

# A Bright future

I am Katie Hollis, the new Executive Director of the Down Syndrome Association of West Michigan (and its Foundation). I am the mother of two children, a 13-year-old girl I adopted in 2019 and a two-year-old boy with Down syndrome. Prior to accepting the role of Executive Director, I was a member of the organization who regularly attended the Moms' Night Out support group and Baby Sign Language classes; and was wondering how I could get more involved. I am trained as a molecular biologist, with experience in teaching and project management, a heart for fundraising, and a hobby in event planning. When the opportunity to lead this organization came up, I knew this was where I was meant to be. I am excited to lead DSAWM and see how we can grow together to support our loved ones with Down syndrome and advocate for their full inclusion in the West Michigan community.

2021 brought a lot of big changes for the Association. At the start of the fourth quarter, the Executive Director of almost 5 years, Jennifer DeVault, accepted an amazing opportunity as the new director of

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2022-2023 Members-At-Large

Rich Kloeckner\*

\*elected in 2022

Thank you to Jennifer DeVault, Nate Clark, and Tom DeVault for their tenures. We wish them the very best in their new career adventures.

We extend a heartfelt thank you as well to our Board members whose service ended in 2022: Niurka Diaz, Tom Hackett, Molly VandeWater, Michael Wolfe, and Dave Wood.

Kent District Libraries. We thank her for her years of service and for helping the organization to thrive during her tenure.

Under her direction, we started 2021 by moving into the Special Olympics Unified Sports and Inclusion Center. The new space has allowed for growth in many ways, including launching our Connecting Communities capital campaign, a summer day camp, and the 21 Shades support group for multi-racial families. While some of our programming remained virtual, we took advantage of our new home's Life Skills room to host our popular Cooking Capers program, providing just a small peak at what this new facility has to offer.

DSAWM also began focusing on collaborative programming opportunities for our members. We hosted a dance program with Reaching with Rhythm and wrapped up the year with a new collaborative therapeutic playgroup with Comprehensive Therapy Center.

As I began my new role in February of 2022, I saw the Association was positioned in an environment full of opportunities. Through townhall meetings with our members and analysis of the organization, we began preparing a new strategic plan. We are now in the process of re-launching committees and improving the efficacy of the organization. We have also been hard at work growing our staff and board, while also embracing

collaboration with community partners

and other local non-profits. I look forward to continuing the great success of this organization and ensuring a bright future of

opportunities for those with Down syndrome in our community!

Katie Hollis
Executive Director



## Hola. Estrella's!

It's not often that you see Diane Estrella still. Whether she's at the head of the class teaching her students, behind the wheel chauffeuring a group to afterschool activities, or at a beloved nonprofit volunteering her time, Diane is almost always in motion. The same is true of many moms who, like Diane, have a child with Down syndrome. Days are spent moving from the classroom to therapy visits to social programs to doctor's offices and more. It's a busy life, but, Diane says, she wouldn't have it any other way.

In 2003, while living in the Dominican Republic, Diane welcomed her son Elias. A native of Michigan, Diane had moved to the Dominican Republic to teach and start a family with her husband Naun. The couple had previously welcomed three daughters and were excited to have a boy in the family. It was only after his birth and on a visit back to the United States, that they learned Elias had Down syndrome. "We were introduced to the Down Syndrome Association of West Michigan pretty quickly," Diane recalls. "It was so nice to connect with other families."

The Estrella's continued to foster those connections on their return visits to Michigan every summer. Building a community aided their decision to permanently move back to the U.S. when Elias was diagnosed with leukemia at age 6. "We loved our home in the Dominican Republic and that Elias had full inclusion there, but the need to be close to family and have more support played a big role in our decision to return." Thankfully, Elias' leukemia entered remission, and with his health no longer a barrier Diane and her family's involvement with DSAWM flourished.

Elias is now 19 years old and a regular at DSAWM programs including Rapid Runners, Cooking Capers, Music Jamboree, Buddy Up Tennis...the list goes on and on! "He's made great friendships over the years," Diane says. It's these relationships that Diane credits to enriching her family's experience with DSAWM. "Elias enjoys having friends to go to activities with...and his involvement has created opportunities for his siblings to volunteer and build their own friendships with people with

disabilities." Elias' sisters are now pursuing careers in special education and recreational therapy, paths that Diane says are directly impacted by having Elias as a brother.

