THE CHROMOSOME 18 REGISTRY & RESEARCH SOCIETY

2021

ANNUAL REPORT

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MEET THE 2021 BOARD

BOARD OFFICERS

President & Founder	**Jannine Cody**
Vice President for Member Relations	**Liz Woodfield**
Vice President for Development	**Kathy Glascock**
Treasurer	**John Drymala**
Secretary	**Dave Aldrup**

DIRECTORS AT LARGE

Director at Large	**Carol Connor Cohen**
Director at Large	**Tom Kunkle**
Director at Large	**Meredith Moore**
Director at Large	**Brad Jensen Sheppard**

Chromosome 18 Board of Directors & Staff, August 2021

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

A MESSAGE FROM THE CHROMOSOME 18 FOUNDER & PRESIDENT

In this Annual Report, we have the opportunity to thank the many people who help move the goals of the chromosome 18 families forward. 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, as an organization, have an awesome responsibility. A responsibility which could not be met without the generous commitments of many many friends who value generosity and hope and who know they "helped that one;" as so well illustrated in the Starfish story from the book Chicken Soup for the Soul.

The Starfish story continues to inspire us and is a theme for our many events. Sadly, with 2021 being yet another pandemic year, we were again unable to hold in-person Starfish events. That that did not stop us, however, from expanding our virtual outreach work. While we were saddened to not be able to share hugs with our friends, we were heartened to be able to meet so many new families from across the globe using virtual platforms. In adversity lies opportunity, and we used this opportunity to find new ways to reach out to people.

This Annual Report highlights the new and expanded programs for the record number of new members in 2021. It also shares the challenges of supporting that work when fundraising events can't happen. But, most importantly, we wish to share a public thank you to the many individuals, foundations, and corporations whose generosity enabled continued support to our very special families. We all work together to make lives better for people with chromosome 18 abnormalities. Thank you all!

SINCERELY,
DR. JANNINE CODY

2021 IN REVIEW

THE CHROMOSOME 18 REGISTRY & RESEARCH SOCIETY

JOINED CHROMOSOME 18 IN 2021*

77

18q- Families

26

18p- Families

24

Trisomy 18 Families

27

Tetrasomy 18p Families

10

Ring 18 Families

*DOES NOT INCLUDE 22 FAMILIES WHO SELECTED "OTHER" AS A SYNDROME TYPE.

200

NEW MEMBERS

CHROMOSOME 18 SAW A RECORD HIGH OF 200 NEW MEMBERS JOIN THE REGISTRY THIS YEAR AS WE WERE ABLE TO REACH MORE NEW FAMILIES THAN EVER BEFORE!

OVER 6,700 HOURS OF SUPPORT FROM OVER 30 WORK-STUDY AND VOLUNTEER INTERN STUDENTS IN 2021

6,700

HOURS

14

MEDICAL PROFESSIONALS JOINED THE REGISTRY TO LEARN MORE ABOUT HOW TO BETTER CARE FOR THEIR PATIENTS

CHROMOSOME 18 WAS AWARDED OVER $59,000 IN GRANT FUNDING IN 2021. THIS FUNDING IS THE COMBINATION OF SEVERAL LOCAL AND NATIONAL GRANTS OF VARIOUS SIZES.

