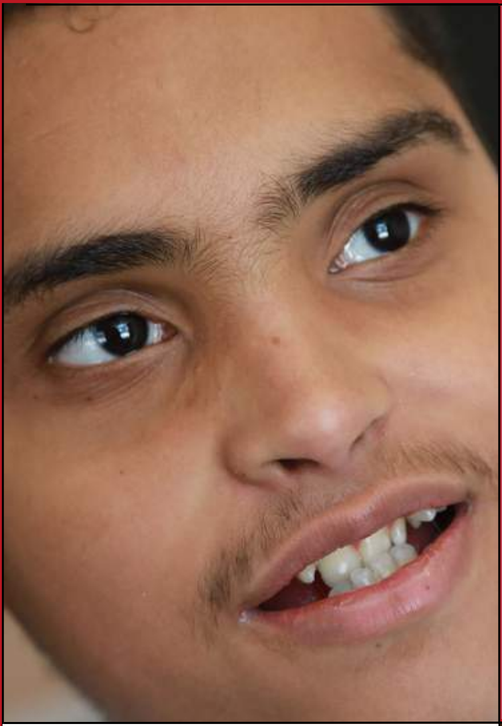

THE
CHROMOSOME 18
REGISTRY &
RESEARCH
SOCIETY

2022 ANNUAL REPORT

The Chromosome 18
Registry & Research Society

Photo credit- cover and page 2:
Rick Guidotti and Positive Exposure

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MEET THE 2022 BOARD

BOARD OFFICERS

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

DIRECTORS AT LARGE

Director at Large	**Carol Connor Cohen**
Director at Large	**Tony Haney**
Director at Large	**Kristy Holder**
Director at Large	**Keri Lamont**
Director at Large	**Meredith Moore**
Director at Large	**Brad Jensen Sheppard**

Chromosome 18 Board of Directors, October 2022 **not all members pictured**

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

A MESSAGE FROM THE FORMER AND CURRENT PRESIDENTS

The goal of an annual report is to highlight accomplishments and to thank the many people who facilitated those accomplishments; be it through donating their precious time or by providing monetary support or both. This year's report is particularly special because it is about more than just the last year's accomplishments. This report marks an important milestone as the Chromosome 18 Registry completes the transition from the kitchen table to the board room. The final step in the transition is thanks to an amazing staff who do the day-to-day work of providing the services so important to our membership. This includes, Neale Parker, CEO; Amelie Simons, Director of Operations; Holly Hollingsworth succeeded by Jessica Sanders, Program and Events Coordinator and Victoria Ochoa succeeded by David Garza, Accounts Supervisor. These talented and dedicated individuals with the help of numerous volunteers and work-study staff create the solid foundation for all that goes into making the organization function. Their competence allows the Board of Directors to not be involved in the day-to-day running of the organization but to focus on those things that a Board of Directors should be focused on, strategic planning, networking, and fundraising. I am excited to report that we have such a board! And I am ecstatic to report that 2023 begins with the transition to a new board President, Dave Aldrup. This is such an important moment for the organization and completes the evolution from the kitchen table to the board room.

SINCERELY,
JANNINE CODY,
FOUNDER & BOARD PRESIDENT 1990-2022

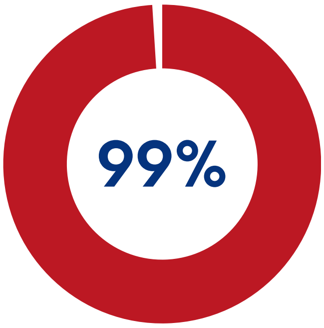
Looking back on 2022, among the many people who helped move the goals of the Chromosome 18 Registry forward, I would like to highlight the significant contributions over the years from our Founder, Dr. Jannine Cody. Even as Jannine transitions out of the role of Board President, she continues to advise the board and lead the clinical research team. The accolades continue to pour in for her, and we all owe her a huge debt of gratitude. To many, we may look like a small organization. But if your child has a chromosome 18 condition, we are the lifeline: to an embracing community, to guidance for survival, to hope for a better future. To these families we are not small; to them we are essential. And for them, we have a tremendous responsibility. A responsibility which could not be met without the considerable contributions of time, talent, and treasure from many supporters who value generosity and hope.

SINCERELY,
DAVE ALDRUP, BOARD PRESIDENT 2023

! *Curious about where Chromosome 18 is headed in the future? Check out the* ***2022-2024 Strategic Plan*** ***HERE.***

2022 MEMBER SURVEY

In our 2022 all-member survey...



