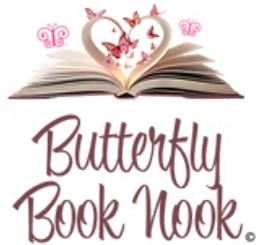


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TSSUS is the largest organization – worldwide – supporting the Turner syndrome community.

The Turner Syndrome Society of the United States: Advancing knowledge, facilitating research, and supporting all those touched by Turner syndrome. [Contact](#)



Hi, Butterflies! **Welcome to the Butterfly Book Nook**, a collection of educational and entertaining books by women with TS. The internet is so very alluring, but sometimes it's great to curl up with a good book. [Visit the Nook](#) and be inspired by the TS experiences of others.



Cindy Scurlock
President & CEO,
M.A., L.P.C.



Visit

TSSUS Turner Syndrome Programs & Services

- [Attend the largest TS meet-your-peers conference in the world](#) • [Join us at local Butterfly Walks](#)
- [Join a private TS Facebook group](#) • [Apply for a scholarship](#) • [Support TS Research](#)

¡Nuevas Guías de Práctica Clínica en Español!

¡Nos complace compartir las Pautas de atención clínica recientemente revisadas para el síndrome de Turner! Esta versión está diseñada para pacientes, familias y cuidadores. También puede acceder a la versión completa para proveedores de atención médica. Explora ambas versiones [aquí](#).



Cuidado Clínico en el Síndrome de Turner

Una versión para pacientes y públicos en general de las Guías de práctica clínica para niños y adolescentes con síndrome de Turner; así como para sus padres, cuidadores y hermanos.

New Clinical Practice Guidelines!

We're excited to share the newly-revised Clinical Care Guidelines for Turner Syndrome! This version is designed for patients, families, and caregivers. You can also access the full version for healthcare providers. Explore both versions [here](#).



Clinical Care in Turner Syndrome

A patient/family version of the Clinical Practice Guidelines For Girls And Women With Turner Syndrome: Data, Patients, Caregivers And Families

California Expands Prenatal Screening Program

The service is offered to all pregnant women in California to detect potential genetic conditions and birth defects in their developing fetus. California is the only state providing voluntary, standard-of-care prenatal screening to all individuals.

[Read more](#).

TSSUS is committed to providing information to those who receive prenatal test results indicating Turner syndrome. Unfortunately, information is limited because there is little research related to outcomes and testing accuracy. We are launching a new TS survey in May to obtain information about TS during pregnancy-early life. The survey announcement will be shared in the newsletter.

Data from TSSUS and UT Health surveys inspires & facilitates patient-powered TS research.

[Read March 2024 Updates](#).



“ TS Quotable ”



"An unforgettable experience that truly helped me to have a feeling of a sense of belonging would be when I was able to attend my first Annual TSSUS conference last year!" **Adriana Garcia**

Thank you, Adriana, for your thoughtful response in our discussion about the potential for those living with Turner syndrome.



Check out the
[May 2025 Issue](#)

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Next Steps

- **Use the Search Bar:** The search function at the top of the page makes it easy to find specific topics.
- **Contact Us for Questions:** If you have any specific questions, feel free to contact us—we're here to help!
- **Connect with the Community:** Gain insights from others by contacting us to connect with a peer via phone, email, or by joining the TSSUS private Facebook group.
- **Browse Resources:** Explore our wide range of educational materials.

Contact TSSUS

Board Member Login

The Turner Syndrome Society's mission is to advance knowledge, facilitate research, and support all those touched by Turner syndrome.



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