

annual report 2019





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GREATER
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HD

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Looking back at 2019 it's hard to not look back at the whole decade. Can you believe that when the decade began, HDYO wasn't yet even an idea on paper? From an idea, to an entity to an international organization that's shifted the paradigm on how young people impacted by Huntington's Disease are supported, educated and empowered. Wow! I'm grateful, thankful, blessed and indebted to so many who have helped make our mission a reality!

HDYO continued to take exponential strides in 2019 offering more programs, shifting more energy into research and building a stronger team to bring everything to life!

Programs:

Every program we ran was created and implemented with young people's needs at the heart of that decision. For the first time ever we had over 1,000 young people from around the world directly reach out to us with their questions or support needs. As you read through this report keep in mind that we are still a small team doing a whole heck of a lot of work around the entire globe.

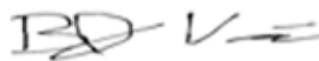
Research:

It's amazing to see the increased interest for HD treatments from the biotech industry. When this decade began, Lundbeck was really the only company showing any interest to HD. That has drastically changed with more companies than ever before with programs for HD. That's also changed how HDYO thinks about research. We have taken major strides to make sure that the young person's voice is heard and respected by these companies as the young people are the future of these treatments. We have a lot of work to do to fully get the young person's voice into the ear of these companies, but are excited to keep working hard.

Team HDYO:

We continue to search for dedicated individuals who are passionate about positively changing the lives of young people through the HDYO platform. In 2019 added seven amazing board members from around the world to take on different roles and look to keep growing that number. We are super excited about all the work we've put into the creation of our Young Leaders Network that will officially launch in January 2020 with two pilot cities in the USA (Philly and Washington D.C.). If you are a young adult and looking for a cause to get involved in, please reach out to join or start an HDYO team!

2020 is going to be an amazing year for HDYO. We are hosting an international congress in Glasgow, we will be launching an international patient registry for the Juvenile HD community, we will be growing our Young Leaders Network and we will continue to put our hearts and souls into everything we do for young people! Join our team, support our work, or spread the word about HDYO!



BJ Viau

Board Chairman and Co-Founder
BJ@HDYO.org / 952-270-5428



Executive Director Report

02

A year of partnerships is probably the best description of 2019 for HDYO. As a team we have strengthened, developed and re-affirmed our commitment to work with the whole HD community to improve education, services and understanding of the true impact of HD on young people.

I have the privilege of representing HDYO at many events around the world and 2019 HDYO certainly made its mark. The year started with myself & Chandler speaking at the first South East USA HD Symposium in Nashville and Lauren creating an interview series at CHDI therapeutics conferences in Palm Springs.

In **April** BJ, Eric & I did a whirlwind tour of Boston pharma companies (6 companies in 24 hours!). We are so excited by the commitment and passion that these partners have in bringing effective and safe therapies to market for those impacted by HD and will be actively promoting young people being involved at all stages of development and delivery.

Our Latin American volunteer, Laura represented HDYO at Help4HD conference in Puerto Rico and did an amazing job.

In **May** I was honoured to be the keynote speaker at Huntington's Queensland Planning for the Future forum. We also delivered a fab workshop for professionals on working and engaging young people. Thank you to Huntington's Queensland for securing funding to allow HDYO to be there. HDYO also held our very first fundraising event in Washington DC, we are hoping to make this a more regular event!

In **June** the USA team were out in force at HDSA

convention in Boston and we are so grateful to Alan Rotberg for his amazing event before hand that raised vital funds for HDYO and our partners HDSA NYA.

August was busy with camps. I delivered three workshops at HDANI Summer Camp in Northern Ireland on a range of topics designed specifically for the group. It was then off to San Diego for HDYO North American Camp followed by a trip to DC for some planning time with Chandler then speaking at Tennessee Education Day.

September took us to Prague, Czech Republic for a development meeting with Heathe RND team (exciting new project) followed by a trip to Nice, France for Movement Disorder Conference where I met with the young MD specialists committee and spoke with professionals from all over the world.

In **October** Matt flew the flag for HDYO at EHA conference in Bucharest and also delivered a workshop to professionals in partnership with Romanian HDA.

November saw most of the HDYO board and staff team head to Sacramento for HSG conference where we were delivering workshops, giving presentations, meeting with partners on exciting joint projects and meeting lots of members of the HD community at the HDYO table. HDYO also held an ice cream and beers happy hour at the start of HSG...this could be an annual event? We ended that week with an in person board meeting and development session.

The next 3 year strategy along with lots of exciting announcements will be launched in Spring 2020. November also saw Matt head to Czech Republic HDA conference to deliver a presentation on HDYO

and share his family experience of PGD.

December didn't slow down much with a trip to Rome to present HDYO JoHD Registry update to LIRH annual family conference before Ashley for HDANI came to visit HDYO office for 3 days to do some job shadowing.

As always it's been a busy year but 2020 is shaping up to be bigger, busier and better. We are all looking forward to welcoming so many young adults to Glasgow, Scotland in May. Our new look website will be launched, our registry projects will go live, we have two new projects for young people that we are launching and so much more.

Make sure you are signed up to our mailing list so that you don't miss out on any updates and opportunities!

Thank you all for your support and sending us your stories, questions and feedback.