Respondents reported that Chromosome 18 is doing important and meaningful work.

Respondents report that research is important for improving quality of life for those with chromosome 18 conditions

Respondents report the Chromosome 18 website is a useful resource.

Respondents report their life has been affected in a positive way by the Registry.

2022 HIGHLIGHTS

JOINED CHROMOSOME 18 IN 2022*

60

18q-
Families

35

18p-
Families

27

Trisomy 18
Families

21

Tetrasomy 18p
Families

10

Ring 18
Families

**Does not include 37 families who selected "other" as a syndrome*

190

NEW MEMBERS

190 NEW MEMBERS JOINED THE REGISTRY, OUR SECOND-HIGHEST YEAR EVER AS WE WORK TO REACH MORE NEW FAMILIES FOLLOWING DIAGNOSIS

OVER $114,000 CONTRIBUTED FROM OUR MEMBERS IN END-OF-YEAR GIVING

$114K

30

COUNTRIES

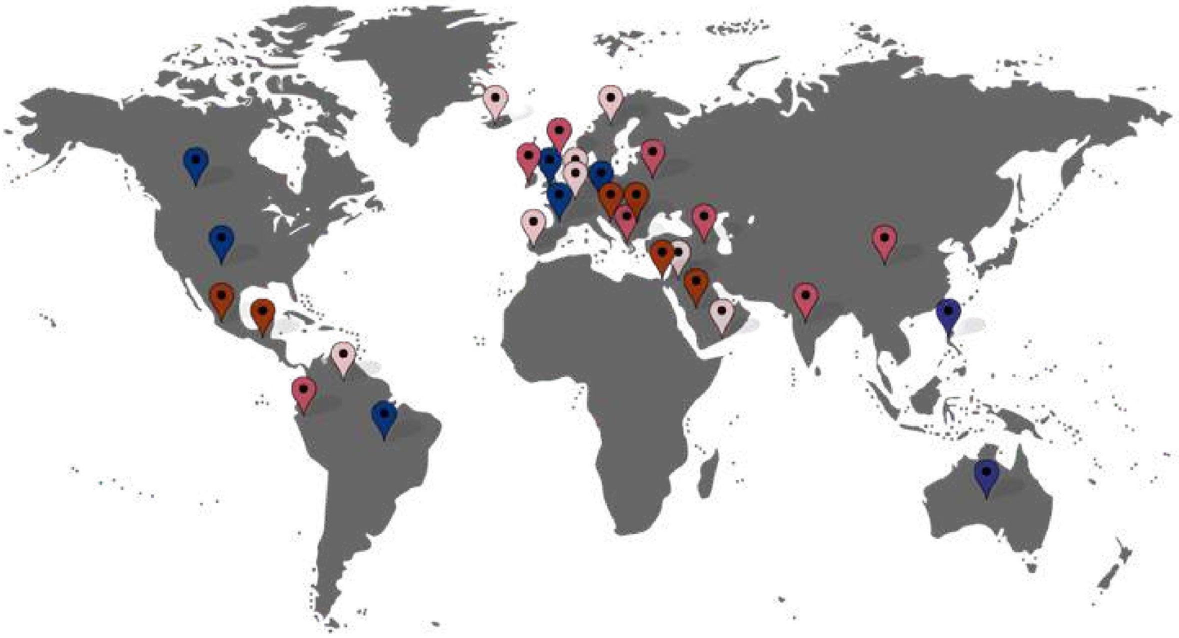
REPRESENTED AMONG THE NEW MEMBERS THAT JOINED THE REGISTRY IN 2022. INTERNATIONAL FAMILIES WERE 36% OF NEW MEMBERS IN 2022.