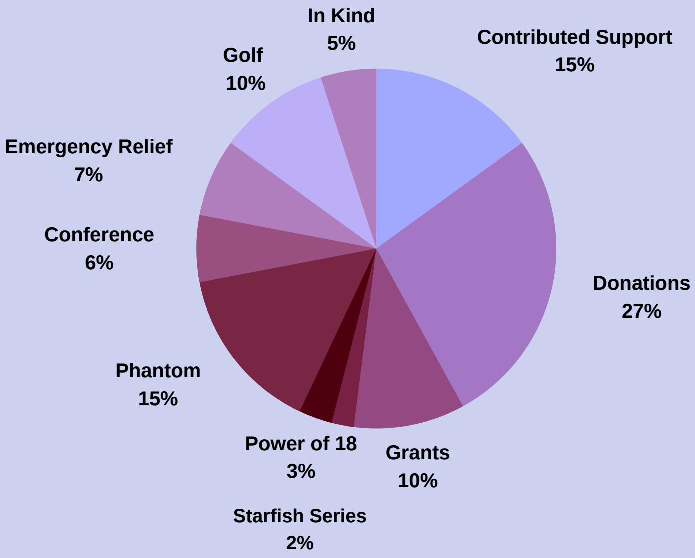
$59K

GRANT FUNDING RECEIVED

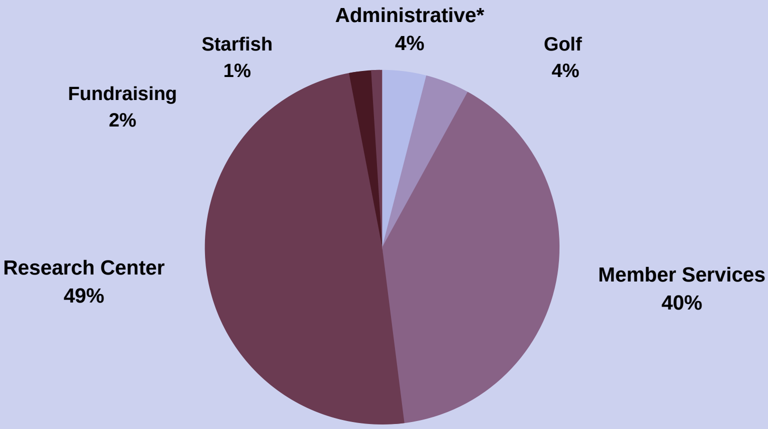
CHROMOSOME 18

2021 INCOME AND EXPENSES AT A GLANCE

2021 INCOME



2021 EXPENSES



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

5-YEAR ANNUAL INCOME OVERVIEW

OVER $590,000 RAISED IN 2021

2021 marked another challenging financial year for Chromosome 18 in light of the global pandemic. While our finances have yet to return to their pre-pandemic high, we were still able to raise over $590,000 despite severe limitations to our in-person fundraising events. Thanks to the help of our members, grantors, sponsors, and volunteers, income has remained consistent and Chromosome 18 was able to persist through two unprecedented years.

Did You Know?

In 2021 we raised almost

$50,000

Through Facebook Fundraisers!

Facebook fundraising accounted for almost 10% of our total income for 2021. These fundraisers are vital to our efforts to support chromosome 18 research and put on amazing in-person and virtual events for our families. Interested in fundraising on Facebook but don't know where to start? Just email us to get started at **office@chromosome18.org**

COVID-19 IMPACT & Virtual Events

OUR CONTINUED RESPONSE & RESOURCES

CLINICAL RESEARCH CENTER VIRTUAL TOURS

The staff from the Chromosome 18 Clinical Research Center provided 5 virtual tours of the facilities throughout 2021. Dr. Cody took us on a tour of the offices, introduced the staff & volunteers, and lead us through the lab where the research is done. View one of the virtual tours here.

FACEBOOK LIVE EVENTS

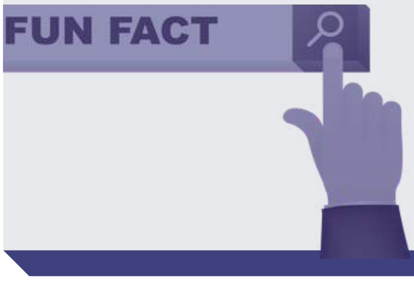
Our Facebook Lives were a big hit! We had almost 4,000 views on our Facebook Live videos in 2021. We highlighted Self-Advocates' hobbies and job experiences, as well as Research Updates, Fundraising Opportunities, Virtual Conference, and more!

SELF-ADVOCATE FEATURES

In 2021, we highlighted several Self-Advocates. David Aldrup Jr. talked about his job at the movie theater. Meagan Rush shared her experience working in childcare and with self-employment. Matthew Peter Bates discussed his job and his triathlon training. Cierra Fudala told us about her job at Bake Ability and her love for working out.

COVID 19 VACCINE UPDATES

In December 2021, Paul Niolet, MD, an Allergy & Immunology Specialist, as well as a parent of a child affected with 18p-, shared another update on COVID-19 Vaccines & individuals affected by chromosome 18 conditions. View the video here.



