

February is Turner Syndrome Awareness Month



WHAT IS TURNER SYNDROME?

Turner syndrome (TS) is a chromosomal condition that affects girls and women. Females typically have two complete X chromosomes (the sex chromosomes). Turner syndrome is caused by the absence of all or part of the second X chromosome in some or all of the cells of the body.

Early diagnosis is vital so that treatment can begin for any TS related health issues.

- 🦋 **Short Stature** of unknown cause
- 🦋 **Kidney irregularity** (horseshoe shape, absence, or unusually small)
- 🦋 Unexplained **delayed puberty**/menstruation
- 🦋 **Reduced** fertility
- 🦋 Numerous **moles**
- 🦋 **Educational** difficulties



- 🦋 Droopy **eyelids**, low set/prominent **ears**, narrow **roof of mouth**
- 🦋 **Neck webbing** (from fetal cystic hygroma)
- 🦋 **Hearing Loss** (both conductive and sensorineural)
- 🦋 Recurrent **middle ear infections**
- 🦋 **Excellent** verbal skills/vocabulary

How Common is Turner Syndrome?

TS affects approximately 1 in every 2,000-4,000 live born baby girls. It is the **second** most common genetic/chromosomal disorder in females.

Believe in Miracles!

It is estimated that **98%** of Turner syndrome pregnancies end in miscarriage, many even before diagnosis. Those who are living with TS truly are **miracles!**

Down Syndrome 1 in 700

Turner Syndrome 1 in 2,000 - 4,000

Cystic Fibrosis 1 in 3,500

Did You Know?

TS can be diagnosed during **fetal life**, in **infancy**, during the **late pre-teen** period (8-12 years), or in **late adolescence/early adulthood**. A simple blood test is used to determine the karyotype (chromosome makeup).

FOR MORE INFORMATION VISIT US AT
WWW.TURNERSYNDROME.ORG OR CALL:
800-365-9944

SOME HEALTH ISSUES RELATED TO TS