OVER $36,000 RAISED BY FAMILIES WHO HOSTED REGIONAL FUNDRAISERS

$36K

NEW MEMBERSHIP

2022 New Members by Country



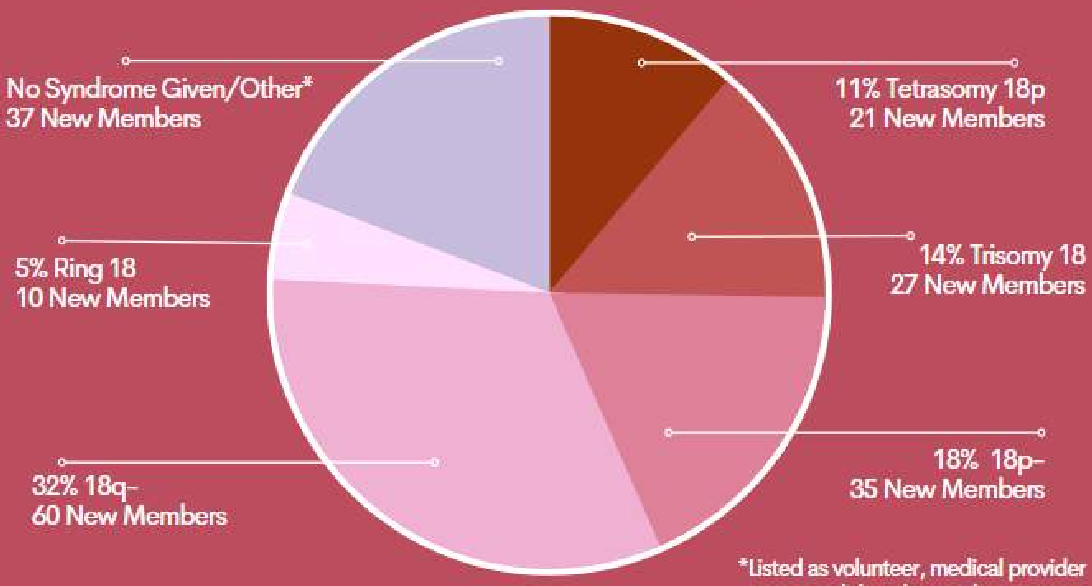
NUMBER OF MEMBERS FROM EACH COUNTRY & PERCENTAGE OF 2022 OVERALL NEW MEMBERSHIP

1. United States - 124 or 65%
2. United Kingdom (in total) - 12 or 6%
3. Australia - 7 or 4%
4. Canada - 6 or 4%
5. Brazil - 6 or 4%
6. France - 6 or 4%
7. Phillipines - 4 or 2%
8. Germany - 3 or 1.5%
9. India - 2 or 1%
10. Ireland - 2 or 1%
11. Scotland - 2 or 1%
12. Poland - 1 or .5%
13. Greece - 1 or .5%
14. Ecuador - 1 or .5%
15. China - 1 or .5%
16. Azerbaijan - 1 or .5%
17. Iceland - 1 or .5%
18. Jordan - 1 or .5%
19. Portugal - 1 or .5%
20. UAE - 1 or .5%
21. Belgium - 1 or .5%
22. Venezuela - 1 or .5%
23. Netherlands - 1 or .5%
24. Sweden - 1 or .5%
25. Saudi Arabia - 1 or .5%
26. Israel - 1 or .5%
27. Guatemala - 1 or .5%
28. North Macedonia - 1 or .5%
29. Mexico - 1 or .5%
30. Austria - 1 or .5%

Total international new members in 2022: 62

2022 New Member Syndrome Breakdown

2022: 190 New Members Joined the Registry



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

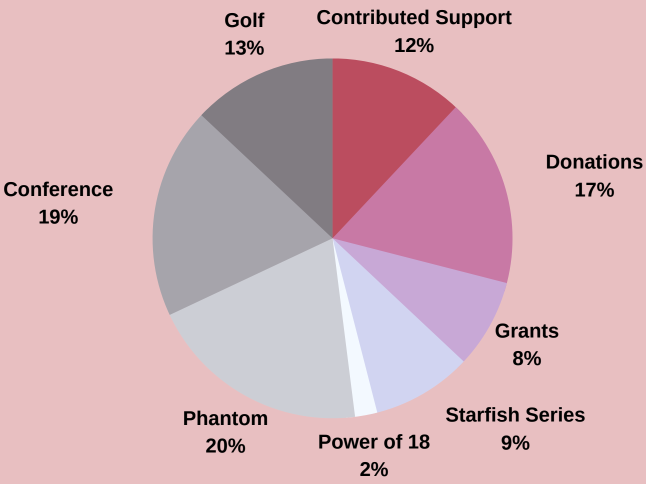
Attention chromosome 18 families:

We have quarterly New Member Support virtual meetings and regular Sydrome Support groups scheduled for you in 2023. Check out our events page ***HERE*** *to see when the next call is scheduled!*

