

About Turner Syndrome

What is Turner Syndrome?

Turner syndrome (TS) was discovered in 1928 and defined as a chromosomal condition describing females* with common features, physical traits, and medical conditions caused by the complete or partial absence of the second sex chromosome. TS affects people differently, so each person has unique health needs and characteristics. Early diagnosis is vital to treat TS-related health issues and health maintenance is necessary throughout life.

*Although TS is defined as a female condition, our organization understands males with a 45 X cell line may be diagnosed with TS.

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

- Short Stature (under 5')
- Unexplained delayed puberty/menstruation
- Cardiac issues & Kidney irregularities
- Non-functioning ovaries/reduced fertility
- Numerous moles
- Droopy eyelids, down-turned eyes, low-set, prominent ears, narrow roof of mouth
- Neck webbing
- Hearing loss (both conductive and sensorineural)
- Recurrent middle ear infections



- Hip dysplasia
- Upturned fingernails/toenails
- Educational and behavioral difficulties
- Slower Processing speed
- Repetitive thoughts
- Anxiety/Depression
- Excellent verbal skills/vocabulary
- Educational and behavioral difficulties
 - Slower Processing speed
 - Repetitive thoughts
 - Anxiety/Depression

Believe in Miracles! Approximately 98% of Turner syndrome pregnancies end in miscarriage, many even before diagnosis. Those who are living with TS truly are miracles!



TS can be diagnosed

- During fetal life
- In **infancy**
- During childhood
- During late pre-teen period (8-12 years)
- Late adolescence/ adulthood.

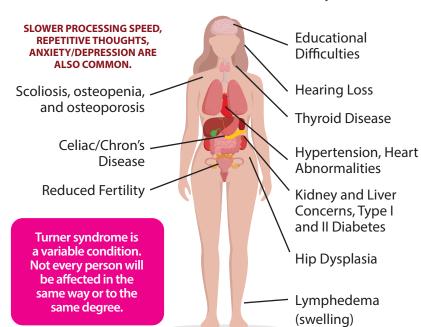
A simple blood test is used to determine the karyotype (chromosome makeup). Turner Syndrome: 1 in 2,000 - 4,000

Cystic Fibrosis: 1 in 3.5k

Down Syndrome: 1 in 700

How Common is Turner Syndrome?

Some Health Issues Related to Turner Syndrome





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