ANNUAL REPORT 2023

The Chromosome 18 Registry & Research Society

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Rick Guidotti and Positive Exposure

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MEET THE 2023 BOARD

BOARD OFFICERS

President	**Dave Aldrup**
Founder	**Jannine Cody**
Vice President for Member Relations	**Liz Woodfield**
Vice President of Development	**Claudia Traa**
Treasurer	**John Drymala**
Secretary	**Brad Sheppard**

DIRECTORS AT LARGE

Director at Large	**Kristy Holder**
Director at Large	**Ashley George**
Director at Large	**Elizabeth Shaffer**
Director at Large	**Tony Haney**
Director at Large	**Keri Lamont**
Director at Large	**Meredith Moore**

Chromosome 18 Board of Directors, October 2023 *not all members pictured*

For more information about the board members above, click on their names or visit www.chromosome18.org/board-of-directors/

LETTER FROM THE PRESIDENT

The Chromosome 18 Registry & Research Society wrapped up a successful year in 2023 with several achievements to highlight. We enjoyed a wonderful conference in Columbus, highlighted for me by a ‘spirited’ live auction at the Starfish Dinner and Dance. We continued to offer strong programming to bring more attendance to over 50 virtual events. Our network of regional and syndrome coordinators was very active in supporting our membership and new families. We expanded the board to include more members from the adjacent community (not parents), and now have the largest board in our history. Our staff didn’t change and continues to grow in experience, led by CEO Neale Parker with over 7 years of service as well as Director of Operations Amelie Simons with 3 years. And we achieved >15% growth in corporate giving as we continue to seek ways to diversify our funding sources. However, donations from individuals and family fundraising events remain our most significant source for funding our program services. Thank you to all who bring your time, talent, and treasure to support our mission.

Dave Aldrup

PRESIDENT

**pictured far right*

2023 HIGHLIGHTS

4558 CHROMOSOME 18 REGISTRY MEMBERS IN THE U.S.

GRANT FUNDING RECEIVED **$108,000**

24 COUNTRIES REPRESENTED AMONG NEW MEMBERS

THANK YOU VIDEOS SENT TO REGISTRY DONORS AND SUPPORTERS **261**

$659,412 TOTAL FUNDING CHROMOSOME 18 RECEIVED IN 2023

JOINED CHROMOSOME 18 IN 2023*

62	18q- Families
30	18p- Families
22	Trisomy 18 Families
17	Tetrasomy 18p Families
9	Ring 18 Families

*145 Total New Members that joined the Registry in 2023 *(including 3 new members with no syndrome listed)*

Help Chromosome 18 accomplish even more great things in 2024. Use the QR code to show your support!

