Schwannomatosis (SWN)

is an umbrella term for a group of genetic conditions that cause tumors to grow on nerves throughout the body.

There are many types of schwannomatosis,

which are referred to by their specific gene variant. So far, we know of:

- NF2-related schwannomatosis
- SMARCB1-related schwannomatosis
- LZTR1-related schwannomatosis
- 22q-related schwannomatosis
 schwannomatosis NOS
- (not otherwise specificed) schwannomatosis NEC
- (not elsewhere classified)

Schwannomatosis is characterized by the development of benign tumors called schwannomas usually on Spinal peripheral nerves or vestibular (hearing) nerves.



Clin 70,000 excluding NF2-related SWN Schwannomatosis is caused by pathogenic variants in genes located on chromosome

Approximately 1 in

Y 🕜 `

diagnosed with SWN

will one day be

births

Signs and symptoms of **SWN** usually develop during the teen to **adulthood years**, though they can **OCCUP at any time**.

The most common symptom is **Chronic pain** which can occur

anywhere in the body.

It is believed to be caused at least in part by **schwannomas** pressing on nerves.

Pain management is usually an integral part of care.

Schwannomatosis affects all populations, genders, and ethnicities equally.

THERE IS NO CURE FOR SCHWANNOMATOSIS YET but promising advancements in research are underway.

We're here to help. For more information on schwannomatosis, contact the Children's Tumor Foundation at 1-800-323-7938 or go to ctf.org