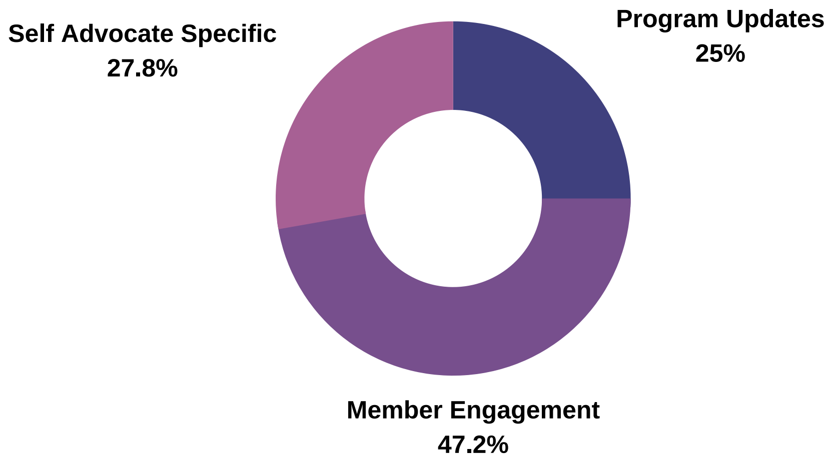
YOUTUBE HIGHLIGHT

In 2021, Dr. Cody created a series of 10 FAQ videos to answer some of the most common questions about the chromosome 18 conditions. These videos were distributed in the monthly Research Center newsletter, and are available here to view anytime!

Member Engagement Events

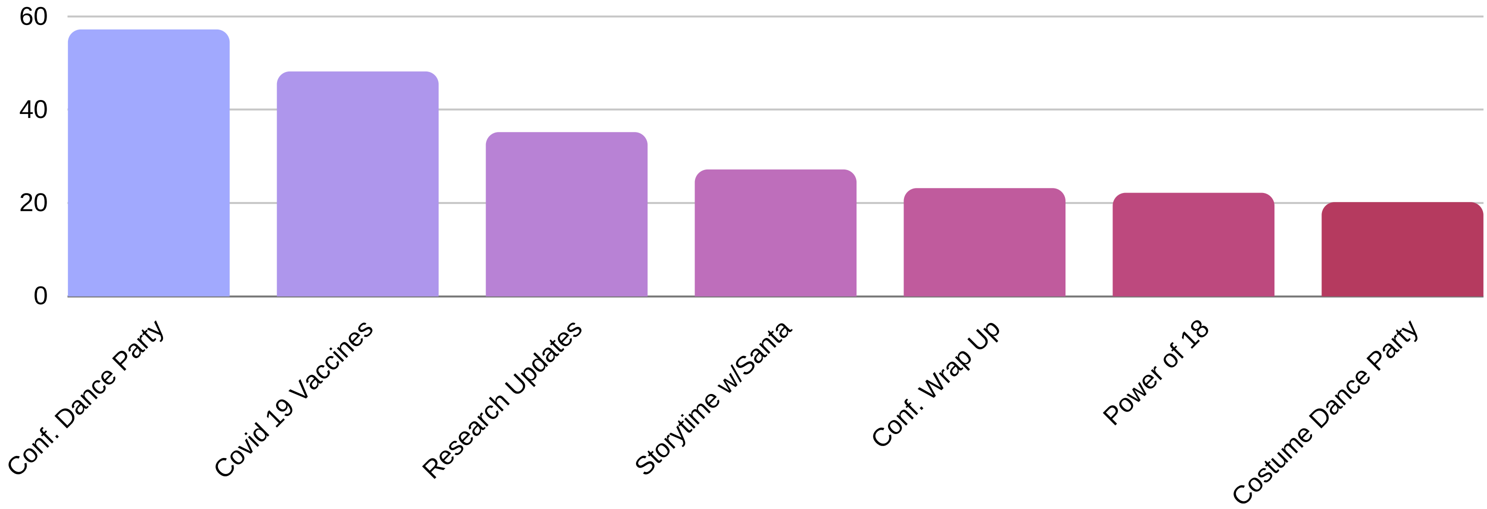
36

Virtual member engagement and educational events were hosted by Chromosome 18 in 2021. This averages to be three events a month!



2021 continued to be a difficult year for both Chromosome 18 and our families. We were grateful for the opportunity to continue our virtual event series that brought together families from around the world and provided connection opportunities for our families in a time of isolation. We are excited to continue this series into 2022!

Most Popular Events by Attendance



472

Unique Participants

11

Volunteer Coordinator Led Events

36

Activity & Engagement Events

Virtual Events

We continued hosting Virtual Events throughout 2021 to help stay connected with our global Chromosome 18 community. Some of the highlights include Self-Advocate Features, Parent Social Hours, Research Updates, New Member Welcome Calls, the Family Costume Dance Party, Storytime with Santa, and Virtual Tours of the Clinical Research Center. Even though we couldn't be together in person, we enjoyed meeting monthly for informational Zoom calls as well as fun social events.

