomyopathy webinar, it allows CCF members the opportunity to ask follow up questions. Four virtual chats were held in 2023:

- Meet and Chat with the CCF team
- Awareness Month Kick-off
- Support for Newly Diagnosed Families
- Sudden Cardiac Arrest



“CCF has been an amazing resource for my son and the entire family as we have been on the obstructive HCM journey. It’s so nice to have an organization that has experience and an understanding with heart support. We appreciate the connection we have with CCF knowing that they are there for us.”

KAREN WELLMAN
CCF PARENT

PATIENT & FAMILY SUPPORT

CCF HEART KIDS

As part of an ongoing social media campaign, the Children's Cardiomyopathy Foundation profiled 8 children living with cardiomyopathy.



SEDONA



ALEX



AHSHA



BEN



LUKE



GAVIN



OWEN



KELLY ANNE

CONNECTIONS



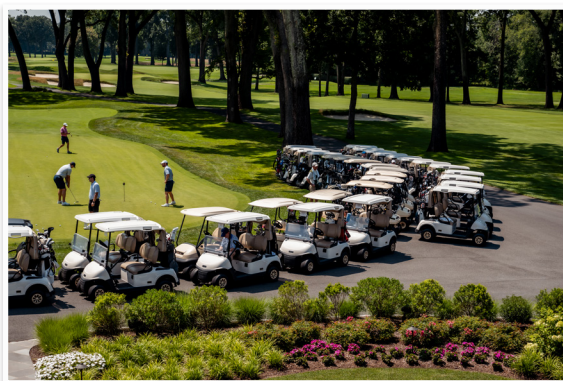
FUNDRAISING

FUNDRAISING

The Children's Cardiomyopathy Foundation's signature fundraising events brought families and supporters together to raise funds for CCF's research, education, and family support programs.

GOLF FOR A CURE

The Children's Cardiomyopathy Foundation (CCF) 19th Annual Golf for a Cure event was held July 24 at the championship Ridgewood Country Club course in Northern New Jersey. In 2023, 150 attendees and 54 sponsors raised \$344,390 for research and education programs on pediatric cardiomyopathy.



BOTTOM LEFT Maureen D'Alleva, Lauren Murphy, Shea Parrish

BOTTOM RIGHT Nick Levine, Rob Steelman, Casey Gaarn, Bobby O'Brien

FUNDRAISING

WALK FOR A CURE

As part of the Children's Cardiomyopathy Awareness Month in September, the 9th Annual Walk for a Cure was promoted in August and September. This email campaign included tips on forming a walk team, setting up an event fundraising page, and utilizing CCF's walk materials and resources to ensure a successful community event walk. The small but mighty group included 84 walkers and volunteers from the National Honor Society which helped to raise \$19,745.



TOP LEFT AND BOTTOM Skinner Family Be kind for Noah Walk for a Cure
TOP RIGHT Owen's Walk for a Cure

FUNDRAISING

BAKE FOR A CURE

Bake for a Cure was promoted in January and February to celebrate National Heart Month. Students across the U.S. organized bake sales to raise community awareness and funding for pediatric cardiomyopathy research.



TOP LEFT Emily Peattie

BOTTOM RIGHT Montclair, NJ Bake for a Cure

FUNDRAISING & FINANCIALS

FAMILY FUNDRAISERS

Families supported CCF by raising \$42,425 in other creative ways, ranging from company/employee gatherings and tricky tray events to social media fundraisers.



“CCF connected me to the correct people and resources when I had questions about my daughter’s cardiomyopathy, as well as supported us with own non-profit, the Ellis Ann Cosby Foundation.”

MEGAN HOPKINS COSBY
CCF PARENT

ABOVE, CLOCKWISE FROM TOP LEFT Rivera Family 4th Annual Butterfly Bee Baking Competition, Kashmala Zaman and MonoSol's LLC Employee Fundraiser, Riley Family Annual Casen's Crew Golf Tournament

CONNECTIONS



FINANCIALS

2023 FINANCIAL SUMMARY

Fiscal year ending December 31, 2023

STATEMENT OF FINANCIAL POSITION

ASSETS

Cash & Cash Equivalents	\$1,542,049
Investments	\$1,074,956
TOTAL ASSETS	\$2,617,005

LIABILITIES + NET ASSETS

Accounts Payable & Accrued Expenses ...	\$12,045
Unrestricted Net Assets	\$2,604,960
Restricted Net Assets	\$0
TOTAL LIABILITIES + NET ASSETS	\$2,617,005