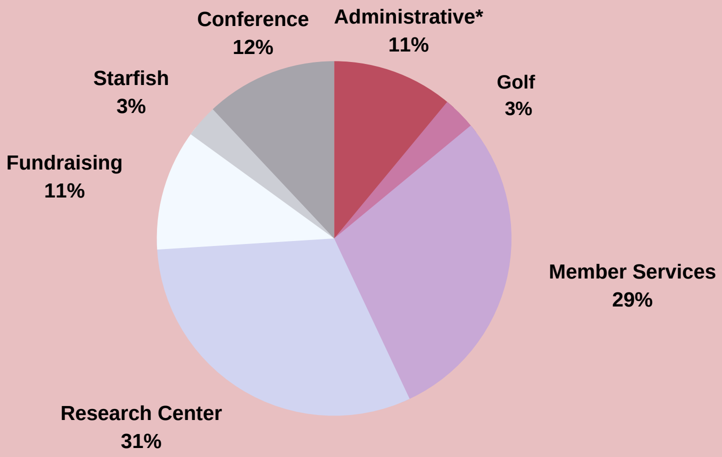
CHROMOSOME 18

2022 INCOME AND EXPENSES AT A GLANCE

2022 INCOME

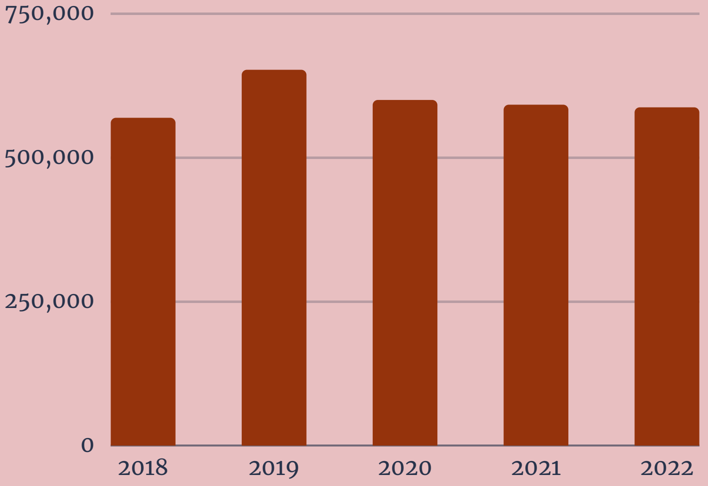


2022 EXPENSES



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

5-YEAR ANNUAL INCOME OVERVIEW



OVER $585,000 RAISED IN 2022

2022 marked another challenging financial year for Chromosome 18 following the global pandemic. While our finances have remained steady since 2020, they have yet to return to their 2019 high. Nevertheless, we were still able to raise over $585,000 in 2022. Thanks to the help of our members, grantors, sponsors, and volunteers, income has remained consistent over the past three years and we were able to overcome one of the biggest global challenges in modern times togther.

In 2022 our members raised over

$30,000

with Facebook Fundraisers!

Thanks to our amazing members, Facebook fundraising accounted for 5% of our total income for 2022. These individual fundraisers, especially birthday fundraisers, are a vital part in funding research and programming for our families. Facebook fundraisers are easy, anyone can do it! Just email us to get started at **office@chromosome18.org**

27TH ANNUAL CHROMOSOME 18 DECODED FAMILY CONFERENCE

JULY 9-12, 2022 - San Antonio, TX

27TH ANNUAL FAMILY CONFERENCE
CHROMOSOME 18 DECODED
SAN ANTONIO, TX ★ JULY 7-9, 2022

239

Attendees

45

Educational sessions

45

First-time attendees

23

Medical Experts

1

Unforgettable Dance Party!

2022 saw the return of our beloved Chromosome 18 Decoded Annual Family Conference in San Antonio, TX. It was a magical three days of community, education, and fun! The Chromosome 18 family came together to enjoy some incredible learning and time together. Some of the highlights included:

- Mariachi band at the welcome reception
- Special workshops and training sessions, like Love & Logic
- Syndrome breakout groups
- Parent, sibling, and self-advocate panels
- Sibling scavenger hunt and dinner out
- Self-Advocate night out at Dave & Buster's
- Ladies' line dancing night out
- Men's beer and BBQ night out
- Cosplayers and princesses at the Starfish Dinner & Dance
- And so much more!