NEW MEMBERSHIP

2023 New Members by Country

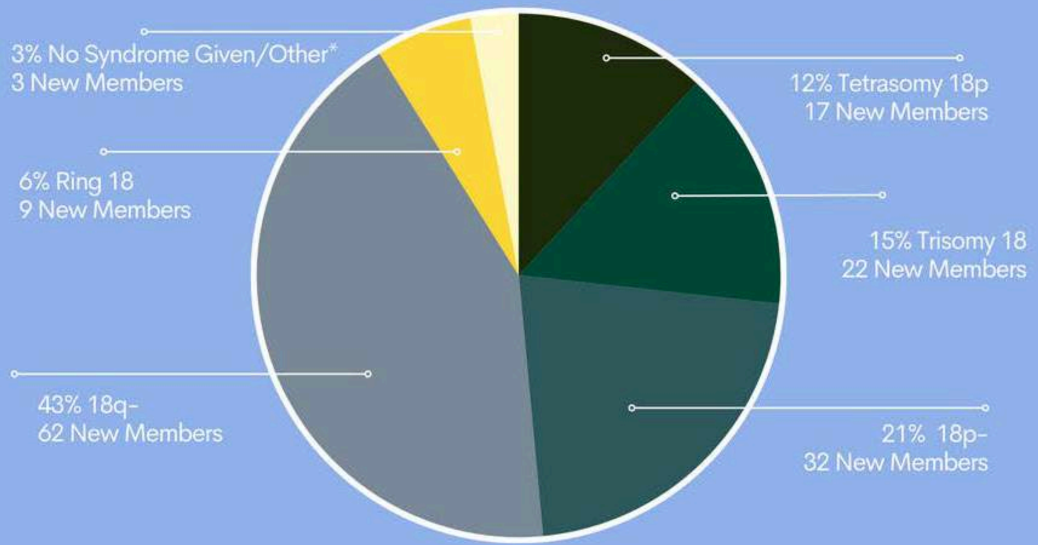
NUMBER OF MEMBERS FROM EACH COUNTRY & PERCENTAGE OF 2023 OVERALL NEW MEMBERSHIP

1. United States - 99 or 67%
2. United Kingdom - 7 or 4.8%
3. Canada - 4 or 2.75%
4. Germany - 4 or 2.75%
5. Brazil - 3 or 2%
6. Chile - 3 or 2%
7. Spain - 3 or 2%
8. Argentina - 2 or 1.4%
9. Belgium - 2 or 1.4%
10. Norway - 2 or 1.4%
11. Philippines - 2 or 1.4%
12. China - 1 or .7%
13. Greece - 1 or .7%
14. Hungary -1 or .7%
15. Israel - 1 or .7%
16. Mexico - 1 or .7%
17. Netherlands - 1 or .7%
18. Oman - 1 or .7%
19. Pakistan - 1 or .7%
20. Portugal - 1 or .7%
21. Romania - 1 or .7%
22. Rwanda - 1 or .7%
23. South Africa - 1 or .7%
24. Switzerland - 1 or .7%
25. Uganda - 1 or .7%

Total international new members in 2023: 46

2023 New Member Syndrome Breakdown

2023: 145 New Members Joined the Registry



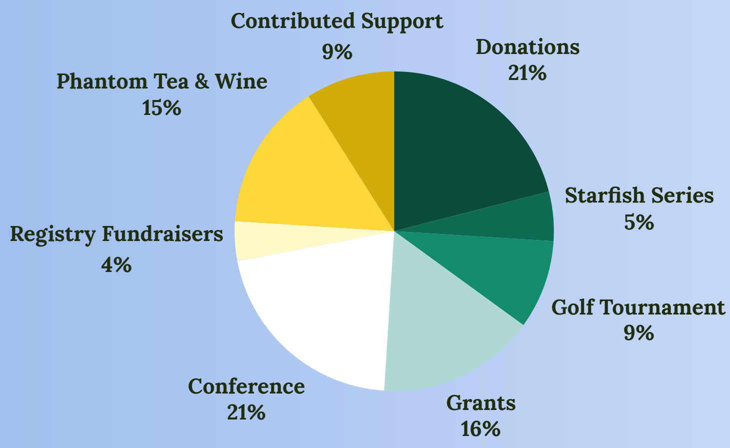
*Listed as volunteer, medical provider or did not list syndrome group

Attention chromosome 18 families!

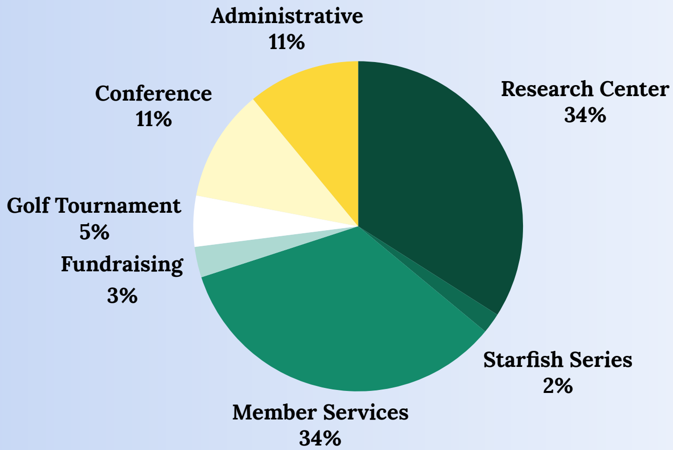
We have quarterly New Member Support virtual meetings and regular parent support groups scheduled for you in 2024. Check out our events page **HERE** to see when the next call is scheduled!

