

C.I.T.Y. OF SUPPORT

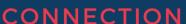
ANNUAL REPORT 2023

MISSION

TO CREATE A DYNAMIC COMMUNITY, BOTH ONLINE AND IN PERSON, THAT SUPPORTS ALL FAMILIES NAVIGATING THE NEEDS OF THEIR CHILD IN THERAPY, REGARDLESS OF DIAGNOSIS.

CORE VALUES







COLLABORATION



CELEBRATION

- WE WELCOME <u>ALL</u> FAMILIES & CHILDREN, <u>REGARDLESS OF DIAGNOSIS</u>. WE UNDERSTAND NOT ALL DISABILITIES ARE VISIBLE, FAMILIAR, OR UNDERSTOOD, AND WE BELIEVE ALL FAMILIES CAN BENEFIT FROM THE SUPPORT OF THIS UNIQUE COMMUNITY.
- RESOURCES & PROGRAMS ARE PROVIDED AT <u>NO COST</u> AND IN VARIOUS FORMATS (<u>IN-PERSON & ONLINE OPTIONS</u>) TO INCREASE ACCESSIBILITY FOR FAMILIES.
- WE STRIVE TO CONNECT FAMILIES TO OUR RESOURCES <u>AS SOON AS POSSIBLE</u>
 BASED ON CURRENT RESEARCH THAT SHOWS THAT CARING FOR CHILDREN WITH
 MEDICAL COMPLEXITY OFTEN TAKES A PHYSICAL, EMOTIONAL, & PSYCHOSOCIAL
 TOLL ON PARENTS AND CAREGIVERS.













MEET SOME OF THE AMAZING CHILDREN IN OUR C.I.T.Y.

C.I.T.Y. CLOSE-UPS

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"My youngest son Silas is the toughest and most resilient kid I know. What was supposed to be a routine C-section ended up with him being born in distress and needing to be transferred to a level 4 NICU. Six weeks later we were discharged with an NG feeding tube and a lot of unanswered questions! I'm so thankful for the C.I.T.Y. Facebook group as it helped me navigate his first year of life. We found out he has not only a pituitary disorder, but also a rare genetic syndrome. He's been in therapy both privately and through Early Intervention. He charms everyone he meets and all the doctors and nurses say he is the best patient!"

"We are proud of our son, Marc, for making many great strides. One, is his communication. Although non-verbal, he has increased the amount of gestures and ability to use his communication device. Secondly, he's now more independent such as taking the school bus by himself. Lastly, no matter how he is feeling, he always has a positive attitude."

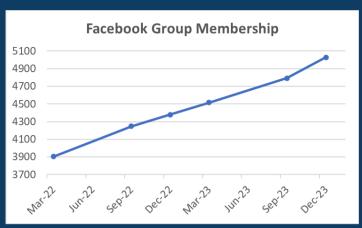


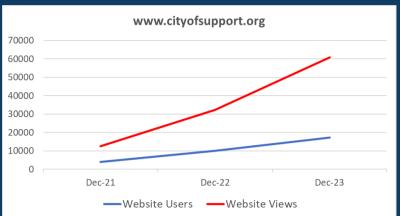


"Theo has a rare genetic condition known as Pelizaeus Merzbacher Disease. Severely affected, we were told Theo would likely not survive his first 2 years. We are extremely proud of Theo for his continued perseverance against staggering odds. He is now 6 years old and has learned to walk with a gait trainer and even ride an adaptive tricycle. Theo is a thrill seeker who loves to ride fast in his wheelchair and be pushed high on the swing. He wins the heart of everyone who meets him with his determination, his adventurous spirit and his signature smile. Though Theo's health is declining, he continues to shine as the brightest star at the center of our universe and we treasure every memory and every smile he shares with us."

CONNECTION

ALL OF OUR PROGRAMS & RESOURCES ARE OFFERED AT NO COST TO FAMILIES, AND MANY ARE AVAILABLE 24/7 ON OUR WEBSITE AND ACTIVE SOCIAL MEDIA CHANNELS.





