

Awareness

Did you know that Turner Syndrome occurs in 1 out of every 2,000 girls?

Treatments and early Interventions are critical to a young girls future growth & development

Early diagnosis and access to care is essential, and we need your help!



Our Mission in Action!

The goal of the Turner Syndrome Foundation (TSF) is to support research initiatives and develop educational programs to increase professional awareness and enhance medical care of those affected by Turner Syndrome. Early diagnosis and comprehensive treatments over the lifespan will lead to a brighter and healthier future for all young girls and women with Turner Syndrome.

TSF is a registered 501 (c)(3) organization



Turner Syndrome Foundation

P. O. Box 726
Holmdel, NJ07733

(732) 847-3385 (800) 594-4585

Turner Syndrome
Awareness Month
FEBRUARY



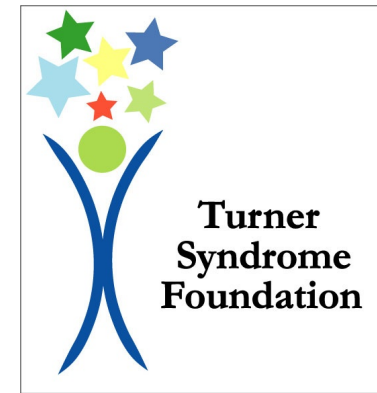
Collaborate

www.TurnerSyndromeFoundation.org

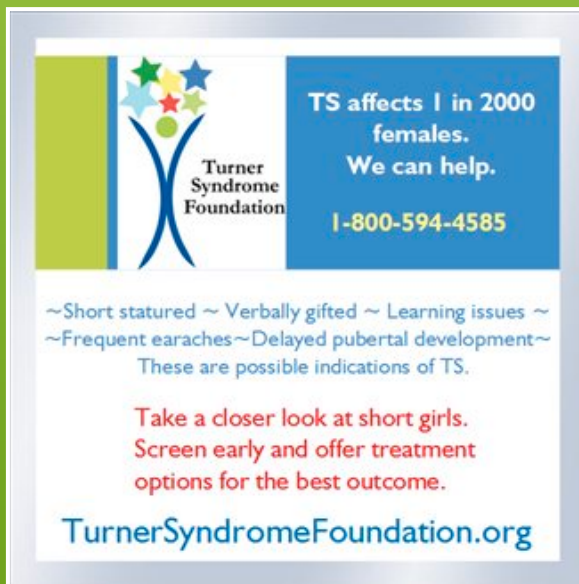
Collaborating for a Cause

Turner Syndrome Awareness Month is February

Thanks to our team of advocates, Turner Syndrome may finally get the attention it needs to identify babies earlier in life. You can help us reach important stakeholders including; administrators, institutions, healthcare providers, philanthropy and consumers.



FEBRUARY - TURNER AWARENESS



TS affects 1 in 2000 females. We can help. 1-800-594-4585

~Short statured ~ Verbally gifted ~ Learning issues ~
~Frequent earaches~Delayed pubertal development~
These are possible indications of TS.

Take a closer look at short girls. Screen early and offer treatment options for the best outcome.

TurnerSyndromeFoundation.org

There is an opportunity to collaborate with your organization on this health initiative to spread the word about this often under-diagnosed and overlooked condition! Here are some of the ways you can get involved:

- Place a banner on your website
- Link with our website
- Post an awareness article in your newsletter
- Host an awareness event

If you would like to get involved, please contact us at:

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