Cat Martin

Executive Director

Meet the HDYO Team of 2019

03



Staff Team

Cat Martin

Executive Director – UK Based
catherine@hdyo.org

Matt Ellison

Project Co-Ordinator – UK Based
matt@hdyo.org

Chandler Swope

Director of Youth Services – USA Based
chandler@hdyo.org

Paula Richmond

PA to Executive Director – UK Based
office@hdyo.org

Board of Directors

BJ Viau – Chairperson

Donna Spence – Vice Chairperson

Danielle Valenti – Secretary

Bonnie Hennig-Trestman – Research Co-chair

Lauren Byrne – Research Co-chair

Clare Braithwaite – Marketing Co-Chair

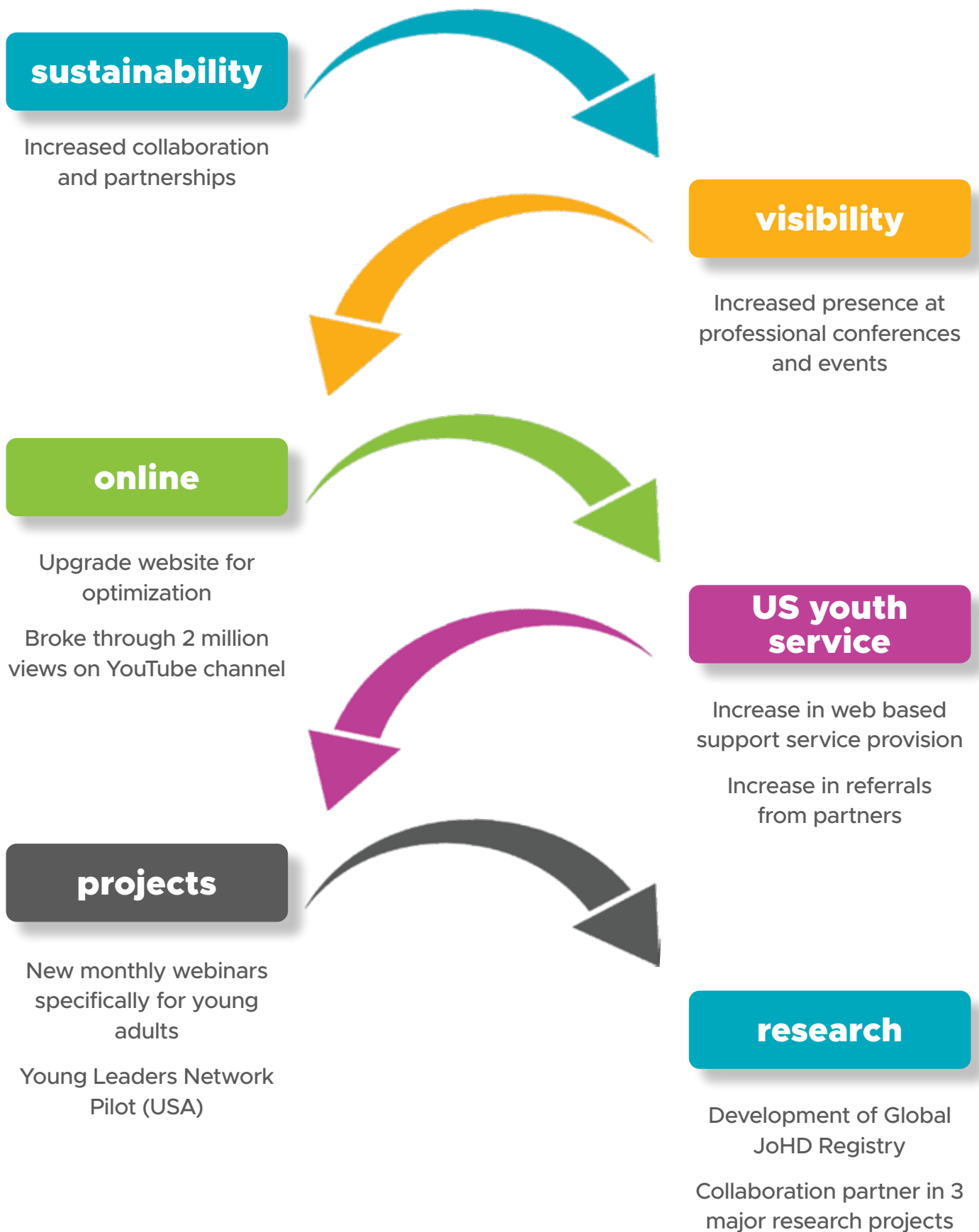
Ana W – Marketing Co-Chair

Eric Miller

Lindsay Morrison

Seth Rotberg

Jimmy Pappadeas



Organizational Sustainability

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New Partners

HDYO has developed partnerships with 12 new organizations and pharmaceutical companies.

Education Events

HDYO attended and facilitated 18 education and training events.

HDYO ran weekly clinical training programme with Vanderbilt University Medical Centre HD COE.

New Funders

5 new Industry partners.

Online donations through social media & website continue to grow reaching \$35,000.

New UK partnership with grant writer to develop funding bids through Trusts & Grants.

HDYO held its first fundraising event in Washington DC in May 2019 and raised over \$6,000

Conferences

Attended a record number of 14 conferences in 8 countries.

From these conferences we have increased awareness of HDYO services, gained new partnerships and attracted interest from outside HD community for collaboration.

Consultation & Advisory Board Meetings

HDYO sits on 5 advisory boards providing consultation on protocols and engagement for 3 clinical trial partners.

HDYO manages 2 advisory boards that are developing patient and caregiver registries for

Juvenile onset HD and Young People Impacted by HD.

Research Programs

HDYO partners in 3 research programs. One of the programs is being led by HDYO in partnership with NHS England looking at the impact of Double Disclosure genetic testing on families.

HDYO is leading on two patient and caregiver registries for the HD community. The most important of these is the Juvenile onset HD Registry.

New date: March 12th – 14th 2021

Our first ever International Young Adult Congress was due to take place on May 9th through 11th May 2020 where we expected to welcome over 400 young adults to Glasgow, Scotland. However, due to the global threat from COVID-19 and with the health and wellbeing of everyone we work with as a priority we took the decision to postpone Congress until March 2021.

Congress is aimed at young adults aged 18-35 impacted by HD from around the world to bring them together to learn, support, empower and educate the HD community on the true impact of HD. We are also delighted that professionals from associations, clinics, research and industry are supporting Congress and see the benefits of engaging with young adults directly.

Congress planning has been a huge project for HDYO over the last two years with securing funding, venues, sponsorship, speakers and activities. We have 6 young adult led teams that have been helping us make congress happen and with their help we have a programme featuring more than 50 speakers, the Scottish premier of Dancing at the Vatican documentary, activities throughout Glasgow including some inner city White Water Rafting, escape rooms, city tours and much more. Our final event party will be a huge celebration of the amazing participation we will witness over 3 days as well as some interesting dancing, singing and accent challenges!

HDYO has been extremely humbled by the support from partners for Congress. Over 20 national and local HD Association are providing funding for young adults to attend Congress, this along with HDYO's

community raised Scholarship Fund will support over 150 individuals. A special thank you to the following individuals who have collectively donated more than £20,000 toward HDYO Scholarships Dr Bonnie & Dr Bob Trestman, Lindsay & Kyle Morrison, Gimbel Family Scholarship, Jacqui Harrison. To each of those who have donated, fundraised or sponsored scholarship we truly appreciate every single penny!

Congress is a truly unique event for the HD Community it will be special that is why we will be live streaming the plenary events for those who can't join us in person. The Congress app will also allow EVERYONE to participate.