- VIRTUAL PARENT MEET-UPSDADS HANGOUT GROUPS
- 1ST-EVER PARENT RETREAT
 FAMILY PLAYDATES





7,433

FOLLOWERS IN SOCIAL MEDIA NETWORKS OF ENGAGED PARENTS, PROFESSIONALS, & COMMUNITY MEMBERS





864

PARENTS WHO
CONNECTED IN-PERSON AT
C.I.T.Y. EVENTS THIS YEAR

"The network of families and therapists that CITY brings together is priceless.

Being a parent to a special needs kiddo can be scary, and joyful, and overwhelming. For any of these moments, thanks to CITY, I know right where to go for advice, for expertise, for friendship, and for celebration."

CONNECTION & OUTREACH

CONNECTING C.I.T.Y.'S
NEWEST FAMILIES IN A
COMMUNITY-WIDE EFFORT

"WELCOME TO OUR C.I.T.Y."

This program was developed specifically for families whose newborn is referred for therapy following their NICU stay. Upon discharge from the hospital, parents are immediately connected with a giftbag filled with resources & support from our CITY.





500

BAGS DISTRIBUTED WITH
RESOURCES, WELCOME CARD,
PARENT LETTER, BOARD BOOK &
SMALL GIFTS ASSEMBLED BY
COMMUNITY VOLUNTEERS IN 2023

ONE-OF-A-KIND LEARNING EXPERIENCE WHERE PARENTS TEACH THE THERAPISTS

V.I.P. PROGRAM: "VALUING THE INSIGHT OF PARENTS"

"I wish this training could be mandatory for every Provider and Service Coordinator in our system. Whether you are new to Early Intervention or have been in the field for a long time, this experience will both reaffirm and reshape your practice. The information shared by these families is invaluable."





108

GRADUATES FROM THE V.I.P. PROGRAM THIS YEAR

COLLABORATION

- "WE LOVE OUR C.I.T.Y." MONTH COLLABORATIVE THERAPY TIPS
- ACCESSIBILITY SUPPORT LIBRARY COLLECTION
- BI-ANNUAL PROFESSIONAL NETWORKING EVENTS
- ONLINE CLASSES/WEBINARS & COLLABORATIVE BLOG POSTS





27

VIDEOS OF THERAPY TIPS SHARED BY PROFESSIONALS DURING FEBRUARY 2023 FOR "WE LOVE OUR C.I.T.Y." MONTH





803

NUMBER OF MATERIALS
CHECKED OUT FROM THE
ACCESSIBILITY SUPPORT
COLLECTION AT THE
ARLINGTON HEIGHTS
MEMORIAL LIBRARY THIS YEAR





146

PROFESSIONALS WHO
ATTENDED BI-ANNUAL C.I.T.Y.
NETWORKING EVENTS THIS
YEAR

CELEBRATION

THIS YEAR, 119 GRADUATES REGISTERED TO PARTICIPATE IN OUR ANNUAL, FREE EVENT WHERE ALL ACCOMPLISHMENTS ARE CELEBRATED.

- ANNUAL GRADUATION CELEBRATION TRIBUTE CONCERT/PARENTS NIGHT OUT CITY CLOSE-UPS
 - **BIRTHDAY BUDDIES**















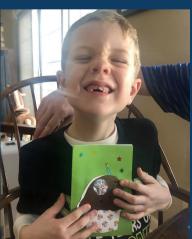
"We've never been to an event like it!"