The 18/80 Challenge!

Dan Vasquenza accomplished the 18/80 Challenge by running 80 miles on foot in under 18 hours (16 hours and 16 minutes), in honor of his son Jack (18p-), on April 18, 2021. Dan completed the 18/80 challenge to raise awareness and funds for Chromosome 18 Registry & Research Society. Thank you Dan for raising $22,837.83 for chromosome 18 research and for inspiring us all!

If you would like to lead a fundraiser for Chromosome 18, please email office@chromosome18.org

Regional & Fundraising Events

18 Fore 18

On August 7, participants gathered in memory of Jay T Parker, to spend a day filled with fun, food, golf, and lots of laughs. Thank you to everyone who participated & donated to the 18 Fore 18 Event in Oklahoma. It was a huge success! Thank you Holly Hubbell for organizing this event!

Brayden's Benefit

In September 2021, members held a 3-day festival with live music, good food, and a 5K at the Stony Fork Campground in Pennsylvania in honor of Brayden Blackwell. Everyone had a great time raising awareness and funds for chromosome 18 research. Thank you to everyone who helped make this event possible!

Ontario Family Picnic

On August 29, families gathered together for a Chromosome 18 family picnic in Ontario, Canada. Everyone enjoyed a beautiful day filled with good friends, food, and fellowship. Thank you Shelley Hunt for helping organize this event!

New Jersey Starfish Scramble

On November 7, participants enjoyed an afternoon of golf, comradery, food and fun at Top Golf in Mt. Laurel, NJ. Thank you for raising awareness and funds for Chromosome 18. Thank you to Liz Woodfield and Mike Fudala for helping put this event together!

Membership Resources

CHROMOSOME 18 CLINICAL RESEARCH CENTER PHYSICIAN MANAGEMENT GUIDES

UT Health
San Antonio
The Chromosome 18
Clinical Research Center

Released in 2016, the Physician's Management Guides were updated and expanded in 2020. They summarize 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.

Program Service Coordinator

Holly Hood
2021 Program Service Coordinator

During the "Snowpacalypse" of 2021, I gratefully accepted the position as Program Service Coordinator (PSC). Thank you to the Gordon Hartman Family Foundation, the MZ Foundation, and the generosity of a Chromosome 18 family for fully funding this position in 2021.

As the PSC, I enjoy reaching out to new families and long-time members to connect them with resources and connection opportunities. Through virtual events and our Closer Than Ever Virtual Conference, members were able to connect, share resources, and learn about the latest research.

I am very excited about our Virtual Library Project which will be available in 2022 to help families looking to hear more from other chromosome 18 parents and caregivers.

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

THE CHROMOSOME 18 CLINICAL RESEARCH CENTER

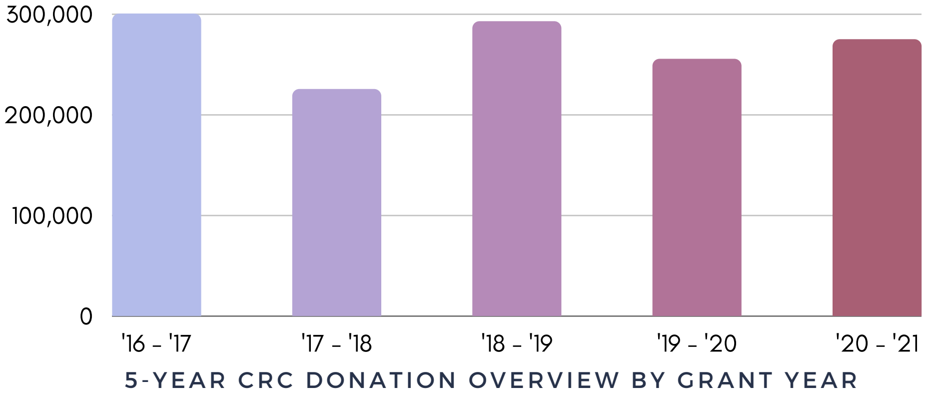
View the Chromosome 18 Clinical Research Center's 2021 Annual Report here.