Diane's eagerness build relationships has also shaped her role in the organization. As one of DSAWM's most active bilingual members. Diane has connected dozens of Hispanic families with DSAWM programs and services over the years. In August 2021, she helped launch 21 Shades, a support group for families of color and multicultural families. The group meets for social activities and workshop opportunities every few months, with most families currently participating identifying as Hispanic or Latino. "It's important for families to see other people like them at programs," Diane says. "It's been exciting to see families who come and discover the value in DSAWM and the connections it offers, and are now

Diane and the Estrella's continue to live lives in motion. As Elias grows into adulthood, new opportunities are emerging, and Diane will continue to embrace her roles as teacher, chauffer, volunteer, cheerleader, and friend...aka Mom. She'll also add advocate to that list as she finds more opportunities to connect with families in the community, having recently joined DSAWM's Step Up for Down Syndrome Planning Committee and the First Call Committee. "I want

convincing others to join and participate as well."



# Our Impact

While the Covid-19 pandemic continued to impact our operations in 2021, widespread distribution of the vaccine gave DSAWM confidence to cautiously resume some in-person programming and events. In the spring and summer, popular skill-building programs including Cooking Capers, Rapid Runners, and Book Club began meeting outdoors. Summer Day Camp kicked off with 27 school-age members enjoying a week of fun activities and socialization with new friends, Important member events like the Summer Picnic on the lakeshore and Holiday Party in downtown Grand Rapids were also well-attended and gave families the opportunity to connect after being apart for over a year. Step Up for Down Syndrome was, as always, a highlight of the year and over \$89,000 was raised to support DSAWM.

We are grateful to our staff, board, and volunteers for working hard to ensure the challenges of the pandemic did not hinder

our services and outreach. It takes

Greating Employment opportunities

in the office including writing donor and volunteer thank you cards, assisting with event

preparations, and engaging in advocacy opportunities within the community. In September 2021, DSAWM hired Eric Willinger for the role. Eric has a sharp sense of humor and is extremely personable, traits which make him an engaging advocate both in and out of the office. Eric also works at Chick-fil-A and was the first person with Down syndrome to be hired at any West Michigan location. In his free time, he's an avid bowler, enjoys basketball, and plays several musical instruments. We're thrilled to have him on the team!

•• We are brodram! -Member Candace Martin

287 member families engaged and connected

unique programs and events offered



1.269

program and event attendees



## The foundation

The Down Syndrome Association of West Michigan Foundation operates alongside DSAWM to support the West Michigan Down Syndrome community. Through the management and growth of its endowment, the Foundation supports the annual operating budget of DSAWM while also supporting other organizations that consistently and actively serve the Down syndrome population.

While the pandemic forced the Foundation to cancel it's annual springtime fundraiser, the Winners Cup Benefit, a new event was introduced in October. The Halloween Casino Night invited Foundation supporters, old and new, to a lively "friendraiser" at Kent Country Club. Organized by Winners Cup Committee Co-chairs William and Bobbi Blanton, the event gave attendees the opportunity to mingle and network while enjoying a fun night out.

The Halloween Casino Night was also an opportunity to celebrate the launch of the Foundation's Connecting Communities Capital Campaign. The \$850.000 fundraising effort will allow the Foundation and Association to invest in their new headquarters at the Special Olympics Unified Sports & Inclusion Center, fund program expansion,

> and enhance the endowment fund. During the campaign's public announcement in August

2021, campaign co-chairs Rich Kloeckner

and Jim Leach shared that over \$800,000 had already been

raised!



### foundation Board of Directors

**Bob Bovlen Board President** 

Katie Hollis Executive Director Robin Zondervan Treasurer

William Blanton, Cameron Dolbow, Tom Hackett, Jim Scarlett 2022-2023 Members-At-Large



The Foundation is grateful to the following donors who have contributed to the Connecting Communities Capital Campaign.