2023 INCOME & EXPENSES At a Glance

2023 INCOME

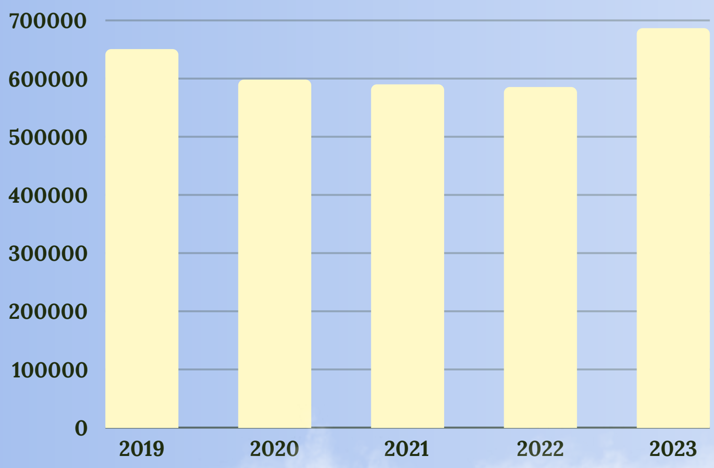


2023 EXPENSES



**Administrative costs include taxes, processing fees, insurance, and donor development costs.*

5-YEAR ANNUAL INCOME OVERVIEW



OVER $686,000 RAISED IN 2023

2023 marked a fantastic financial year for Chromosome 18. While our finances have remained steady since 2020, they have surpassed their 2019 high. We were able to raise over $686,000 with an additional $20,000 in in-kind donations. Thanks to the incredible efforts of our members, grantors, sponsors, and volunteers, income has remained consistent over the past four years and hit the highest revenue yet in over the last 5 years.

IN 2023 OUR MEMBERS RAISED OVER

$71,000

WITH REGIONAL FUNDRAISERS!

Thanks to our amazing members, regional family fundraisers accounted for 10% of our total income for 2023. These regional fundraisers hosted by our members are a vital part in funding research and programming for our families. Anyone can host a regional fundraiser! Just email us to get started at office@chromosome18.org

THE CHROMOSOME 18 CLINICAL RESEARCH CENTER

UT Health
San Antonio
The Chromosome 18
Clinical Research Center

For 33 years, the Chromosome 18 Clinical Research Center (CRC) at UT Health San Antonio has been the only institution in the world dedicated to the study of the chromosome 18 conditions.

2023 saw many changes take place at the CRC. Annice Hill retired after serving as Project Manager for 13 years. Dr. John Gelfond, long-time collaborator, is getting more involved in his new role as Assistant Director with his expertise in biostatistics and data analysis. We also welcomed three new staff members to the team: Catherine McCarthy, Participant Coordinator; Brittany Antopia, Program Data Manager; and Aamuktha Pentala, Research Associate.

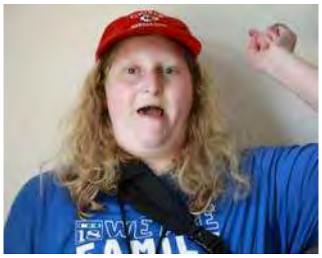
Every day important research is taking place but last year the CRC received a prestigious research grant from the National Institutes of Health (NIH) to enhance the clinical database for the 150 participants who were selected to participate in the Gabriella Miller Kids First Pediatric Research Program (GMFK). The GMFK program provides whole genome sequencing for 150 chromosome 18 research participants with structural birth defects.

The Chromosome 18 Registry & Research Society is the primary funder of the CRC and provided $240,000 in funding in 2023, a $40,000 increase from 2022. With your support, we can continue to increase funding for the CRC and secure vital chromosome 18 research into the future.

Elizabeth & Jannine Cody
SA Light Award Reception
October 2023

732	33	800+	$240,000
Participants enrolled in the research study	Years as the world leader in chromosome 18 research	People found the CRC through Google	Funding received from the Registry in 2023

2023 ANNUAL FAMILY CONFERENCE



The 2023 Chromosome 18 Family Conference was three fun-filled days of learning, community, friendship, and dancing in Colmubus, OH. Conference Co-Chairs Matt & Julie Miller brought us movie nights on the lawn, a meet-and-greet with zoo animals, a dance performance, and a truly memborable summer experience. Conference is the highlight of the year for our families and nothing compares to the experience of being surrounded and supported by the chromosome 18 family.

278 attendees
39 first-time attendees
31 Candian attendees

14 experts
22 educational sessions
1 amazing dance party!

Join us July 7th - 10th in Charlotte, NC for the 2024 Chromosome 18 Family Conference! Registration is open; information available at chromosome18.org.

19TH ANNUAL GOLF TOURNAMENT & STARFISH SOIREE

OCTOBER 5-6, 2023

96

Golfers

$14,500

Funds raised in silent & live auction

$76,500

Total funds raised

2023 was the 19th year for the Chromosome 18 Annual Golf Tournament, as well as the 2nd Annual Starfish Soiree. The Soiree was a magical evening at Oak Hills Country Club with live music, cocktails, hors d'oeuvres, silent auction, live auction, a wine pull, and much more. Thanks to **Soirée Co-Chairs and Board Members Claudia Traa & Lisa Gurwitz** for organizing this incredible pink-themed evening. The weather was perfect the next day for the Golf Tournament and memorial balloon release at the Quarry Golf Course in San Antonio with Valero as Title Sponsor. The golfers had fun playing 18 holes with their family and friends cheering them on. The evening concluded with refreshments and an awards ceremony with a beautiful view of the course.