2022 Conference Chairs
Dave & Tam Aldrup

*This year's conference (2023) has a new name, but will be the same amazing experience: **Camp Chromosome 18 - Annual Family Summit**. Join us **July 9th - 12th in Coumbus, OH** for an experience you won't want to miss! Click **HERE** for more information and to register, we can't wait to see you there!*

THE CHROMOSOME 18 CLINICAL RESEARCH CENTER

For 32 years, the Chromosome 18 Clinical Research Center (CRC) has pioneered chromosome 18 research as the only dedicated research center in the world for these conditions. Because of the CRC, San Antonio is the epicenter of the chromosome 18 community and stands as beacon of hope for anyone with a chromosome 18 condition. 2022 saw many exciting developments for the CRC. Dr. John Gelfond, long-time collaborator, joined the CRC as Assistant Director and brings his expertise in biostatistics and data analysis to research on the chromosome 18 conditions. The CRC also continued work in key areas like studying long-term survivors of Trisomy 18, and the Kids First Program in which 150 study participants with structural birth defects were selected for whole genome sequencing. Thanks to generous grant funding from community partner Valero, the CRC was able to purchase a new NanoDrop One device (pictured top right above) that researchers use to accurately measure DNA concentrations from the blood samples donated by chromosome 18 families. Finally, with the return of the Chromosome 18 Decoded Annual Family Conference in San Antonio, the CRC was able to resume in-person enrollment and testing for the longitudinal study.

The Chromosome 18 Registry & Research Society is the primary funder of the CRC and provided $200,000 in funding in 2022. Unfortunately, funding has not yet been able to recover to pre-pandemic levels. However, with your support, we can invest in essential chromosome 18 research by increasing CRC funding. Increasing funding for the research would mean both preserving the important discoveries and expanding research projects into the future.

UT Health
San Antonio
The Chromosome 18
Clinical Research Center

709

Participants enrolled in the research study

60

Peer-reviewed articles published

32

Years as the world leader in chromosome 18 research

22

Research updates and educaitonal videos for families

19

Expert collaborators from various disciplines

Congratulations Dr. Cody!

In October 2022, Dr. Cody was celebrated as a winner of the prestigious Morgan's Wonderland Hearts of Gold Wall of Fame award. The award recognizes individuals who have dedicated their lives to those with disabilities and special needs. Dr. Cody was honored for her over three decades of work to improve the lives of those living with chromosome 18 conditions. Thank you to Morgan's Wonderland for this special honor, and thank you Dr. Cody for a lifetime of serving children, individuals, and families affected by chromosome 18 conditions.

Photo credit: Morgan's Wonderland

> *"The contributions you have made to science and medicine are significant."*
> -Dr. William Henrich, President of UT Health, October 2022

GOLF TOURNAMENT 18TH ANNIVERSARY

OCTOBER 6-7, 2022

In 2022 we celebrated the 18th anniversary of the Chromosome 18 Golf Tournament in a very special way! The evening before the tournament, golfers, community partners, sponsors, and supporters enjoyed a beautiful evening at Oak Hills Country Club for the Starfish Soirée. Thanks to **Soirée Co-Chairs Claudia Traa & Lisa Gurwitz**, the celebration included specialty drinks, a silent and live auction, and wine pull surrounded by starfish-themed décor.

The next day, the 18th Annual Golf Tournament started at the Quarry Golf Course with a presentation by **Tournament Co-Chairs Claudia & Jim Traa** and a memorial balloon release. The golfers enjoyed perfect weather on the renowned course cheered on by chromosome 18 families and self-advocates from the sidelines. The evening concluded with refreshments and an awards ceremony.

$73,000 total raised!

over $7500 in silent & live auction donations

100 golfers

Click here for more information about the 19th Anniversary of the Chromosome 18 Golf Tournament, scheduled October 6th, 2023 at the Quarry Golf Course.

Thank You Valero Energy!

A very special thank you to the title sponsor of the 18th Anniversary of the Annual Golf Tournament - Valero Energy. Valero has been an amazing supporter of the Chromosome 18 mission for over 25 years and has donated $550,000+ to help our families. John Drymala, Chromosome 18 Treasurer and Valero employee, has been instrumental in raising funds to support our organization and has dedicated countless hours over the years to helping our operations and events run smoothly. Thank you John for all your hard work and support!

The Starfish Dash 5K

200 runners, walkers, and members of the community came out to support the return of the Starfish Dash 5K and Fiesta in San Antonio, TX! Chromosome 18 families in Texas joined in the active morning, live music, and a health & wellness fair. We also had numerous families from across the country and around the world participate from afar in the virtual Starfish Dash. Don't miss out on joining us at the 6th Annual Starfish Dash on April 29, 2023, either in person or virtually! Click HERE to learn more!