For all the details about congress go to www.hdyocongress.org or email events@hdyo.org



Meet the Congress Teams

website & marketing team



This team is responsible for the design and development of all online content for congress, including the website, new logo and social media videos

programme team



This team is responsible for development of the Congress programme, speakers and workshop content

scholarship team



This team is responsible for developing, fundraising, awarding and supporting HDYO Congress scholarships.

fundraising team



This team is responsible for securing sponsorship and community fundraising to ensure we have enough money to deliver the best event for young adults impacted by HD

volunteer team



This team is responsible for recruiting and managing volunteers who will be helping us over the week of Congress.

events team



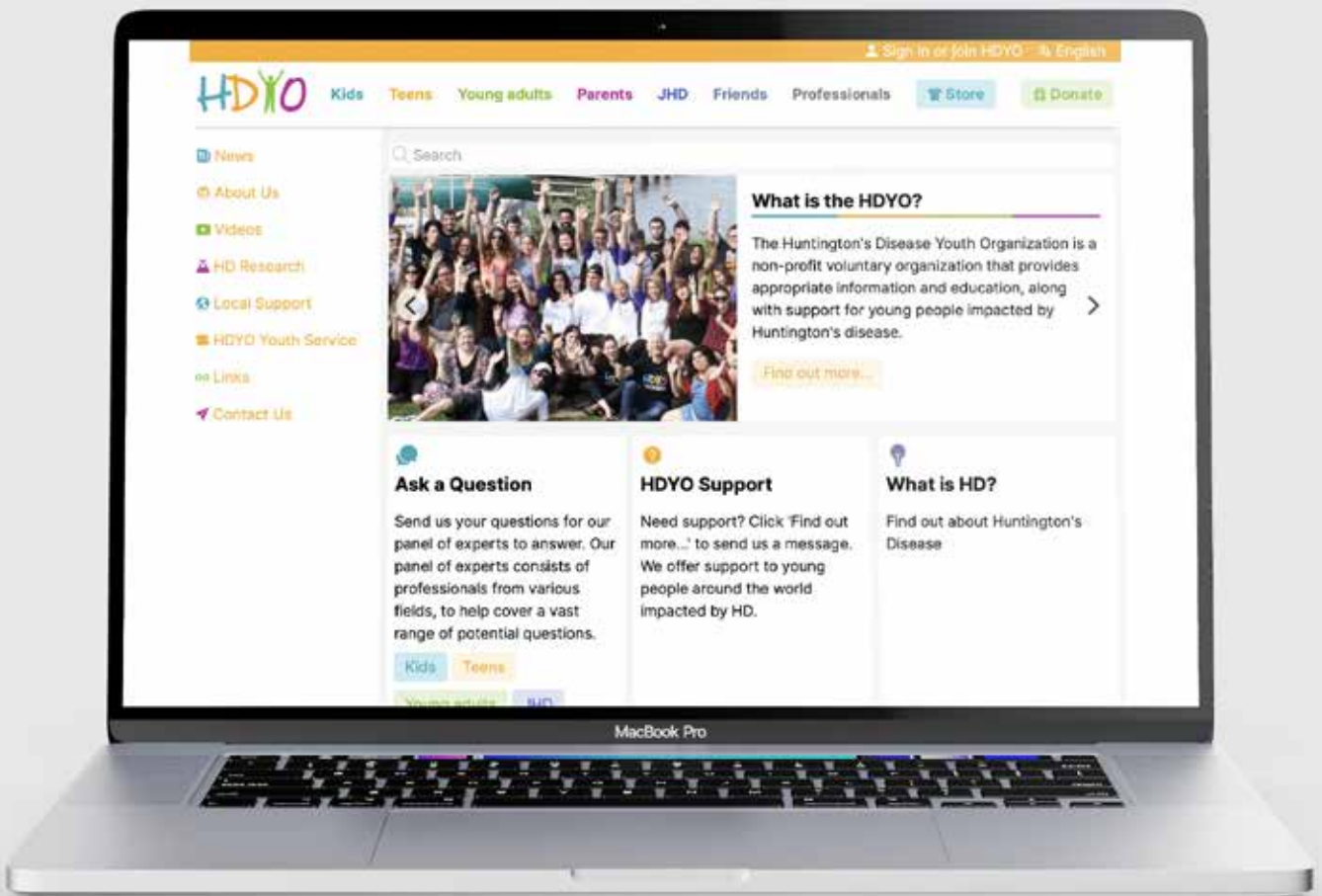
This team is responsible for ensuring all surrounding events, activities and final party are a success. They have helped source activities, venues, entertainment and food!

We have been working for months with a web developer Marc to update our website, a complicated undertaking given the massive amount of content and the translation to 14 languages. The first part of the update has been completed, and our website is now easier to read on any device — but we aren't stopping there!

The next updates are in progress now and we should be launching the website in May 2020.