THE CHROMOSOME 18 CLINICAL RESEARCH CENTER CELEBRATES 30TH ANNIVERSARY

The Chromosome 18 Clinical Research Center (CRC) saw a milestone year in 2021 - the 30th anniversary. Although we were unable to celebrate this momentous occasion in person due to the global pandemic, this anniversary holds tremendous importance. For the past 30 years, the CRC has stood as the only dedicated research center in the world for chromosome 18 conditions and continues the only decades-long longitudinal study of these conditions thanks to the families that have enrolled in the research and contributed to this vital project. This year, the CRC was accepted as a participant in the prestigious National Institutes of Health (NIH) Gabriella Miller Kids First Pediatric Research Program. This program funds the Broad Institute of MIT and Harvard to perform whole genome sequencing on 150 of our participants.

The Chromosome 18 Registry & Research Society is the primary funder of the CRC. As you can see from the graph below, 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 growing CRC funding. Increasing funding for the research would mean not only preserving the important discoveries, but moreover expanding the project in the future.

CHROMOSOME 18 CLINICAL RESEARCH CENTER FUNDING TIMELINE



UT Health San Antonio
The Chromosome 18 Clinical Research Center

705

participants enrolled in the research study

30

years as the world leader in chromosome 18 research

12

Research newsletters published in 2021

5

Virtual Chromosome 18 Clinical Research Center Tours conducted in 2021

Celebrating 30 Years with UT Health San Antonio!

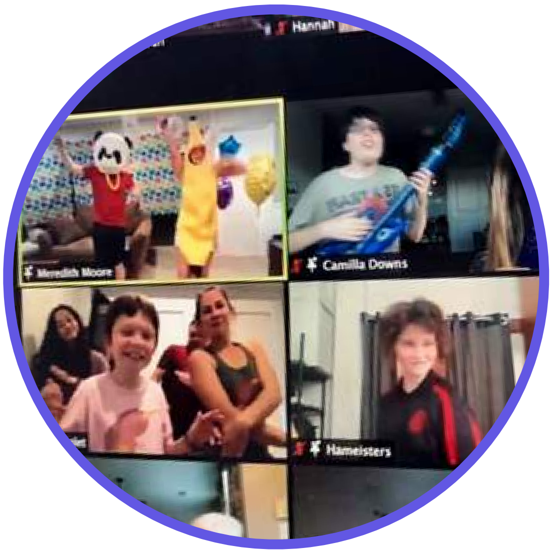
Thank you to UT Health San Antonio for your support as a partner to both the Chromosome 18 Registry & Research Society and the Chromosome 18 Clinical Research Center for the past 30 years. We are truly grateful to you as we look back and celebrate the impact of the research discoveries for our families over the last three decades. It is with your continued partnership that the Clinical Research Center at UT Health San Antonio has been a beacon of hope as the only dedicated research center in the world for chromosome 18 conditions over the last three decades.

Closer Than Ever Virtual Conference

JULY 11-13, 2021

GROUP PHOTO OF SELF-ADVOCATE BREAKOUT SESSION

2021 was the second year that the Chromosome 18's Annual Family Conference was held on a virtual platform. The international representation at our global conference was one of the highlights of the event. This year's conference saw over 11 countries in attendance, representing all continents except Antarctica! The 3-day event brought together over 40 expert speakers, and gave attendees opportunities to forge new connections and visit with old friends across multiple breakout sessions.



HIGHLIGHTS FROM THE VIRTUAL DANCE PARTY

LEFT TO RIGHT:
JANNINE CODY (PRESIDENT & FOUNDER),
NEALE PARKER (EXECUTIVE DIRECTOR),
MEREDITH MOORE (CONFERENCE CO-CHAIR)

Even though we would have loved to see one another in person, we were grateful for the chance to connect to our families from around the world during this special event.

Be sure to visit the 2021 Closer than Ever Virtual Conference page to access recordings of all of our sessions. Click **here** to get started.

For more details about the return of our in-person conference in San Antonio, TX, July 7-9 2022, click **here**.

2,300+

Chat messages sent between participants during conference

129

Total conference attendees

56

Conference sessions

44

Medical & professional experts presented

6

Continents represented (all continents except Antarctica)

17th Annual Golf Tournament

AUGUST 27, 2021

2021 saw the return of the Chromosome 18 Annual Golf Tournament to great success, it was our biggest golf tournament yet! Attendees enjoyed 18 holes and a fun-filled day at the beautiful Hill Country Golf Club. The tournament started with a presentation and balloon-release ceremony in memory of those chromosome 18 individuals we lost in 2020 and 2021. Meredith Moore, one of the Chromosome 18 Board members, emceed the evening dinner and awards ceremony, and the silent auction offered attendees a variety of fun items for bid. Thank you to our Golf Tournament Chairs Jim & Claudia Traa for hosting the return of this special event.