Virtual Events

2022 was packed with virtual events, some returning favorites and other brand new ways to have fun together from afar. We enjoyed a pet show and tell, virtual game night, boys' and girls' hang outs, a Halloween dance party, and storytime with Santa. We also started Syndrome Support Groups for chromosome 18 parents and caregivers, which are returning for 2023!

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

REGIONAL FUNDRAISING HIGHLIGHTS

STARFISH SCRAMBLE EVENT SERIES

STARFISH SCRAMBLE NAPERVILLE

In early May 2022, The Borello Family hosted a Starfish Scramble at Topgolf in Naperville, IL. With 40 people in attendance and some amazing silent auction items and sponsors including Andromeda Technology Solutions and Pete's Natural Market, the Scramble raised nearly $7000. Check out the Facebook album **here**.

STARFISH SCRAMBLE HOUSTON

In late May 2022, the Aldrup Family held a Starfish Scramble in Houston, TX. Thanks to the 30 attendees and family sponsors including the Ashley, Shively, Larson and Gardner families, the event raised over $2500. View photos on Facebook **here**.

STARFISH SCRAMBLE ATLANTA

In June, the Linman Family hosted a Starfish Scramble at Topgolf in Atlanta, GA. They had 12 wonderful family sponsorships and 30 people came out to golf and support the event. Together, the Linmans and their friends and supporters raised $3,000. View photos **here**.

STARFISH SCRAMBLE FT. WORTH

In October 20, 2022, The Hight Family hosted a Starfish Scramble at Topgolf in Ft. Worth, TX. The Scramble saw 34 participants and raised $5000 for chromosome 18 research. View the photos in the Facebook album **here.**

STARFISH SCRAMBLE TUCSON

In November, 2022, the Robold Family organized and hosted the Tucson Starfish Scramble in Arizona. With over 60 participants, sponsors like the Kennedy, Schafer and Knox families, and in celebration of Jessica Robold's birthday, the Tucson Scramble raised $7500. View photos **here.**

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

PROGRAM SERVICE COORDINATOR

From Holly Hood, Program Service Coordinator 2021-2022:

It has been wonderful getting to know you all virtually and meeting some of you at conference! You and your families are amazing, and it has been a complete joy to work with each one of you! Thank you for welcoming me as part of the Chromosome 18 family.

The community here is unlike any other organization I've worked for. I look forward to volunteering at the golf tournament, conference, and other opportunities to stay involved, so I will still see ya'll around.

Thank you again for welcoming me into the Chromosome 18 family.

Jessica Sanders
Current Program & Events Coordinator

I joined the Chromosome 18 team in December 2022 and I am excited to grow as the programs and events coordinator. I moved to San Antonio in 2022 and have enjoyed getting to know the area and the Chromosome 18 community.

We are looking forward to Syndrome Support Groups and self-advocate events resuming to give more connection to our families. We hope to continue to provide engaging, educational and community events for our members. This year, I hope to grow our Blog and Family stories so other families can relate and feel less alone in their journeys. We hope to continue to grow as a organization.

Being able to reach out to families and learn about their lives, has allowed me to see how impactful the registry can be to new families and families who have been with the registry for years.

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

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 30 years of research and can be shared directly with your physician or specialist. **Find them here.**

SOCIAL MEDIA

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

ONLINE RESOURCE PORTAL

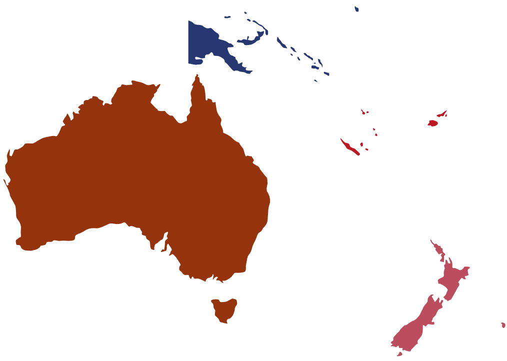
Chromosome 18 has consolidated member resources into one central location. Take a few minutes to review our resources **here**.

*View all membership benefits & resources **here.***

2022 Blog and Family Highlights

> ***Stay up to date with all things Chromosome 18, meet our new families, and hear stories from other parents in our monthly newsletter and blog. Click HERE to check out our blog, and email office@chromosome18.org to be added to our mailing list.***

CHROMOSOME 18 AUSTRALASIA



Chromosome 18 Australasia supports people with a chromosome 18 condition and their families to connect and share information to help overcome the obstacles they face. We encourage families to join us at regional events and online. Many of our self-advocates participate in community activities as they endeavour to lead healthy and productive lives.