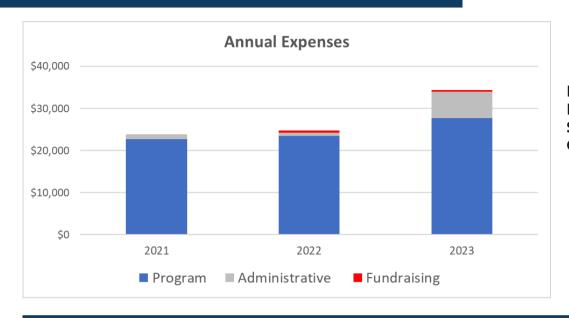








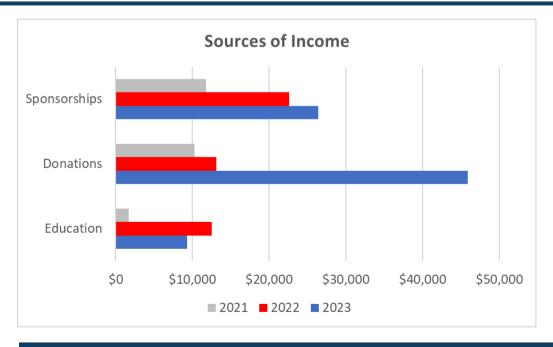
2023 FINANCES



IN 2023, 84% OF EXPENSES WERE USED SPECIFICALLY TO FUND C.I.T.Y. PROGRAMS

C.I.T.Y.'S BOARD CHAIR SPENT <u>965</u> VOLUNTEER HOURS DEVOTED TO THE RUNNING OF THIS ORGANIZATION IN 2023.

THAT'S AN EXTRA 80 HOURS/MONTH (20HOURS/WEEK)
IN ADDITION TO HER WORK AND FAMILY RESPONSIBILITIES



OUR COMMUNITY HAS DEMONSTRATED INCREASINGLY STRONG SUPPORT FOR C.I.T.Y, AS THEY RECOGNIZE THE NEED & IMPACT OF OUR ORGANIZATION

NOTABLY, C.I.T.Y. OF SUPPORT <u>DOUBLED</u> ITS RESERVE FUNDS WITHIN THE FY2023 ALONE.

WHAT'S NEXT FOR C.I.T.Y.

CREATING A PHYSICAL, CONVENIENT, AND INCLUSIVE COMMUNITY CENTER THAT SPECIFICALLY WELCOMES CHILDREN WITH DISABILITIES <u>AND</u> THEIR CAREGIVERS:

THE C.I.T.Y. WELCOME CENTER.





Parents could:

Meet new friends Attend classes together Enjoy quiet time Grab a quick fitness break

Savor a cup of coffee Plan private zoom meetings Answer emails & phone calls Explore a library of resources WITH STRONG PARENT
INTEREST AND FULL
SUPPORT FROM
COMMUNITY
STAKEHOLDERS, THIS
CENTER WOULD BE THE
FIRST OF ITS KIND TO
INCREASE ACCESS TO
MEANINGFUL SUPPORTS
& RESOURCES THAT
MEET THE NEEDS OF
THE ENTIRE FAMILY.



Children could:

Meet new friends Have playgroups with peers Bond with new caregivers

Enroll in family-fun classes Explore different play spaces Enjoy options for all ages OUR NEXT STEP IS TO IDENTIFY & SUSTAIN FUNDING SOURCES THAT CAN SUPPORT SUCH A SPACE WITH ONGOING REVENUE STREAMS, DONORS, & VOLUNTEERS

- "This is a fantastic idea! It's so hard to make adult friends as a parent with our kids. I would love to be able to connect with others that have similar struggles that I could hang out with and foster a friendship."
- "It would be life-changing. Being a parent is lonely, being a parent of a kid who is different is so isolating that it's nearly impossible to socialize. My kids are isolated."
- "I would love something like this. As my daughter's disease progresses I no longer feel that our favorite play places are meant for us. Having somewhere we can go that is specifically for children with disabilities and their families would be so meaningful!"

C.I.T.Y. OF SUPPORT

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BOARD OF DIRECTORS

ALL OF OUR MOTIVATED VOLUNTEERS HAVE A PERSONAL CONNECTION WITH C.I.T.Y, WHETHER IT BE AS A PARENT, PROFESSIONAL, OR COMMUNITY ADVOCATE



BETH DEITER

Founder & Board Chair

Speech-Language Pathologist



JILLIAN BURGARD

Business Owner in Applied
Behavior Analysis



DR. JAMES WEEDON

Developmental & Behavioral

Pediatrician



MARIA
PAPANASTASSIOU
Youth Librarian



SUSAN CHAVEZ
Parent Advocate



JEREMY JOSEPH
Fiduciary Financial Advisor



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www.cityofsupport.org
Contact us at: hello@cityofsupport.org