$40,000 total raised

100 golfers

$4000 in silent auction donations

Click here for more information about the exciting 18th Anniversary of the Chromosome 18 Golf Tournament, scheduled October 7th, 2022.

A Special Thank You to Valero Energy

Chromosome 18 is proud that Valero Energy once again was the title sponsor for the Golf Tournament and over the last 30 years has donated over $500,000+ to help our Chromosome 18 families. Huge thanks to Valero Energy and to John Drymala, our Treasurer and Valero employee who has helped steward this relationship over the last few years.

CHROMOSOME 18 EUROPE

Formed back in 2009 as a charity in Scotland, the goal of Chromosome 18 Europe is to bring European families affected by any chromosome 18 abnormalities together to share information and experiences together. Although the Annual Conference planned to take place in Rome had to be postponed because of Covid-19, families are excited to be connected virtually through different, social events, podcasts and the Annual General Meeting. We are proud to be affiliated with Chromosome 18 Europe and continue to ensure research carried out at the Chromosome 18 Clinical Research Center in San Antonio, TX is available to everybody around the World.

Click here to check out an interview with self-advocate Eve (pictured bottom left). She is the first self-advocate on the European podcast and in the interview she chats about her life and work.

Visit **Chromosome 18 Europe's website** for more information.

CHROMOSOME 18 AUSTRALASIA

Chromosome 18 Australasia has a devoted network of Chromosome 18 families. These families come together to support one another and to raise awareness of Chromosome 18 conditions in the community. Families gather in person at various regional activities, and then once every 4 years for an in-person Conference. The uncertainty of Covid-19 has seen these in-person activities postponed, but we are hopeful they will be back on the calendar soon. In the meantime, families stay connected to each other through online video meets and the Australasian Facebook page. Here you can meet our wonderful families and self-advocates like Claudia (pictured), who has mosaic Trisomy 18.

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

CHROMOSOME 18 CANADA

The Canadian Chromosome 18 families have been very active over many years with regional events happening on a regular basis. The families are in the process of forming Chromosome 18 Canada as a new organization and have the whole support of the other organizations in the USA, Europe & Australasia. Look out for some updated news on Chromosome 18 Canada coming soon!