current website



new website

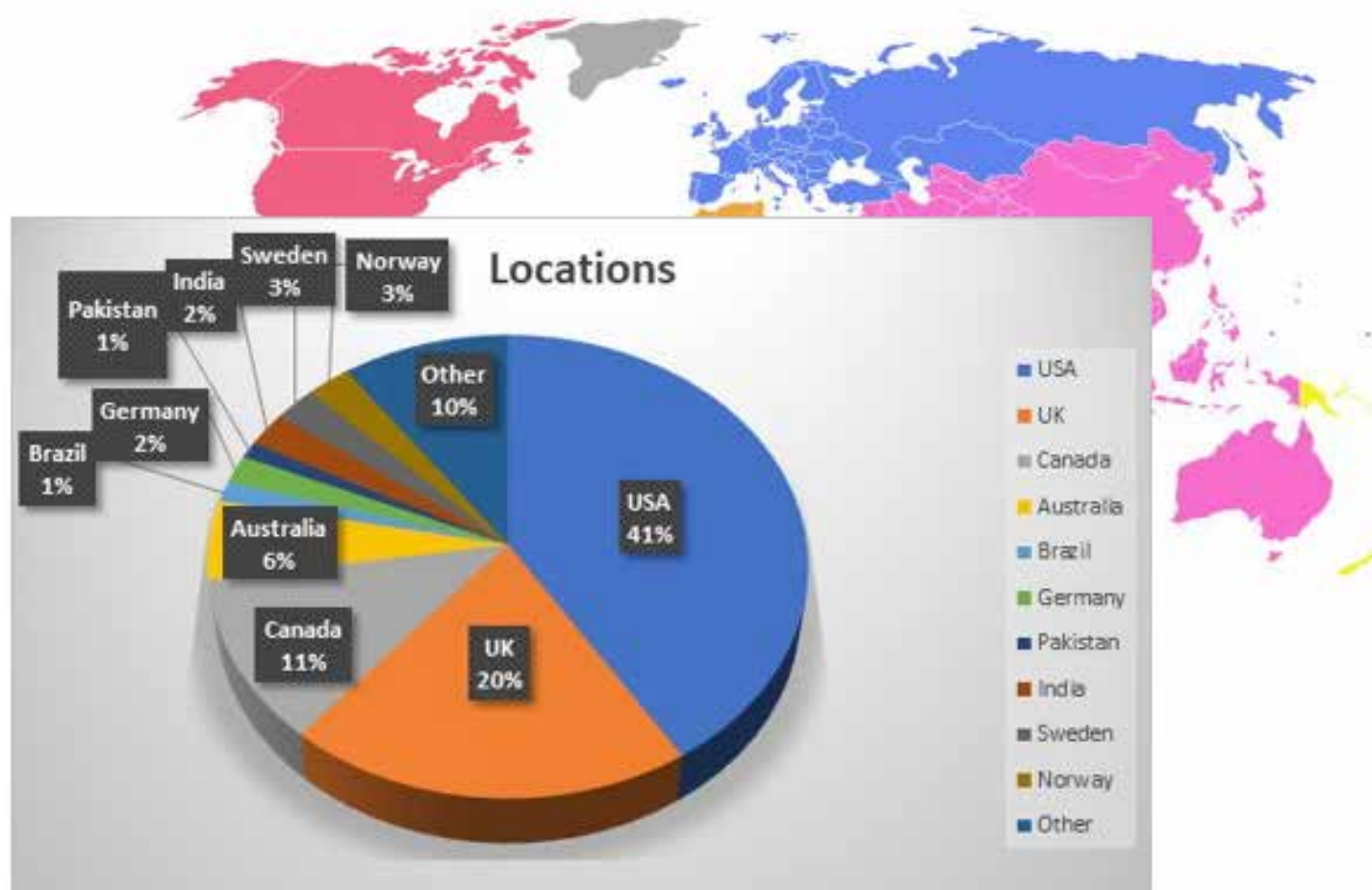
HDYO Direct Support Services

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Contact Story

We've had many impactful contacts such as a young person in Europe who reached out to HDYO multiple times this year as they felt extremely isolated with their thoughts on HD. In a number of contacts they were putting themselves at serious risk of harm so we had to breach 3 times this year to ensure they were ok. During these contacts

we were securing support options for the young person and in communication with their family to try and improve the isolation felt by the young person. They are now in a better place for the time being, it's always difficult, and doing well with the support now around them in their country, as well as HDYO always here if needed.



Help4HD

Our partnership with Help4HD continued this year. The HDYO Youth Worker attended 2 H.I.P.E days and their annual symposium. HDYO also sent a trained, Spanish-speaking volunteer to their Puerto Rico event to discuss the services and support HDYO provides in the patients preferred language.

HDSA and NYA

HDYO and the NYA continue to work together to create and facilitate events for young people in the United States. The Youth Worker attended and facilitated workshops at all the NYA retreats as well as the HDSA Annual Convention. Jennifer Simpson, Senior Manager of Advocacy & Youth Programs, attended and led workshops at the 2019 North American Youth Camp.

HSG

HDYO continued to work with HSG to ensure that the impact and needs of young people in the HD Community were discussed at their annual event. HDYO's Director of Youth Services took the stage at the event to talk about Genetic Testing amongst young adults and Executive Director, Catherine Martin, led youth-friendly workshops at their annual Family Day. This partnership continues to grow and evolve as we see young people more eager to get involved and have a voice in the larger scientific community.



Partnership in Action – Meet Sharmin and Hillary Page

In March, the HDYO Youth Worker received a referral for Sharmin and Hillary from the HDSA Social Worker in Louisville, KY. Sharmin had recently lost her husband and Hillary had been diagnosed with HD shortly after.

After some consultation with the social worker and Sharmin an appropriate plan was developed with the referring social worker, the closest CoE (Vanderbilt) and HDYO. Hillary was referred to the Vanderbilt Clinic for clinical care and support from their clinic team, Sharmin was provided support and services from the Louisville Social Worker and HDYO provided case consultation and supervision on the work. Case consultation happened by HDYO leading a series of weekly calls to assess where the family was at, what their current needs were, reviewing best practices on interventions and helping those involved navigate a complex family dynamic. HDYO continues to assist and provide supervision and technical assistance on the case.

Since the referral, both Sharmin and Hillary have received appropriate clinical care, signed up for a clinical trial and attended the HDSA National Convention together.

Hillary also attended a NYA Youth Retreat where she connected immediately with another young person and was comfortable and confident to share her HD journey.



Matt was invited to Czech for a talk at their HD family event to spread the HDYO message. As usual I encouraged people to come and chat with me whilst I was there, I caught up with Nikola who shared her story. She is 16 years old and was struggling to cope as she had no-one to talk to and felt very isolated. We both chatted for a while to allow Nikola to talk about her experiences, her frustrations and what she needed. We connected Nikola to other young people through our Facebook group and made sure she had the email address for HDYO direct support service.

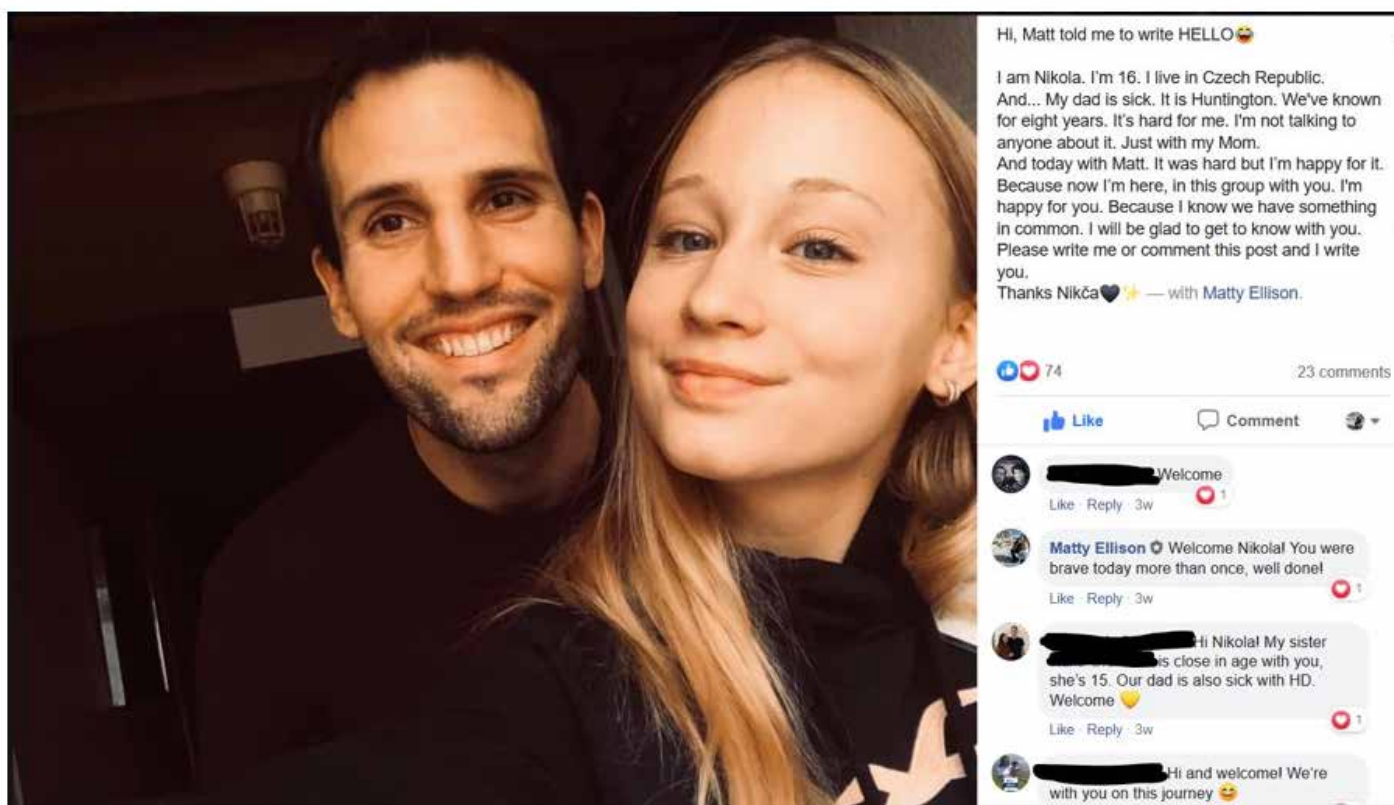
Nikola is now connected with other young people of