STATEMENT OF ACTIVITIES

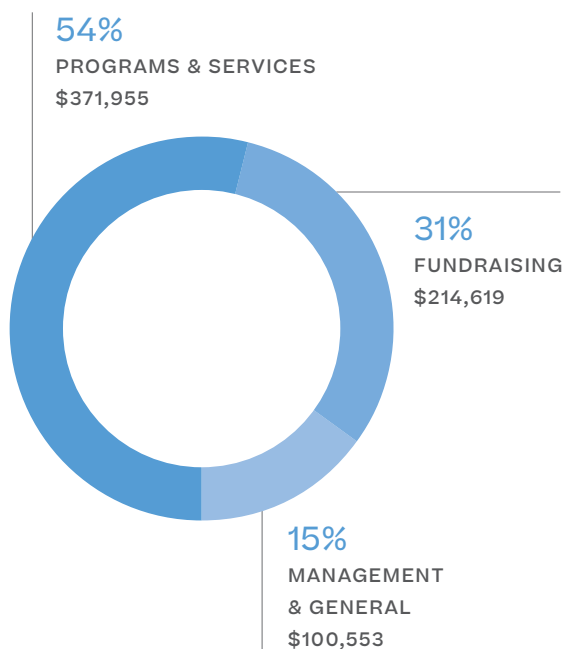
REVENUE

Contributions	\$318,712
Fundraising + Special Events.....	\$378,072
Net Investment Return.....	\$256,094
TOTAL REVENUE	\$952,878

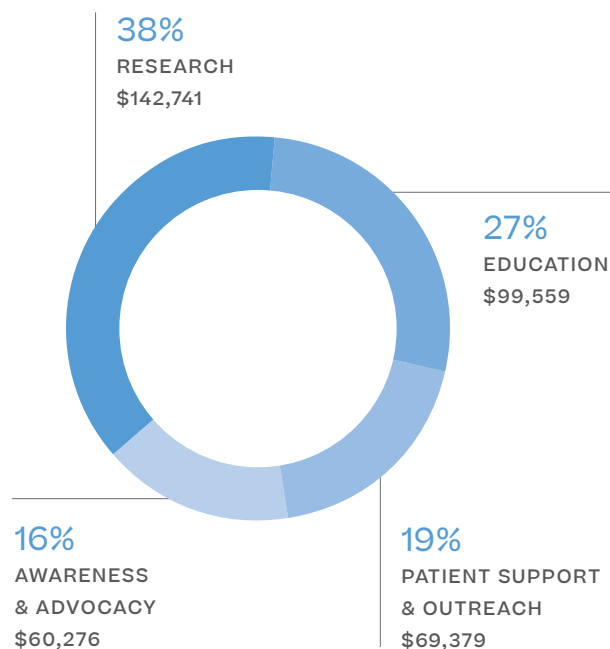
EXPENSES

Programs & Services	\$371,955
Management & General	\$100,553
Fundraising	\$214,619
TOTAL EXPENSES.....	\$687,127
NET INCOME	\$265,751

TOTAL EXPENSES



EXPENSES BY PROGRAM & SERVICES



TOP DONORS

We extend our heartfelt thanks to all who have contributed to the Children's Cardiomyopathy Foundation. The following corporations, foundations, and individuals made contributions of \$500 or more in 2023.

INDUSTRY PARTNERS



DONORS \$10,000 AND OVER

Bloomberg
Bristol Myers Squibb
Casen's Crew
Cytokinetics
Ellis Ann Cosby Foundation
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DONORS \$9,999-\$5,000

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Jonathan Las
Joseph Rossano
Keith B. Hayes Foundation
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Kevin Riggs
Liam Gilmore
Meagher + Geer
Mitchell Salzberg
Philip Schmidt
Polly Huston
Robert Skinner
Sean Monaghan

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Thomas Schneider
Tom & Christine Chun
Victoria Kelly
VMware Foundation
Won Choi

DONORS \$499-\$250

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Elizabeth Poeschl
Gary Stenson
George Khouri
Georgie Vettraino
Ann Vettraino

Jesse Bender
John Pearce
Joo In Kang
Julie Skinner
Karen Aulner
Lisa Tulusaari
Maureen OCallaghan
Michael Ryan
Progressive Insurance Foundation
Raoul Savy
Richard & Susan Thiede Penl
Robert J. Barrett
Sheila Gibbons
Thomas Albertelli
Tracy Erb
TricorBraun
Victoria Bridewell



24 West Railroad avenue, Suite 408, Tenafly, NJ 07670
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