



dada2

FOUNDATION

Rare Disease
Day 2022

DRIVING TO CONSENSUS

February 17, 2022

Rare Disease Day 2022 is right around the corner!

This is a time of year to help raise awareness and instigate change for people living with a rare disease and their loved ones.

A week ahead of Rare Disease Day, DADA2 Foundation founder Dr. Chip Chambers is teaming up with [Nationwide Children's Hospital](#) as the keynote speaker for their virtual RDD event to speak on the importance of linking patients, families, and physicians to find a cure for this disease.



Dr. Chambers will be joined by other research, patient support, and advocacy experts and we encourage you to sign up and listen in on **Tuesday, February 22 at 4 p.m. ET**. This virtual event is free and open to the public, and includes a special program for children from 4 to 5 p.m.

You can [register for the event and see a full agenda here](#). We'd love to see you there. Don't forget to tell others that might be interested in joining too!

As part of our DADA2 Foundation community, we also wanted to invite you to join us in taking action and raising awareness one week later for Rare Disease Day on Monday, February 28. There are a number of ways you can help below:

- **Take a selfie and post your story about DADA2** on social media on February 28 using the hashtag **#RareDiseaseDay**. This can be on Twitter, Facebook, LinkedIn, or Instagram!
- **Download social media visuals from Rare Disease Day** to post on February 28 with the hashtag **#RareDiseaseDay**. There are even some visuals available in multiple languages.
- **Create a social post about why The DADA2 Foundation is important to you** using the hashtag **#RareDiseaseDay** on February 28.
- Sign up for a [Rare Disease Day virtual event](#) and let others know about The DADA2 Foundation. Many of these are being held around the world so you should be able to find one in your native language.

The [Rare Disease Day website](#) has all sorts of great visual and educational resources for you to share online leading up to the big day. Take a look and find something you'd like to share.

We appreciate all of you and your work to make The DADA2 Foundation community strong. If you're interested in further supporting our work, consider making a [donation today](#).

With much enthusiasm,



Chip Chambers, M.D.
Founder and President, the DADA2 Foundation

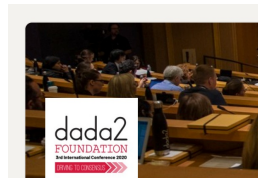
[DONATE to The DADA2 Foundation](#)

We would love to be a part of your year-end giving. Please consider a donation to the DADA2 Foundation if you are able. Otherwise, know that we are immensely grateful for your engagement!

GET INVOLVED


You shop. Amazon gives.

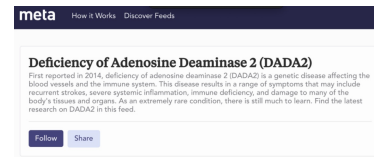
RAISE FUNDS WITH US
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The DADA2 Foundation
Pursuing a cure for DADA2, together.
Non-profit Organization Management · Ne

JOIN THE LINKEDIN PAGE

We've [launched a LinkedIn page](#) to keep you updated.



February 23, 2021
Clinical Features and Outcomes of Childhood Polyarteritis Nodosa: A Single Referral Center Experience.
Modern Rheumatology
Nilsen Tekatze MD, Bana Acar

LATEST DADA2 PAPERS
We're partnering with Meta to [bring the latest papers](#).