HDYO Hangout group on Facebook and has other who understand her situation. Nikola is hoping to come to Congress in Glasgow so that she can meet more young people and get the support from her new virtual community in person.

Nikola even sent me this picture we took when we met and used it to introduce herself to HDYO Hangout group.

Facebook group: <https://www.facebook.com/groups/1950616678545840/>

Direct support Service support@hdyo.org



Meet the Chichester Family:

Bianca (seen in picture with her dad Aaron) began working with the youth service in 2015. She and her siblings were referred by the social worker who was working with the family. Initially, the family wanted some support in talking about HD with their children and providing a safe space for them to process their emotions.

The youth worker helped ensure the family was connected to a full HD team, had information on trials they were eligible for and worked to get the young people more involved in the community. The youth worker and local HD social worker also worked together to ensure the family was able to access all the services available to them since Aaron is a proud veteran of the US Army.

In 2018, Bianca was ready to attend the HDYO North American Camp. Bianca had expressed some interest in previous years, but stated she wasn't quite ready to dive into a 5-day trip focused around HD. When she decided she was ready and attended, she flourished being able to open up and share her experiences with her peers.

Although dad was moved to a nursing home some years ago, the youth worker and Bianca surprised him this past June with Bianca in her cap and gown since he was unable to attend the ceremony.

The amount and type of support the family has wanted has fluctuated over the years, but they know the HDYO Youth Service will be there when they needed it and will coordinate with the other providers that were administering care.



Background:

The Youth Service, since its inception in 2014, has had continuous growth each year. HDYO US Youth Service is managed by Chandler Swope, Director of Youth Services.

Stats:

Data base currently sits at 784 individuals

We gained 180 new contacts in 2019.

Genetic testing, general resources, HD 101 info, North American Camp and getting connected into the community are the main things young people, parents and professionals are reaching out for.

Impact:

In 2019, Chandler provided over 600 hours of individualized support to young people, families and clinicians.

Chandler helped connect young people with University of Iowa when they expressed interest in the relaunch of the Predict Study.

Chandler helped connect young caregivers to local resources – the local HD Social Worker, HD clinic or other support to find appropriate resources.

Number of Contacts (784 Total)

Continued growth of service

10% increase in referrals from 2018, with a number of those coming from our nine new referral partners

Strong partnerships and increased collaboration

Bi-weekly case consultation with Vanderbilt Interdisciplinary Team

Case consultation with HDSA youth workers

Quarterly supervision calls with global youth work partners

Sharing training programs for new social workers within advocacy organizations

Focus on service delivery at the point of need

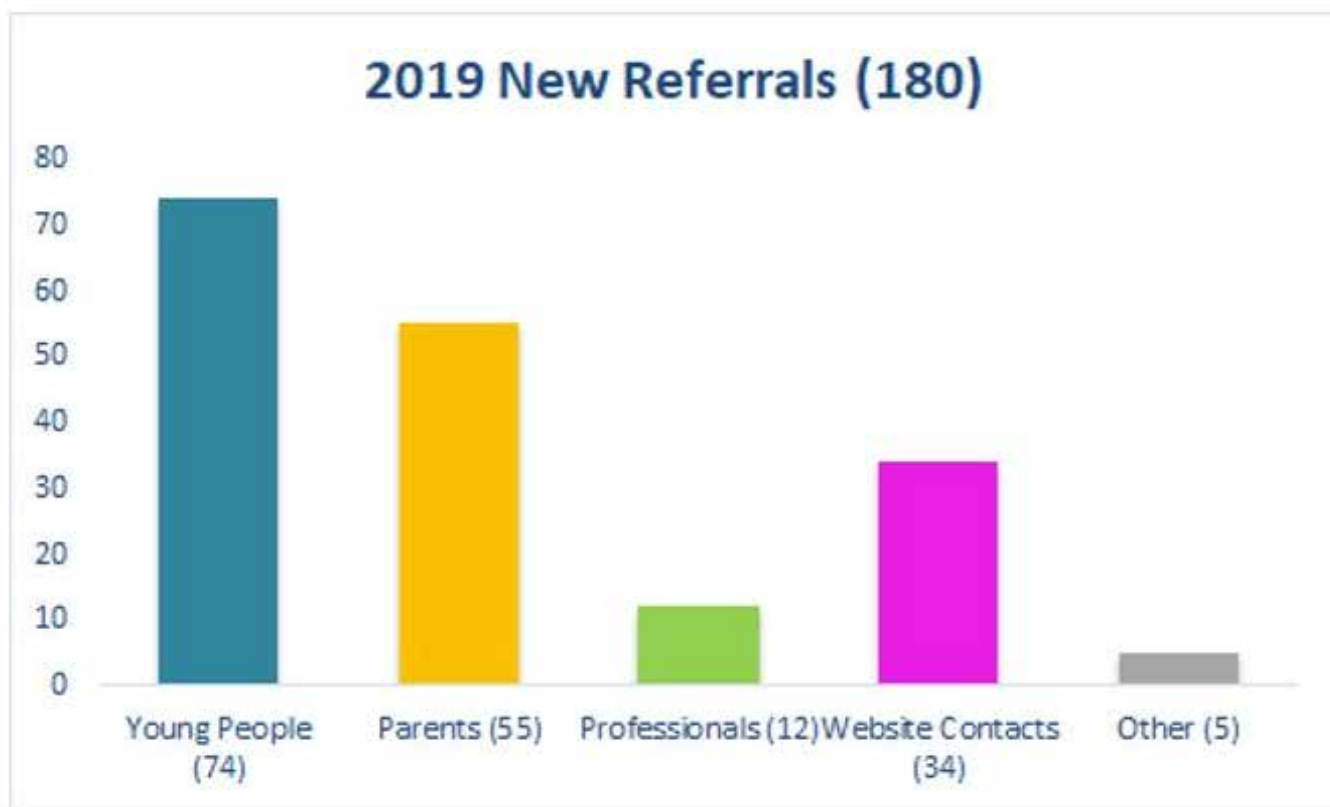
Service users can contact the service using their preferred method of communication at a time when

