

OEBIUS VOICES OCTOBER newsletter

A SEASON OF CHANGE, A SEASON OF VOICES

Dear Friends.

As the seasons change, so too does our community. This summer and fall, we've been making exciting updates at MFOMS. One of the biggest is that all our new programs now begin with Moebius Voices.

Since 2009, together we have been the many faces and many voices of Moebius Syndrome. These new programs are designed to help us share our stories even more widely.

We invite you to follow us on social media and visit our website. Together, we have a bright future ahead.

Take care, Tim Smith





A HOLIDAY GIVING TREE

Share Joy This Holiday Season



This holiday season, we're connecting children and adults with Moebius Syndrome to donors who want to give a little extra.

If even one person is touched through this campaign, it's worth it. It's all about kindness, connection, and giving joy where it's needed most.

Learn More →

PROCLAMATION, PRESS RELEASE & MEDIA TOOLKITS



MSAD 2026 Resources Are Ready

The holidays can get busy, so we've prepared everything early this year:

- Press Release (ready to share)
- Proclamation Toolkit (step-by-step guide for requesting proclamations)
- Media Toolkit (easy instructions to reach local media)

Every outreach matters. Together, we can make MSAD 2026 the most impactful yet.

Download Toolkits



AWARENESS STOREFRONT

Awareness You Can Wear

From Halloween shirts to special 15th Anniversary designs, our storefront is open! Every purchase supports awareness and strengthens our community. All proceeds go to the MSF.

<u>←Shop Awareness Shirts</u> →





* COMMUNITY HIGHLIGHTS

Celebrating Moebius Voices

We have two new programs designed to help your voice be heard.

- Moebius Voices: Interviews with Sophie Taylor, where we highlight the real experiences of people living with Moebius Syndrome. Let us interview you: <u>Learn more</u>
- Moebius Voices: Story Board! Here's how it works: you answer a list of questions in as much detail as you'd like, and we'll turn your responses into your very own story. These stories are more than words on a page — they connect our community, spread awareness about Moebius Syndrome, and remind us that no one is alone in their journey. <u>Learn more.</u>





We're proud to feature talented writers from our community, including Jessica Maher, Gavin Fouché, Theresa McMillian, Kari Storm, Sandy Goodwick, and Leslie Diane Dhaseleer.

Check out their stories here — and if you have one to share, email us at tim@mfoms.org. FRead Stories

GET INVOLVED

Be Part of the Movement

Volunteer, share your story, or join our private Moebius Voices group. <u>—Join Now</u>. Every action helps strengthen our community.



We couldn't have come this far without you. Here's to 16 years behind us and the best is still to come.

Visit Our Website



WWW.MFOMS.ORG