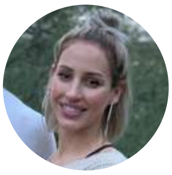
Chromosome 18 Office

MEET THE TEAM

Chromosome 18 Staff


Neale Parker
Executive Director


Holly Hood
Program Service
Coordinator


Amelie Simons
Manager of Operations

Accounting Students


Victoria Ochoa
Accounts Supervisor


David Garza
Accounts Assistant

Team Lead Students


Jude Casanova
Communication
Specialist


Mylinh Du
Communication
Specialist


Jagg Dotson
Communication
Specialist

Community & Development Students


Alyssa Avalos
Community Outreach
Assistant


Lillian Oss
Development
Assistant


Larissa Gonzales
Development
Assistant

Grants Students


Don Draper
Grants Assistant


Camila Zapater
Grants Assistant

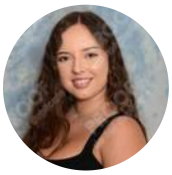

Leilani Garza
Grants Assistant


Adriana Pena
Grants Assistant


Nerma Pasic
Grants Assistant

Social Media & Marketing Students


Jaylen Machacon
Graphics Assistant


Mariaignacia Larrain
Marketing/Social
Media Assistant


Benjamin Creasy
Marketing/Social
Media Assistant


Rachael Graham
Marketing/Social
Media Assistant


Jose Hernandez
Marketing/Social
Media Assistant


Miller Burns
Marketing/Social
Media Assistant


Lisa Vetyuhova
Marketing/Social
Media Assistant


Ananya Lertpradist
Marketing/Social
Media Assistant

IT & Web Development Students


Kevin Nguyen
IT/Web Development
Assistant


Christina Kennedy
IT/Web Development
Assistant


Delon Perry
IT/Web Development
Assistant

Health Special Projects Student


Jessica Alvarez
2021 Conference Assistant

AS A COMMUNITY PARTNER, CHROMOSOME 18 HOSTS INTERNS AND WORK-STUDY STUDENTS FROM LOCAL UNIVERSITIES

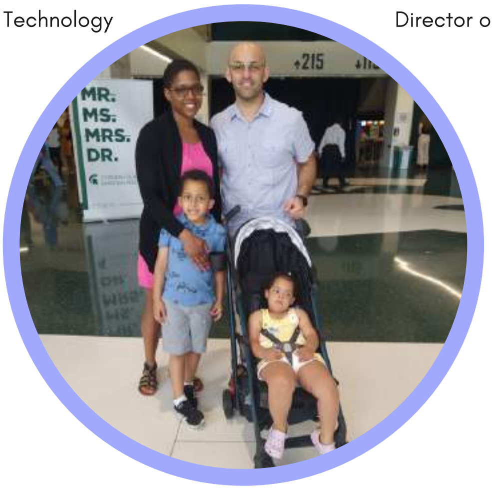
Chromosome 18 is proud of our partnerships with local colleges, universities, and community programs in San Antonio that allows us to host interns and work-study students on our team in various areas of specialty such as grants, social media and marketing, and development. In 2022, Chromosome 18 hosted 19 work-study students and 6 interns. During their time on our team, the interns and work-experience students get real-world experience in nonprofit work, professional development training, and the opportunity to hone their skills in their areas of specialty. It is with help from these students that Chromosome 18 is able to organize virtual and in-person events, keep members up-to-date with our social media platforms and our website, and better serve the chromosome 18 community. 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

Our Volunteer Directors donate an enormous amount of their time and effort to support Chromosome 18. We are incredibly grateful for them volunteering their expertise to help us better serve our members. Thank you to our amazing team Volunteer Directors and for all your hard work in 2021!

2022 MAJOR EVENTS

April 9th, 2022 - The Starfish Dash 5K Run/Walk & Health Fair at UT Health San Antonio
July 7-9, 2022 - The 27th Annual Chromosome 18 Decoded Family Conference, San Antonio, TX
October 7, 2021 - The 18th Anniversary of the Chromosome 18 Golf Tournament, The Quarry Golf Course, San Antonio, TX
December 4, 2022 - Chromosome 18 Virtual International Day (new!)
December 2022- 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 2022!

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

2021 Leadership Team

BOARD OFFICERS

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

DIRECTORS AT LARGE

Director at Large	Carol Connor Cohen
Director at Large	Tom Kunkel
Director at Large	Meredith Moore
Director at Large	Brad Jensen Sheppard

REGIONAL COORDINATORS

Great Plains	Kerry Connolly/ 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-	Katharine Newman/ Camilla Downs

ADDITIONAL COORDINATORS

Sibling Coordinator	Kendall Powell
Bereavement Coordinator	Camille Hammond
Self-Advocate Coordinator	Racine Ghiz

EX-OFFICIO MEMBERS

Europe	Bonnie McKerracher
Australia	Tracy Healy
Canada	Shelley Hunt

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

THANK YOU TO OUR DONORS!

$10,000 and Up:

Boys of '88 Fund
Jeff & Carol Connor Cohen
Marilyn DeMars
The Gordan Hartman Family Foundation
Kristin Palmer
Dan & Shea Stokes
Shea/Stokes Family Fund
Stephanie Stokes
Valero Energy Foundation
Dan Vasquenza
The Michael and Gina Wey Gift Fund
Liz Woodfield & Bill Carlucci
Carolyn Zaletsky

$5,000 to $10,000:

Dave & Tam Aldrup
Katie Bailey/Halo Branded Solutions
Larry & Juana Carter Charitable Fund
Elizabeth Cody
Jannine Cody
George & Traci Dunham
Doug & Julie Masterson & The Masterson Family
Mays Family Foundation
MZ Foundation
Neale Parker & Cynthia Parsons
St. Luke's Letheran Health Ministries
Daniel & Patricia Schmick
Peter B. Sullivan/Snow Family Trust
Claudia & Jim Traa
Chad & Patricia Watson
William & Jude Wolpert & Family

$2,500 to $5,000:

Charities Aid Foundation of America
Michael Fudala
GlaxoSmithKline Foundation
Daniel Hale M.D.
Holly & Kyle Hubbell
Janine Macchia
Meredith & Kent Moore
Tom & Angie Schilly
Brenda & Joseph Schumaker
Sheppard Family
United Way San Antonio/Bexar Cty.