they need help. Although we provide check in and follow up, most young people want to live their lives and reach out when they have a question or need support. We work with a variety of cases, from those that require intense weekly intervention to those who reach out once over the year for short-term support.

Expanding professional practice to support young people locally

In 2019 we have had an increase in the number of non-HD specific organizations asking for training and resources to support young people with Juvenile onset Huntington's Disease (JoHD).

We are currently working with two schools, two children's hospitals and one nursing home.



HDYO has been hosting the North American HD Youth Camp since 2015. The aim of camp is to connect young people impacted by HD, give them respite from caregiving activities, provide a place to learn about HD and get professional and peer support. Camp is made up of educational sessions, time for individual and group support, as well as fun and games. This year's camp was held at Camp Cedar Glen in Julian, CA. Camp is the largest event (to-date) that HDYO hosts and is an important event for young people to connect face to face. The bonds from camp last a lifetime.

92 applicants (20% increase from 2018); 42 campers attended

50% of campers were under the age of 17 and only four were returning campers

48% of applicants/campers were brand new to HDYO

Post-camp we've done ongoing work with 10 campers via virtual support and connecting them to local resources

"It helped educate me about HD. It helped me learn to be more comfortable with talking about HD. It allowed me to meet some amazing people and make amazing friends. It provided me with the opportunity to ask questions that I wanted to and talk about HD. It was very therapeutic being able to get my stresses and worries off my chest. And over all was an amazing experience." – Camper, 16

"Camp helped me open up and create bonds! The people I met were so supportive and amazing! I talked about things that I never really said out loud and hearing other people's stories made me feel like I wasn't alone!" – Camper



End-of-life planning

We developed a new article on end-of-life planning, which is live on the Young Adult section of hdyo.org. This article looks at what to consider when someone is coming to the end of their life. Although this is very difficult and emotional topic, it is important for HDYO that families have access to trustworthy and factual information that helps them plan ahead for their HD journey.

<https://en.hdyo.org/you/articles/594>

have access to this amazing resource.

<https://en.hdyo.org/land/>

Video Subtitles

We've also been providing our videos with subtitles in multiple languages. These subtitles are now live on our YouTube channel with most of our popular videos now having 4-5 subtitles languages options. Check them out at <https://www.youtube.com/user/HDYOFeed>

Things to consider for those adopting a child at risk of HD

Adopting a child is a major life event and one that takes a considerable amount of time and preparation. When the child you are adopting is at risk of HD, there are even more things that prospective parents need to learn and take into consideration. This article contains some useful insight to help prospective parents with their decisions. This article is live on hdyo.org Parents section.

<https://en.hdyo.org/par/articles/593>

Webinars

We introduced monthly webinars this year, with topics including having children, genetic counselling and research updates from pharma companies and Enroll-HD. These were recorded and are now available on our YouTube and Facebook channels.

Are there topics or people you would like to see appear in our webinar series? Contact Matt@hdyo.org with your suggestions.

Translation upgrade

HDYO Land

Our translations team been working to get our fabulous resource for children, HDYO Land, into additional languages. This project will go live in early 2020 on HDYO Land, allowing a larger audience to

HDYO Heroes

This year we launched HDYO Heroes, a small programme designed to encourage young people to engage with HDYO and the HD community and receive rewards for their efforts. We also updated this programme after launch to become more accessible for young people. Check out HDYO Heroes here: <https://en.hdyo.org/eve/articles/583>

Coping with your results video

We produced a video where we interviewed eight young adults who had been tested for HD, discussing how they have coped since their results. We had both positive and negative participants from various countries, all providing insight into coping with results years afterward. This is an underserved area in terms of resources and knowledge, so felt it was important to put this video out there for young people considering getting tested. See the video here: <https://www.youtube.com/watch?v=BOPotvHitcc>

Donation page revamp and feedback video

We improved the look of our donation page to make it easier and to encourage more donations, providing simple and clear facts to show our work and created a short video using feedback from young people and families who have used our services. Donations have increased this year. See the page here: <https://en.hdyo.org/eve/about/587>

Facebook group for parents

After the success of our Facebook group for young people, we launched a group for parents to share concerns or ask for advice from other parents from HD families. So far, we have over 100 in this group, but we expect it to grow. See it here: <https://www.facebook.com/groups/589164611589575/>

Toolkits

Our toolkits contain resources for families and professionals to guide them through specific topics. They include more practical suggestions and shared experiences alongside balanced advice and facts. HDYO will publish two new toolkits in early 2020:

Genetic Testing: A genetic testing guide for young people expands on the Genetic Testing Checklist to provide young people a source of support during the testing process.

School Toolkit: A guide for schools to support young people impacted by HD and those diagnosed with JoHD. The guide provides information to help education professionals understand the impact of HD and how to support the special education needs of someone with JoHD. This was done in collaboration with the global team of youth workers.