Click HERE for more information about the special 20th Anniversary of the Chromosome 18 Golf Tournament, scheduled Sept. 27th, 2024 at the Silverhorn Golf Club.

Thank You Jim & Claudia Traa

2024 celebrates the 20th year of the Chromosome 18 Golf Tournament. From the first day, the success of the Tournament has been due in large part to the Tournament Chairs: Jim & Claudia Traa. Their hard work, dedication, engagement, and support over the years have brought this event to the beloved annual celebration it is today. Thank you Jim & Claudia!

The Starfish Dash 5K

Almost 300 runners, walkers, and members of the community came out to support the Starfish Dash 5K in 2023! It was a beautiful spring morning with live music from the 323d Army Band, coffee, donuts, health testing, Sparky & the local Fire Department, cosplayers, and so much more! We loved seeing our TX chromosome 18 families running and walking in person on our 5K course at the UT Health campus while families around the country and across the globe joined the virtual Dash and walked/ran in their hometowns. **Join us for the 7th Annual Starfish Dash on April 27, 2024, either in person or virtually! Click HERE to learn more!**

Power of 18 - The 18-Day Challenge

2023 saw the return of the Power of 18 campaign, but with a new twist! This was the first year we did the 18-Day Challenge as our way to show support and fundraise from Chromosome 18. From September 1st - 18th, participants set their own challenge that incorporated the number 18 - one family did 18 jumping jacks every day for 18 days, while someone else completed 18 acts of kindness in 18 days. The 18-Day Challenge will be back for 2024 and we can't wait to see what amazing feats you're going to accomplish in 18 days!

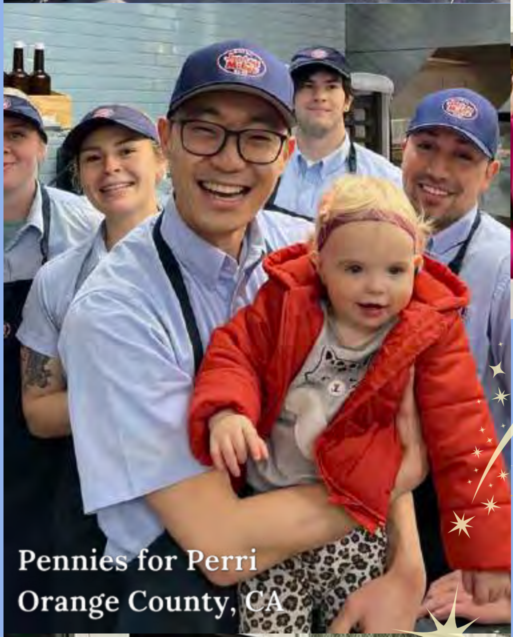
Want to join in the fun and see what virtual events are happening next? Check out our events page HERE.

REGIONAL FAMILY FUNDRAISERS


Southwest Regional Picnic
Yountville, CA


Carter's Cornhole Tournament
Houston, TX


Starfish Strike
Westmont, IL


Pennies for Perri
Orange County, CA


Wheeling Volleyball Team
Wheeling, WV


Starfish Scramble
Richmond, VA


Brayden's Benefit
Wellsboro, PA

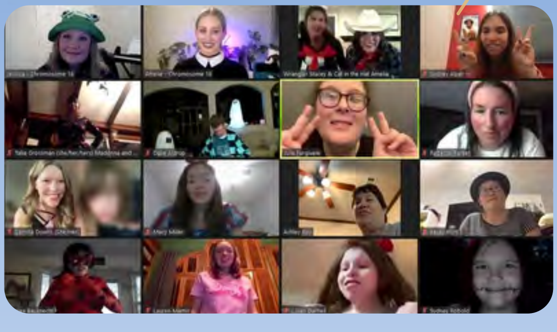

Wine Around the World
Wildwood Crest, NJ

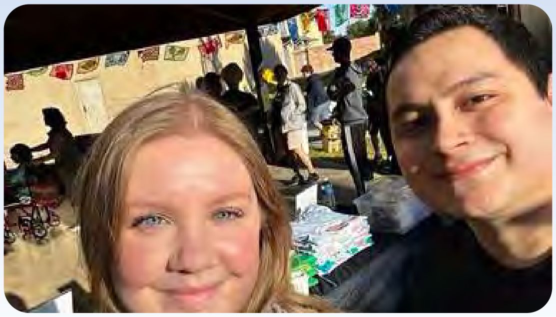

Marvelous Martini Mixer
San Antonio, TX


Danny's Squad
Broomall, PA

Interested in hosting an event in your area? We'd love to help you plan it! Reach out to office@chromosome18.org.