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

CHROMOSOME 18 CANADA

Canadian Chromosome 18 held its first Family Reunion in July 2022, which brought together over 25 families. A great time was had by all in Niagara Falls, Ontario! The Chromosome 18 Canadian committee put together a fun family reunion that included treasure hunts, water park days, crafts, games, and tons of family time for all participants (we were even visited by Santa Claus himself).

We are already planning the next one and hope to start including educational tracks as the reunion grows in the future.

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

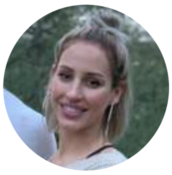
Visit **Chromosome 18 Canada's Facebook page** to stay up to date on future events and for more information

CHROMOSOME 18 OFFICE

MEET THE TEAM

Chromosome 18 Staff


Neale Parker
CEO


Amelie Simons
Director of Operations


Holly Hood
Program Service
Coordinator


Jessica Sanders
Program Service
Coordinator


Victoria Ochoa
Accounts Manager


David Garza
Accounts Supervisor


Tracy Gibbons
Development Manager

Grants Students


Chasta Callies
Grants Assistant


Sophie Gifford
Grants Assistant


Thomas Livesay
Grants Assistant


Alejandra Gerlach
Grants Assistant


Adriana Pena
Grants Assistant


Cheyenne Wilson
Grants Assistant

Accounting Student


Selena Garza
Accounts Assistant

Social Media & Marketing Students


Maria Gonzalez
Graphics & Video
Production Assistant


Maria Vazquez
Graphics & Video
Production Assistant


Karina Fernandez
Social Media &
Marketing

Health Special Projects Students


Jordan Bentivenga
Administrative Assistant


Emily Haase
Conference Special
Projects Assistant


Becky Li
Special Projects Assistant

IT & Web Development Students


Daniel Ramirez
IT/Web Development
Assistant


Christian Kennedy
IT/Web Development
Assistant


Delon Perry
IT/Web Development
Assistant

Community & Development Students


Tessa Uviedo
Community Outreach
Assistant


Larissa Gonzales
Development
Assistant

CHROMOSOME 18 HOSTS INTERNS AND WORK-STUDY STUDENTS FROM CITY PROGRAMS AND LOCAL UNIVERSITIES

Chromosome 18 is incredibly proud of the multiple partnerships we have with local universities and city work readiness programs for high school and college students. Through these programs, we host young adults from San Antonio as work-study students and interns and provide them with real-world, hands-on experience in the nonprofit sector. Our work-study students and interns are essential to the work we do at the Registry and help us in so many areas including grants, development, social media and marketing, and IT and web design. In 2022, Chromosome 18 hosted 18 work-study students and was recognized as a standout host and mentoring organization by SA Worx, a city program that provides high school students internships with local organizations. We want to thank all our amazing interns and work-study student who helped us accomplish so much in 2022, particularly the return of our in-person events. We couldn't do it without you! 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

Alex Lima
Director of Marketing

Riddhi Masukar
Director of Member Relations

Val Shepard
Director of Grants

Our Volunteer Directors donated a huge amount of their time and expertise to support Chromosome 18 in 2022. They are vital to what we do at the Registry and are key to helping us better serve our members. Thank you, wonderful Directors, for your vital wealth of knowledge and your incredible efforts!

2023 MAJOR EVENTS

April 29th, 2023 - The Starfish Dash 5K Run/Walk & Health Fair at UT Health San Antonio

July 9-12, 2023 - The 28th Annual Camp Chromosome 18 - Annual Family Summit, Columbus, OH

October 6, 2023 - The 19th Annual Chromosome 18 Golf Tournament, The Quarry Golf Course, San Antonio, TX

December 3, 2023 - Global Chromosome 18 Day (virtual!)

December 2023- Phantom Tea & Phantom Wine

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 2023!