7 million views
2,000 member accounts
5,000 newsletter subscribers



12,400 followers
1,089 members of
youth support group
150 members of
parent support group



2 million video views
4,000 subscribers
Educational videos

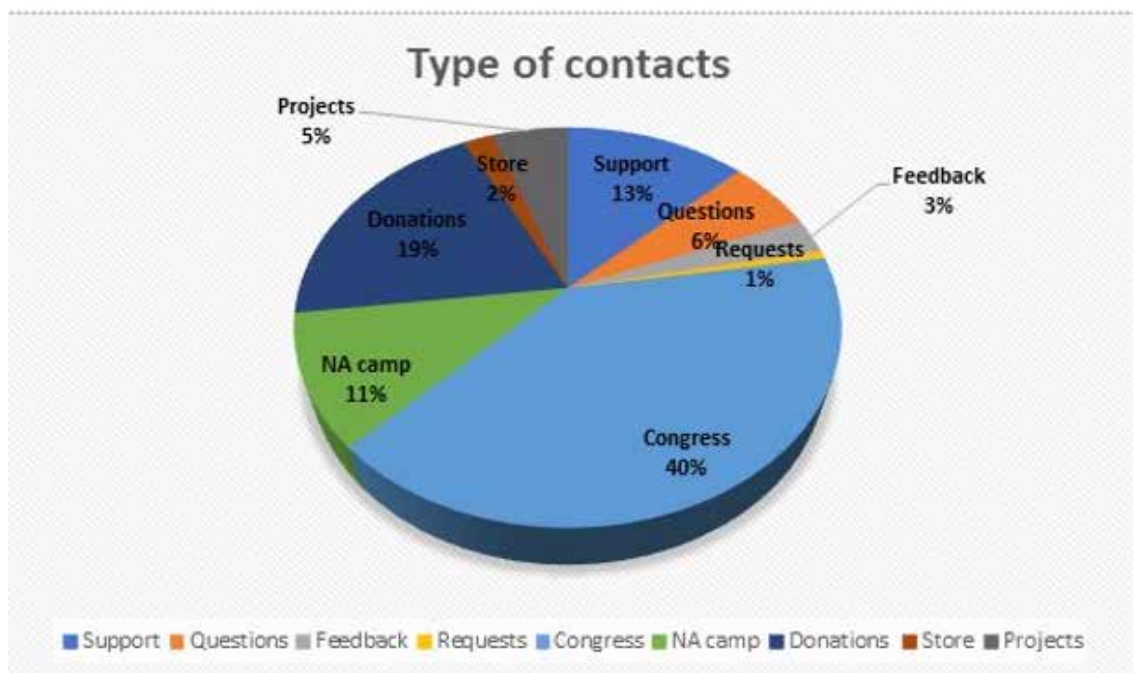
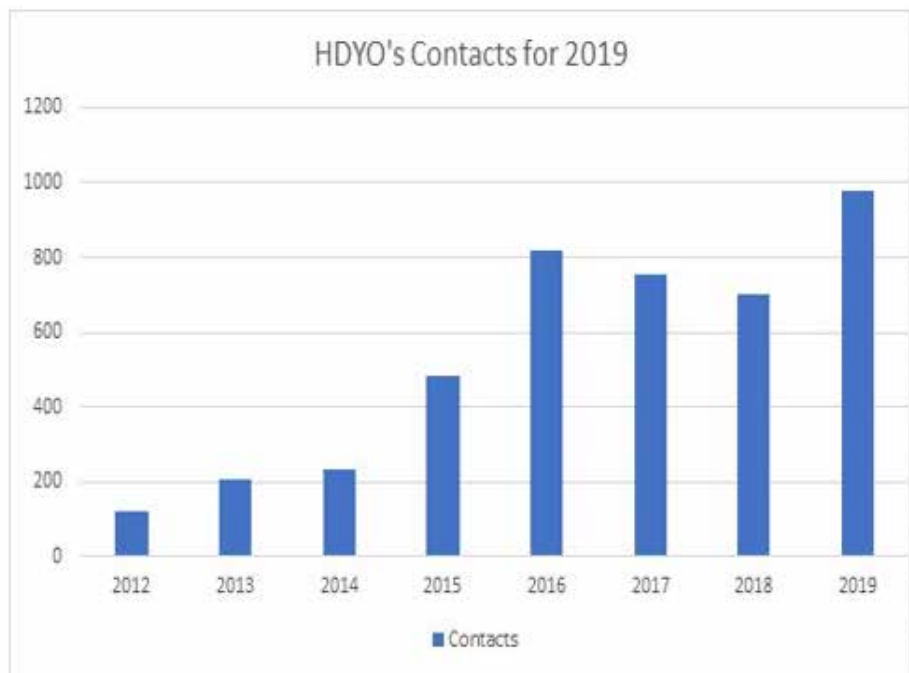