PROGRAM & EVENTS COORDINATOR





I am so excited to be entering my second year with Chromosome 18! Getting to know our families has been absolutely amazing and brings so much joy into my work. Being able to host virtual events, attend my first conference and learn from our regional and syndrome coordinators, makes my job not feel like a job at all.

This year, I am looking forward to Conference in Charlotte, NC and getting to see you all again online and in person.

Thank you for making me feel so welcome and being able to help where I can. The Chromosome 18 community is like no other!

Contact Jessica at ***programs@chromosome18.org*** *for more information about resources!*

-Jessica Sabourin
Program & Events
Coordinator

MEMBER RESOURCES

UT Health
San Antonio
The Chromosome 18
Clinical Research Center

CHROMOSOME 18 CLINICAL RESEARCH CENTER PHYSICIAN MANAGEMENT GUIDES

Released in 2016, the Physician's Management Guides were updated and expanded in 2020. They summarize over 33 years of research and can be shared directly with your physician or specialist. **Find them here.**

SYNDROME & REGIONAL COORDINATORS

As part of Chromosome 18 membership, you have the opportunity to connect with volunteer Regional & Syndrome Coordinators. Find your coordinator **here**.

SOCIAL MEDIA

Chromosome 18 members have access to our closed Facebook pages to connect with other families. Check out the Facebook pages **here**.

View all membership benefits & resources here.

DID YOU KNOW?

Chromosome 18 posted on social media

309

times in 2023! Be sure to follow our FB & IG accounts to stay up to date on our exciting events and developments.

2024 PREVIEW - WEBSITE REDESIGN

We are excited to annouce that our website is undergoing a major redesign, including a brand new "Just Diagnosed" section. The new site will launch spring 2024, stay tuned!

Why Join Chromosome 18?

You're Not Alone, Join Today

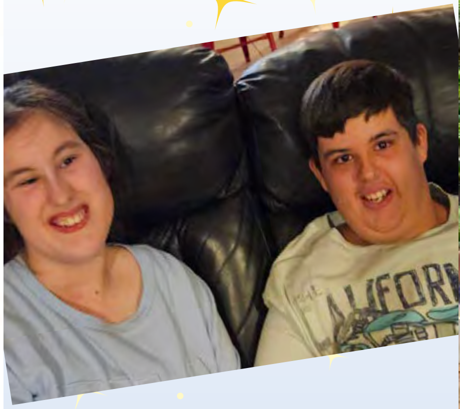
Joining Chromosome 18 is more than just becoming part of a community; it's finding a family that understands, supports, and empowers you. When you join us, you're not alone on this journey. You gain access to a network of individuals and families who have walked similar paths, faced similar challenges, and celebrated similar triumphs. We're here to share knowledge, provide comfort, and inspire hope. Together, we make a difference in the lives of those with chromosome 18 abnormalities. Join us, and let's create a brighter future, one step at a time.

Welcome Video by Dr. Jannine Cody, PhD, Founder

New Parents

Welcome to the Chromosome 18 family! As new parents embarking on this journey, you are not alone. You've joined a compassionate network dedicated to advocacy, support, and research for those affected by chromosome 18 abnormalities. Our Registry is a robust community where you can connect with others through shared experiences, while our Research Center provides cutting-edge medical insights. With educational resources, virtual tours, personalized care guides, and engaging events, we're here to empower you with knowledge and support every step of the way. Welcome aboard, and let's make every moment count together.

CHROMOSOME 18 AUSTRALASIA



In 2023, Chromosome 18 Australasia continued to support individuals and their families to connect and share information. Many of our self-advocates participate in community activities as they endeavour to lead healthy and productive lives. Our committee held several gatherings across the region whilst some of our members preferred to meet at online International events. It was great to see enthusiastic members at all events. We thank all the volunteers who have assisted with organising activities and are extremely grateful to those who are able to support our ongoing fundraising activities.
The committee is looking forward to 2024 and encourages families to join in and connect with other Chromosome 18 families at our activities.

Visit Chromosome 18 Australasia's Facebook page for more information.

Do you live in Europe? Visit Chromosome 18 Europe's website for more information.

CHROMOSOME 18 CANADA

In 2023, Canadian C18 families celebrated and said a big thank you to Shelley Hunt. Shelley was the main face of our Canadian group for over 25 years and we are so grateful for all of the support she has given everyone, from social events to C18 knowledge to butter tarts! In February, the Haney family hosted a wine pairing fundraiser to kick off the year! It was a great time with delicious food and wine for a great cause! To kick off summer, we gathered at Jeff and Natalie Banton's for a BBQ with our U.S. friends, the Parkers. It was a great evening of food, friends and badminton! In August, we had our annual Ontario BBQ in Oakville where we had over 8 families gather. We are so lucky to have a group of families that are so close that we can watch our kids grow together. In November, Susan Douglas arranged a C18 moms getaway in Prince Edward County. There is nothing better than unwinding and enjoying the scenery with your second family! To close out the year we hosted the Canada hour on World Chromosome 18 Day, and had a great time connecting with families from across the country.