Are you planning an event to support Chromosome 18 by holding your own fundraiser?
Let us know by emailing us at office@chromosome18.org

2022 LEADERSHIP TEAM

BOARD OFFICERS

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

DIRECTORS AT LARGE

Director at Large	Carol Connor Cohen
Director at Large	Tony Haney
Director at Large	Meredith Moore
Director at Large	Brad Jensen
Director at Large	Sheppard
Director at Large	Kristy Holder
Director at Large	Dan Pioch
Director at Large	Keri Lamont

REGIONAL COORDINATORS

Great Plains	Anama Family
Northeast	Jason Goodman
Rocky Mountain	Shaunna Keller
Southeast	Allison & Jeremy Linman
Texas Area	Angela Ashley
Great Lakes	Carrie Connolly
Mid-Atlantic	Mike Fudala
Northwest	Joy Wilson
South Central	Bea Powell
Southwest	Jessica Robold

SYNDROME COORDINATORS

Tetrasomy 18p	Cristi Cain
18q-	Nichole McVicker
Trisomy 18	Jude Wolpert
Ring 18	Shaunna Keller
18p-	Camilla Downs

ADDITIONAL COORDINATORS

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

EX-OFFICIO MEMBERS

Europe	Bonnie McKerracher
Australia	Tracy Healy
Canada	Shelley Hunt
Africa	Christine Mutena
Brazil	Dalton Vilela

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

THANK YOU TO OUR DONORS!

$10,000 and Up:
The Harvey E. Najim Charitable Foundation
Elizabeth Cody
Jannine Cody
Peg & Ben Flowe Jr.
Valero Energy Foundation
The Gordon Hartman Family Foundation
Daniel & Patricia Schmick
Dan Shea & Stephanie Stokes

$5,000 to $10,000:
Boys of 88
David & Tambra Aldrup
Kathy & Jeff Borello
Jeff & Carol Connor Cohen
Marilyn DeMars
S & L Marx Foundation
Mays Family Foundation
Michael & Cierra Fudala
Stacey Gallardo
Daniel Hale M.D.
Mike & Erica Johnson
Meredith & Kent Moore
Cynthia Parsons & Neale Parker
Brad & Debra Sheppard
Claudia & Jim Traa
Snow Family Trust
Cowden Charitable Foundation Trust
William & Judith Wolpert
William Carlucci & Liz Woodfield

$2,500 to $5,000:
Katie Bailey
Larry & Juana Carter Charitable Fund
GlaxoSmithKline Foundation
Sundt Foundation
Annice Hill
Kristy & Jason Holder
Duncan Mann
Special People in Need Foundation
Brian & Estee Nelson
Blue Ivy Partners
UT Health Science Center San Antonio
United Way of San Antonio & Bexar County
Joel & Candice Wolf

$1,000 to $2,500:
Alycia Alongi
H-E-B
Scott Carlson
Keith & Helena Christiansen
Fiesta San Antonio Commission
Bryan Cook
Antoinette Cox
Lindy DeGrave
John Drymala
Kristen & Jeffrey Earl
Banton Family
Blackwell Family
Wessman Family
Bettina Finn
Jordan Ford
Jay & Susie Goldberg
Barbara Grossman
Dalton Guthrie
William & Susan Guthrie
Tony & Valerie Haney
Amanda Healey
Jennifer & Thomas Heggie Kunkel
Middlesex United Way Inc.
The Horton Group Inc.
Brian & Shaunna Keller
Sean Kiehne
Patricia Lacey
Judith LaFrombois
Catherine & Ryan Larson
Mary & Scott Lollis
Doug & Julie Masterson
Patricia & John McGing
Jacqueline Ngo
Samantha Norlin
Stephen Oddie
Asel & Associates, PLLC
Puttie Potts
Christine Puckett
Dennis Quinn
Kathryn Ranselben
Brian & Wendy Rathjen
Jessica & Ryan Robold
Tom & Angie Schilly
Deborah Stevens
Lisa Tortolano Mazur

$1,000 to $2,500:
Kelly Ann & John Tremonte
Jeffrey & Brianna Turner
Joseph & Theresa Verstreater
Sullivan County VFW
Chad & Patricia Watson
Carolyn Zaletsky

$500 to $1,000:
Patricia Bailey
Zakary Bates
Andrew & Rachel Bond
James Borello & Co.
LeNora Bredsguard
Heather & David Brogan-Myers
Paul & Kerry Castella
Carrie Colella
Cary & Cindy Collins
Kelly Collins
Carrie Connolly
Rennee Cozzolino
George & Traci Dunham
Carol & Wayne Edwards
Amber Ennis
Carmack Family
Davis Family
The Noonberg Family Fund
Daniel Flanigan
Raul & Sandy Flores
Emily Gosh
Maximilian Grant
Minire Hasi-Zogaj
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Interested in joining Chromosome 18's Legacy Society? Learn more **here**.

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THANK YOU FOR A WONDERFUL YEAR!

2022


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