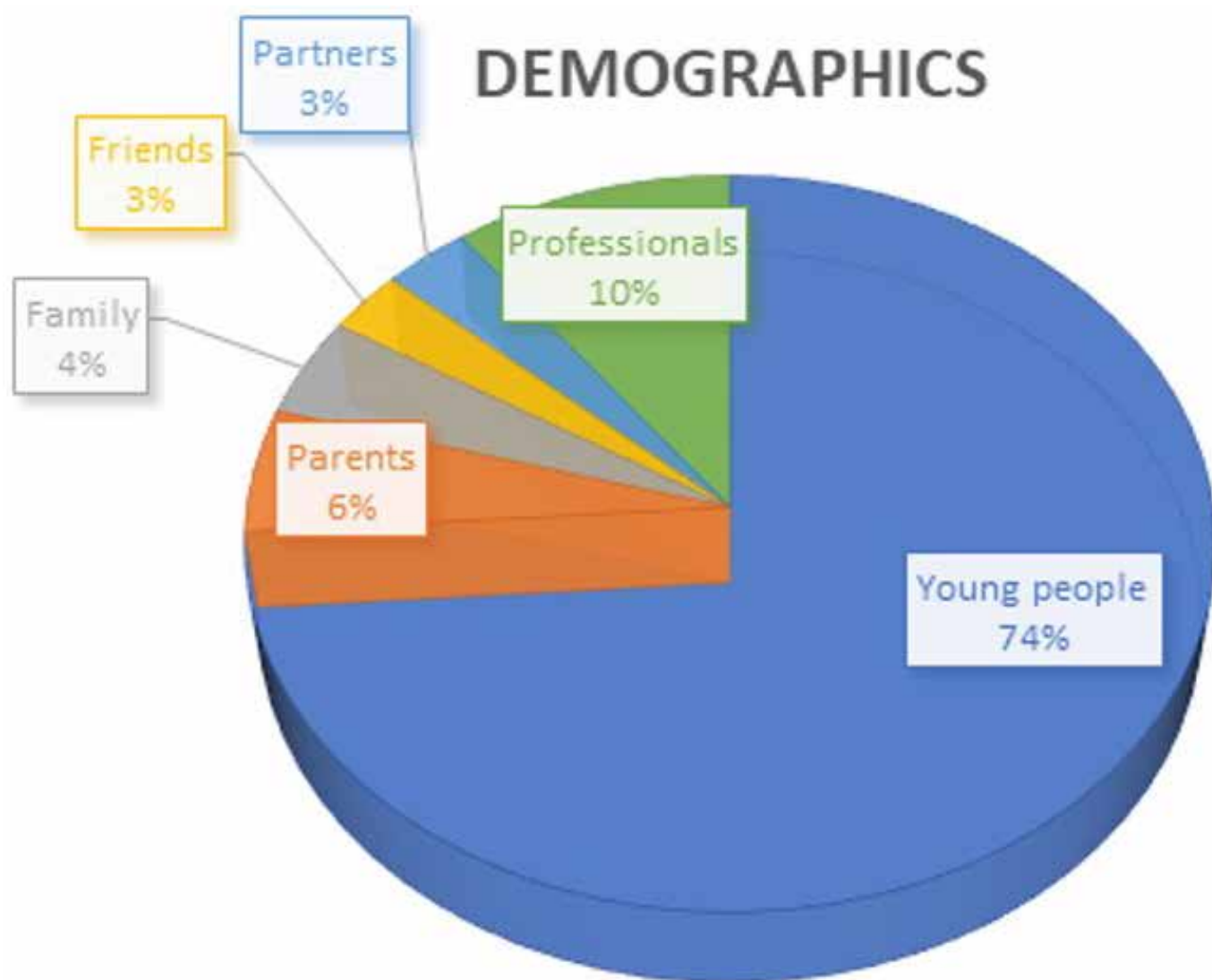
1,164 followers



1,000 followers

We broke our contact record this year with over 976 contacts, beating our previous best in of 819 in 2016.
View the sources of these contacts below.





Translators**German Team**

Michaela
Anne
Clara
Sonja
Clara
Eugen
Julia
Marleen
Janine
Rebecca
Nicole

French Team

Johan
Joel
James
Louis
Jean

Portuguese Team

Sandra
Filipa
Marta
Claudia

Romanian Team

Andrea
Ramona
Dutch Team
Dirk
Dr Lucres
Elisabeth
Monique
Tijl
Wilma
Sanne
Martjin

Social Media

Megan
Rebecca
Doug
Tasha
Gabby
Robert
Jessie
Vicky

Book-keeping

Jessica

Congress**Website & Marketing**

Jaclyn
Megan
David
Caleb
Brad
Hallie

Programme

Emily
Amy-Rose
Nicole
Julianna
Amanda

Scholarships

Brynne
Liam
Amy
Melinda

Volunteers

Jamie
Maggie
Jenna
Dina

Fundraising

Natalie
Jesse
Ashley
Events
Gayle
Karen
Dina
Ross

**North American
Camp**

Lisa
Natalie
Jaclyn
Doug
Erika
Dr Bonnie
Dr Bob
Gwen
Michael
Kris
MaryAnn
Sierra
Jennifer
Corey
Lisa H
Misty
Heather

Sponsors 2019

17

Griffin Foundation

Genentech

CHDI Foundation

Takeda

Georgetown University Medical Centre HD Clinic

Huntington's Disease Association England & Wales

Huntington's Disease Association of Ireland

Deutsche Huntington-Hilfe

HDA Belgium/Franco

Gimbel Family Scholarship

James E "Jake" Hoffman Memorial Fund

Switzerland HD Association



Adam Liebhoff
Alexander Soderlund
Alexandra Bucci
Alan Rotberg
Angelique Marie
Anna Lunsford
Anne Bruns
Annette Carlsson
Anthony & Tammy Miller
Anthony Jackson
Ashley Doak
Barbara Bradshaw
Betsy Ratner
BJ Viau
Bonnie Hennig-Trestman
Brendan Martin
Brianna-Rose Nairn
Brooke Harlowe
Bruce Spiewak
Cari Pitts
Carol Csabafy
Casey Dean Frase
Cassuandra Jones
Catherine Martin
Chad Sipes
Chandler Swope
Charles Austin
Chloe Carr
Chris Brown
Christopher Storgards
CM Moore
Cody Schreier
Corey Janke
Dale & Viki Henry
Dan Murphy
Dan Sharpe
Daniel Mizrachi
Daniella Nelles

Darci Helbling
Dave Hogan
Deborah Sharpe
Deena Schnitman
Denise Seiffer
Donatella Watt
Dr Karen Anderson
Draea Tirschmann
Elizabeth Westbrook
Emilie Liebhoff
Emma Bebbington
Emma Burris
Emma Fickle
Erik Sanches
Fan wu
Felicia Doane
Gail & Bryan Viau
Ganda Setiakurnia
Gil Lewis
Gillian Watts
Gwen Johnson
Harriet Schnitman
Harry Rosenberg
Heidi Klausung Crowl
Holly Matthews
Hope Heller
Jacqueline Riker
Jacqui Harrison
James Lake
James Lieth
Jamie Levey
Jared Piaggione
Jason Potash
Jeanette Clary Braxton
Jeffrey Rotberg
Jesse & Desirae Collett
Jessica Lowe
Jessie Gumbs

Joann Lemasters
Jon Krus
Kaisa Eklumd
Karen Clark
Karen Lenard
Kari Hess
Kari Vinal
Kendell Harrington
Kim Williams
Kirk Oates
Koko Di
Laura Crabtree
Laura Marinari
Leah Ratner
Lindsay Morrison
Lisa Herndon
Lori King
Louisa Winterstein
Maggie Andrews
Maria Arroyo
Mariangels Ferrer
Marilyn Gentner
Marilyn Rye
Mark Murphy
Martina Semacova
Matt Ellison
Matthew Conlon
Megan Carroll
Megan Killegass
Melanie Costa
Melissa Barnes
Michael & Laurel Schnitman
Michal Nowicki
Monica Cazzolli
Nany Watson
Naomi Van Dijk
Nicole Amanda
Nisha Chhabria

Sponsors 2019

15

Oisin Kelly
Patricia Johnson
Patty Romero-Mabry
Peggy Board
Phyllis Gimbel Schnitman
Regina Silver-Koplo
Richard Shiffrin
Robert Macleod
Robyn Oneill
Ryan Bonner
Samuel Karp
Sandra Saenz
Sandra Webb
Sara Hackethal
Savannah Moore
Seth Rotberg
Shannon Dotson
Shiana Darrow
Simone Watt
Solweig Karlsson
Sonia Cruz
Stacey Saladin
Stacey Smith
Stephan Beavis
Steve Ridenhour
Susan Karp
Susan Kelly
Sylvia Pasieka
Sylwia Korszyłowska
Tim West
The Sisu Shop
Tony Miller
Trent Berrier
Vicky Bromby
Definitely missing people