In 2025, we look forward to hosting the conference in July – get your passports ready!

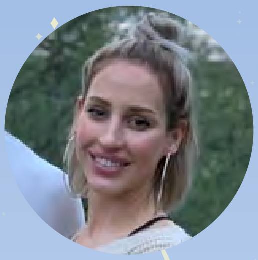
Visit **Chromosome 18 Canada's Facebook page** to stay up to date on future events and for more information

CHROMOSOME 18 OFFICE

MEET THE TEAM

Chromosome 18 Staff


Neale Parker
CEO


Amelie Simons
Director of Operations


Jessica Sanders
Program & Events
Coordinator


David Garza
Accounts Supervisor

2023 Work-Study Students & Interns


Danielle Arreguin
Marketing &
Grants Assistant

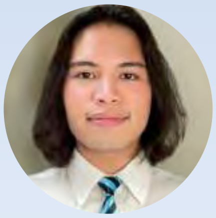

Gabriella Cancino
IT/Web Development
Assistant


Kiran Esani
Conference Special
Projects Assistant


Yonnell Esprit
Accounts Assistant


Karina Fernandez
Social Media &
Marketing Assitant


Alejandra Gerlach
Grants Assistant


Johnny Martinez
Grants Assistant


Adriana Pena
Grants Assistant


Daniel Ramirez
IT/Web Development
Assistant


Isabella Villena
Development Assistant


Jerry Stendebach
Accounts Assistant

CHROMOSOME 18 WORK-STUDY & INTERNSHIP PROGRAMS

Chromosome 18 partners with numerous high schools, colleges, and City of San Antonio programs to create internship and work-study roles within our organization. These roles help us expand our organizational capacity while giving young adults in San Antonio the opportunity to learn about the operations of a “small but mighty” nonprofit organization. Through these programs, we host young adults from our city as work-study students and interns and provide them with real-world, hands-on experience in the nonprofit sector. We hope to both inspire students to work or volunteer for a nonprofit after graduation, as well as encourage them to remain in San Antonio and invest in our growing community. We want to thank all our wonderful interns and work-study students who helped us accomplish great things in 2023!

Click here to learn more about the student experience at Chromosome 18.

CHROMOSOME 18 VOLUNTEER DIRECTORS

Jeff Borello
Director of Technology

Stacey Gallardo
Director of Social Media

Gene Johnson
Director of Data

Val Shepard
Director of Grants

We are incredibly grateful to our amazing Volunteer Directors who donate their time and expertise to support Chromosome 18. In 2023, our Directors helped us with grants, social media, our member database management, and updating our website. They are an invaluable resource and bring their skills to help us raise awareness and reach more new families every year.

WELCOME THE NEW CLINICAL RESEARCH CENTER STAFF

This year, the Chromosome 18 Clinical Research Center (CRC) welcomed three new staff members: (from left to right) Aamuktha Pentala, Brittany Antopia, and Catherine McCarthy. Cathrine is the new Participant Coordinator and will be the first point of contact for families at the CRC. Brittany is the Program Data Manager who will oversee the management of the research study data, and Aamuktha Pentala is the Research Associate. Welcome to the team!

2024 MAJOR EVENTS

April 27, 2024 - The Starfish Dash 5K Run/Walk & Health Fair at UT Health San Antonio

July 7-10, 2024 - The 29th Annual Chromosome 18 Family Conference in Charlotte, NC

September 6-8 - TX Starfish Camp at Morgan's Wonderland Camp (more details to come) in San Antonio, TX

September 27, 2024 - The 20th Annual Chromosome 18 Golf Tournament, Silverhorn Golf Club, San Antonio, TX

December 1, 2024 - World Chromosome 18 Day (virtual and free!)

December 2024 - Phantom Tea & Phantom Wine and End Of Year Giving

Check out our event page on our website here for our monthly virtual events, as well as various other Chromosome 18 fundraisers happening around the country in 2024!