Chris A. Anderson Bill and Carol Ashcroft Dan and Jo Baiema

Chloe Bayer

Bill and Amy Bennett Jim and Judy Blanchard Joel and Laura Blanchard

The Bode Family Robert C. Boylen Ed and Linda Bueche Building Synergy Nate Clark

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Paul and Sheena Vogel

West Michigan Eyecare Associates

Amy and Mike Wigger Eric Willinger

Margaret A. Wilson

Matthew and Laura Witkowski

Dave Wood

Ray and Robin Zondervan

Gifts received in memory of Jim Leach, Sr.

#### Save the Date

### The Winners Cup Benefit

A Kentucky Derby Party and Auction to Benefit the Down Syndrome Association of West Michigan Foundation

May 6, 2023

at Egypt Valley Country Club



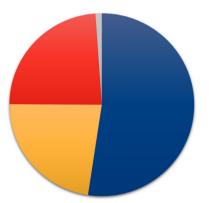
# Financials



# TO DSAWM

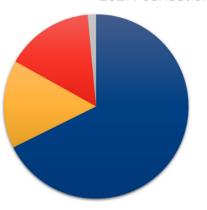
Foundation

#### 2021 Association Operational Revenue



Public Support	\$205,356
Grants	\$93,374
Fundraising Events	\$88,816
Membership Dues	\$4,258
Total Revenue	\$391,804

#### 2021 Foundation Operational Revenue



Public Support	\$293,654
Endowment Fund Investment Income	\$67,294
Unrealized Gain on Endowment Fund	\$67,043
Fundraising Events	(\$6,454)
Total Revenue	\$421.537

#### 2021 Association Operational Expenses



Programs & Services	\$209,437
Admin & Management	\$78,660
Fundraising Costs	\$59,295

ograms & Services \$209,437

Total Expenses \$347,392

### 2021 Foundation Operational Expenses



Programs & Services	\$159,097
Fundraising Costs	\$30,000
Admin & Management	\$8,298
Total Expenses	\$197,395

At the 2021 year's end, the Net Assets of the Down Syndrome Association of West Michigan Foundation totaled \$1,736,219.

## Thank You!



We are deeply grateful to the many donors who support our organization. Because of your contributions, our programming remains free and accessible to families across West Michigan and we are continuously able to expand our outreach and services. We are humbled by your generosity!

A very special thank you to the following sponsors, grantors, and donors whose \$250+ gifts supported us in the 2021 fiscal year.

Tyler Berger

William and Bobbi Blanton

Boardwalk Subs

Mr. and Mrs. Drew Boersma Stephen and Susan Britton

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South Christian High School

Paul and Beth Taylor Jordan Thelen TJX Foundation Tuff Cover

Paul and Sheena Vogel Richard and Julie Wadel Patrick and Ashley Warren Doug and Tracy Warsen Michael and Keri Weber

West Michigan Eyecare Associates



## On the Horizon

Year-round Programs

21 Shades Support Group Aging with Down Syndrome

Book Club

**Cooking Capers** 

Moms & Dads Nights Out
Self-Advocate Board

Winter/Spring 2023

Shape Up (also available in Fall)

Buddy Up Tennis (also available in Fall)

Baby Sign Language Reaching with Rhythm\*

Rock Your Socks Dance (March 18) Winners Cup Benefit (May 6) Annual Meeting (May 8)

Summer 2023 Rapid Runners

iCan Bike Camp (June 12-16)\*
Adaptive Water Ski Clinic (July 21)\*
Member Summer Picnic (July 23)

Oh, Baby!

fall 2023

Step Up for Down Syndrome (September 9)
Member Holiday Party (December 3)

\*partner program

### Welcome. Melissa and Peri!

In August 2022, DSAWM welcomed two new staff members: Melissa Perrin and Peri Phillipi. Melissa oversees the development and expansion of our programs and services in the role of Program Director and Peri provides frontline support and assists with program implementation as the new Program & Administrative Assistant. "I am thrilled for the opportunity to take on a more active role in the programming of the DSAWM and help shape the future of this fantastic organization," says Melissa, who first became involved with DSAWM four years ago when her son Osgood was born with

Down syndrome. Peri similarly has a strong connection to our community. "After assisting in organizing and teaching a work skills class at DSAWM in 2019 I was very eager to find more opportunities to work with this organization. I look forward to taking on the role of the administrative assistant for DSAWM and getting a chance to find new ways to support and be an advocate for people with disabilities in this community."