$1,000 to $2,500:

ABB Sponsorhip
Adolescent Medicine
Alicia Alongi
Americas Best Local Charities
Asel & Associates PLLC
Blue Ivy Partners
Brittney Bas-Telles
Alan Beaton
Thomas Behr
Rhonda Bentley
Rachel Bond
Heather Broganmyer
Cristi Cain
Scott Carlson
CleanScapes LLC
Carrie Colella
Kelly Collins
Antoinette Cox & Rebecca Parker
Howard Davis
Paul DeMars
Documation Inc.
Charles & Robin Eisenbeis & Family
Lorna Feijoo
Peg, Margaret, & Ben Flowe
The Pittsburgh Foundation
Lone Pine Foundation, Inc.
Stacey Gallardo
Becky Gaylor Hobby
Jay & Susan Goldberg
Grande Cares Club
Barbara Grossman
Elizabeth & Seth Grossman
Dalton Guthrie
William Guthrie
Tahira Haque
Myrna & Stephen Harris
H-E-B
Steven Hessing
Jodi Hight & Hight Real Estate
Annice Hill
Constance Hubbell
Thomas Immenschuh
Jay Hirsh/Jumbo Capital Mgt LLC
William Keen
Jennifer Kunkel
Patricia & Devon Lacey
Judith LaFrombois
Catherine Larson
Nora & Eric Leaderschaefer
Julian Maha
Elizabeth Martel
Andrew Meyer M.D.
Marcy Nichols

$1,000 to $2,500:

Tim Ogden
Dean & Jennifer Oltmanns
Owasso Pediatric
Denise Parker
Puttie Potts
Countess & Frances Price
Christine Puckett
Sean Ralson
Jessica Robold
Brenda Roder
Kayvan Romaezi
Kristin Schneider
David Shepard
Todd Shively
Raquel Silhy M.D.
Topnotch Silkscreening
Rebecca Teague
Gene & Martha Schaerr
The Schaerr Family Fund
Sprinkler Fitters Union 692
Yvonne Tritto-Lisella
TX Cavaliers Charitable Fdn.
Joseph & Theresa Verstreater
Walmart Inc.
Eric & Maggie Welle
Jennifer Wessman
Brian & Kathleen Wessman
Anthony White
Lewis & Susan Wiens

$500 to $1,000:

Sarah Adeyeye
John Bailey
Patricia Bailey
John Bottengerg
Robert Burr
Helen Carr
Mike Carr
Steve & Betsy Caviness
United Way of Central Ohio
Damon Torres
DisabilitySA
Carrie Connolly
Camaron Corr
Wendy Crawford
John Drymala
Jeff & Kristen Earl
Carol & Wayne Edwards
Raul & Sandy Flores
Mickey Fluitt

$500 to $1,000:

Michael Furgiuele
Alexis Garneau
Shelby Griffin
Michael Hampton
John Hawley
Judy Heggie
William & Theresa Hill
Olivia Hinden
Greg & Juliette Hitchan
Holly Hood
Donald and Carla Hughes
Anuradha Jaishankar
Shaunna Keller
Amber Koblick
Amyla Lavric
Brian Leivo
Nina & John Lewis
Karen Marcham
Grace Mckee
Jill McLain
Christopher Mosher
Charli Mosteller
Andrew & Shannon Mueck
Chrissie Oh
Owen O'Neill
Scott Peeler
Bea Powell
Tammy Powers-Taylor
Mike Regina
Sandy Rocka
Michael Rush
Thomas & Martha Sakre
John & Elizabeth Santo Salvo
Robert Schmitz
Tracy Sims
Chuck Smith
Deborah Stevens
Amber Styron
Audrey Suchicki
Lewis & Dee Noonberg & The Noonberg Family Fund
Mike & Karen Todd
Patrick Tracey
Michael Uzzell
Janet Verhulst & niece
Cindy & Dan Wetmore
Wetmore Family Charitable Fund
Brian Wunderlin
Kristen & Debra Yerys

Thank you to our amazing donors who supported Chromosome 18 in 2021 with your direct donations and by hosting fundraisings event. This list reflects only direct donations from individuals or corporations. We endeavor to ensure that all donors is 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
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Legacy Society

THANK YOU TO OUR FOUNDING MEMBERS

Jannine Cody
Jeff & Kathy Borello
Marilyn DeMars
Mike Fudala
Dan & Pat Schmick
Rosemary Woodfield

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**.

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

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Thank You for a Wonderful Year!