2023 LEADERSHIP TEAM

BOARD OFFICERS

President	Dave Aldrup
Founder	Jannine Cody
Vice President for Member Relations	Liz Woodfield
Vice President for Development	Claudia Traa
Treasurer	John Drymala
Secretary	Brad Sheppard

DIRECTORS AT LARGE

Director at Large	Tony Haney
Director at Large	Meredith Moore
Director at Large	Brad Jensen Sheppard
Director at Large	Kristy Holder
Director at Large	Keri Lamont
Director at Large	Ashley George
Director at Large	Elizabeth Shaffer

REGIONAL COORDINATORS

Great Plains	Anama Family
Northeast	Rebecca Teague
Rocky Mountain	Kevin and Jennifer Borges
Southeast	Allison & Jeremy Linman
Texas Area	Open
Great Lakes	Carrie Connolly
Mid-Atlantic	Mike Fudala
Northwest	Open
South Central	Bea Powell
Southwest	Jessica Robold

SYNDROME COORDINATORS

Tetrasomy 18p	Cristi Cain
18q-	Nichole McVicker
Trisomy 18	Jude Wolpert
Ring 18	Carmen Tyce
18p-	Camilla Downs

ADDITIONAL COORDINATORS

Sibling Coordinator	Kendall Powell
Bereavement Coordinator	Camille Hammond
Self-Advocate Coordinator	Open

EX-OFFICIO MEMBERS

Europe	Bonnie McKerracher
Australia	Tracey Healy
Canada	Shelley Hunt
Africa	Christine Mutena
Brazil	Dalton Vilela
Mexico	Nancy Chavez

Not sure what region you're in? Click here for more information!

THANK YOU TO OUR DONORS!

$10,000 and Up:

Liz Woodfield and Bill Carlucci
Cowden Charitable Foundation
Jannine Cody
Neale Parker and Cynthia Parsons
Gordon Hartman Family Foundation
Mays Family Foundation
Peg and Ben Flowe
Valero Energy Foundation
Kristy Holder
Botello Lumber Company
Grand Fitness Mgmt, LLC
Jersey Mike's Subs
Dan Shea and Stephanie Stokes

$5,000 to $10,000:

Boys of 88
Dave and Tam Aldrup
Jeff, Kathy, Hannah Borello and Borello Family
Elizabeth Cody
Jeff and Carol Connor Cohen
S & L Marx Foundation
Michael and Cierra Fudala
Stacey Gallardo
Dr. Daniel Hale
St. Luke's Lutheran Health Ministries
Jesse and Jan Poggio
Brad and Debra Sheppard
Peter Sullivan and Snow Family Trust
Blue Ivy Partners - Tim Ogden
Claudia and Jim Traa
Judith and William Wolpert

$2,500 to $5,000:

Heartland Bank
Chuck and Loren Campbell
The Larry and Juana Carter Charitable Fund
Fiesta San Antonio Commission
Paul, Laura, and Morgan DeMars
Christine and Jon DeMars-Victorsen
GlaxoSmithKline
Tony and Valerie Haney
Robbie Mendez
Jacqueline and Alexander Mo
Meredith and Kent Moore
Special People in Need Foundation
Estee and Brian Nelson
Dean and Jennifer Oltmanns
Jessica and Ryan Robold
Tom and Angie Schilly
Elizabeth Shaffer
Kelly and John Tremonte
UT Health
Jessica Verhulst-Biocic
Walmart
Kathleen Wessman

$1,000 to $2,500:

Alycia Alongi
Asel & Associates
Natalie Banton
Cooper Cares
Scott Carlson
Samuel Carmack
Bailey and Liam Carmack
Paul and Kerry Castella
New Moon Charitable Fund
Keith and Helena Christiansen
Thomas Codd
Antoinette Cox
Allison Dale
John Drymala
Turner Family
Armanino Foundation
Jay and Susie Goldberg
Max and Maggie Grant
Barbara Grossman
Liz and Seth Grossman
Kevin Hamblin
Amanda Healey
H-E-B
Jodi Hight and Hight Real Estate
Cheryl Hilgenbrink
Eleanor Claire Hilliard
Jared Kennedy
Sean Kiehne
Patricia Lacey
Nicole and Todd Leslie
Mary Lollis
Scott Lollis
Wayne and Deborah MacMasters
Clean Scapes, LLC - Marcy Nichols
Doug and Julie Masterson
Farm Credit Mid-America
Denise and Tom Parker
Mary and Mark Placey
Puttie Potts
Suzanne Powers
Christine Puckett
Kathryn Ranselben
Brian and Wendy Rathjen
Kayvan Romaezi
David and Valerie Shepard
Alamo Sign Solutions
Lorna Stevenson
Joseph and Theresa Verstreater
Sergio and Alice Viroslav
Chad and Patricia Watson
Brian and Jennifer Wessman
Joel and Candice Wolf

$500 to $1,000:

Jennifer Adams
Lockport Animal Hospital
Zakary Bates
Nancy and Brent Beason
Susan Bentivenga
Arlene Biocic
Andrew and Rachel Bond
Matthew Bouzakis
Joe Brennan
Tim Brunn
Jennifer Bushnell
Cristi Cain
Bryan Cook
Jere Corven
Wendy R. Crawford Fund
Katrina Delaney
Carol Ann and Wayne Edwards
Amber Ennis
Bettina Finn
Raul and Sandy Flores
Melissa Garza
Andrew Gipe
Cathy Gorrell
Lisa Hand
Minire Hasi-Zogaj
Greg and Juliette Hitchan
Donald and Carla Hughes
Laurie and Gene Johnson
Erica and John Keating
Ben Keefer
Nathaniel Kelly
David Lewis and Jack Kelly
River and Amber Koblick
Keri and James Lamont
Karen Lane
Shauna Lapoint
Brian Leivo
Mr. and Mrs. Robert Leonard
Susan Leone
John and Nina Lewis
Eric Miles
Matt and Julie Miller
Andrew and Shannon Mueck
Sheila Newcomb
Kevin O'Kelley
Eagle Print
Rosemarie Quinlan
Tracy Rivest-Keeney
Sandy Rocka
Michael Rush
Karma Sawyer
Christopher and Kristin Schneider
Christopher Scott
Jaron Sheppard
TopNotch Silkscreening
Eli Skarsten
Deborah Stevens
Sheila Stone
Lewis and Dee Noonberg - The Noonberg Family Fund
Yvonne Tritto Lisella
Wheeling University Volleyball Team
Steven and Kim Welborn
Lewis and Susan Wiens
Carol Zaletsky

Thank You

For the overwhelming support you showed us in 2023. It is because of our donors with their direct gifts and families with their tireless fundraising efforts, that we were able to accomplish so many great things this year. We are so grateful to you! Please note, this list reflects only direct donations from individuals or corporations. We endeavor to ensure that all donors are recognized for their support. However, if you notice your name is missing or another discrepancy in our list, please contact our office at **office@chromosome18.org**.

Chromosome 18 Champions

THANK YOU TO OUR MONTHLY DONORS

David Aldrup
Zakary Bates
Jon Bennett
Andrew Bond
Jannine Cody
Kristen Earl
Timothy Egedus
Michelle Faidengold
Michael Fudala
Kathleen Glascock
Dalton Guthrie
Nathaniel Kelly
River & Amber Koblick
Patricia Lacey
Meredith Moore
Elizabeth Shaffer
David Shepard
Michael & Karen Todd

Monthly recurring donations are consistent gifts that help Chromosome 18 plan for long-term success. This list only reflects individuals who give directly to Chromosome 18. We endeavor to ensure that everyone is recognized for their support. However, if you notice that your name is missing from the list, please contact our office at **office@chromosome18.org** so we may thank you properly!

Legacy Society

THANK YOU TO OUR FOUNDING MEMBERS

Jannine Cody
Jeff & Kathy Borello
Terri Lutton
Charles Wirick
Mike Fudala
Dan & Pat Schmick
Rosemary Woodfield
Lorie Stevenson

The benefits of a planned gift to Chromosome 18 are immeasurable by enabling you to support Chromosome 18 and strengthen our mission. Being a member of the Legacy Society gives you the opportunity to belong to a select group of like-minded people who are committed to support the needs of our families.

Interested in joining Chromosome 18's Legacy Society? Learn more here.

Workplace Giving Champions

THANK YOU TO THOSE WHO CONTRIBUTE TO CHROMOSOME 18 THROUGH EMPLOYER MATCHING PROGRAMS

Alycia Alongi
Ashley Engelbach
Eileen Finlay
Raul & Sandy Flores
Nishant Gandhi
Amanda Healey
Joe Higgins
John Hoekstra
Laurie & Gene Johnson
Peter Kniesel
Brian Leivo
Nina Lewis
Heather Maher
Robbie Mendez
Margaret Odem
Dean & Jennifer Oltmanns
Christine Puckett
Mackenzie Reed
Sandy Rocka
Thomas Schilly
Christopher Schneider
Deborah Stevens
Daniel Thiel
Yvonne Tritto-Lisella
Joseph Verstreater
Joel Wolf
Liz Woodfield
Carolyn Zaletsky

Workplace matching gifts are a great way to double your impact when you give to Chromosome 18. Interested in workplace giving? Ask your employer if they participate or have a matching donation program!

This list only reflects individuals who give directly through their employers to Chromosome 18. We endeavor to ensure that everyone is recognized for their support. However, if you notice that your name is missing from the list, please contact our office at **office@chromosome18.org** so we may thank you properly!

THANK YOU FOR A WONDERFUL YEAR!

2023