Gastrointestinal Stromal Tumor Causes, Risk Factors, and Prevention

Risk Factors

A risk factor is anything that affects your chance of getting a disease such as cancer. Learn more about the risk factors for gastrointestinal stromal tumors.

- Gastrointestinal Stromal Tumor Risk Factors
- What Causes Gastrointestinal Stromal Tumors?

Prevention

The only known risk factors for gastrointestinal stromal tumors (GISTs) – older age and certain rare, inherited genetic syndromes – cannot be changed. There are no known lifestyle-related or environmental causes of GISTs, so at this time we do not know of any way to protect against these cancers.
But having a risk factor, or even several, does not mean that a person will get the
disease. And many people who get the disease may have few or no known risk factors.

Currently, there are very few known risk factors for gastrointestinal stromal tumors
(GISTs).

**Being older**

These tumors can occur in people of any age, but they are rare in people younger than
40 and are most common in people older than 50.

**Inheriting certain genetic syndromes**

Most GISTs are sporadic (not inherited) and have no clear cause. In rare cases, though,
GISTs have been found in several members of the same family. These family members
have inherited a gene mutation (change) that can lead to GISTs.

**Primary familial GIST syndrome:** This is a rare, inherited condition that leads to an
increased risk of developing GISTs. People with this syndrome tend to develop GISTs
at a younger age than when they usually occur. They are also more likely to have more
than one GIST.

Most often, this syndrome is caused by an abnormal *KIT* gene that is passed from
parent to child. This is the same gene that is mutated (changed) in most sporadic
GISTs. (See [What Causes Gastrointestinal Stromal Tumors?](#) People who inherited this
abnormal gene from a parent have it in all their cells, while people with sporadic GISTs
only have it in the cancer cells.

Less often, a change in the *PDGFRA* gene causes this genetic syndrome. (Defects in
the *PDGFRA* gene are also found in a small percentage of sporadic GISTs.)

Sometimes people with familial GIST syndrome also have skin spots like those seen in
patients with neurofibromatosis (discussed below). Before tests for the *KIT* and
*PDGFRA* genes became available, some of these people mistakenly were thought to
have neurofibromatosis.

**Neurofibromatosis type 1 (von Recklinghausen disease):** This condition is caused
by a defect in the *NF1* gene. This gene change may be inherited from a parent, but in
some cases the change occurs before birth, without being inherited.

People affected by this syndrome often have many benign (non-cancerous) nerve
tumors, called **neurofibromas**, starting when they are young. These tumors form under the skin and in other parts of the body. These people also typically have tan or brown spots on the skin (called **café au lait** spots).

People with NF1 have a higher risk of GISTs (most often in the small intestine), as well as some other types of cancer.

**Carney-Stratakis syndrome:** People with this rare inherited condition have an increased risk of GISTs (most often in the stomach), as well as nerve tumors called **paragangliomas**. GISTs often develop when these people are in their teens or 20s. They are also more likely to have more than one GIST.

This syndrome is caused by a change in one of the **SDH** (succinate dehydrogenase) genes, which is passed from parent to child.

**References**


What Causes Gastrointestinal Stromal Tumors?

Researchers do not know exactly what causes most gastrointestinal stromal tumors (GISTs). But great progress has been made in learning how certain changes in DNA can cause normal cells to become cancer cells.

DNA is the chemical in our cells that makes up our genes, which control how our cells function. We usually look like our parents because they are the source of our DNA. But DNA affects more than just how we look.

Some genes control when cells grow and divide into new cells:

- Certain genes that help cells grow, divide, and stay alive are called oncogenes.
- Genes that normally keep cell growth under control, repair mistakes in DNA, or cause cells to die at the right time are called tumor suppressor genes.

Cancers can be caused by DNA changes that keep oncogenes turned on, or that turn off tumor suppressor genes. These types of gene changes can lead to cells growing out of control.

Gene changes that can lead to GISTs

The gene changes that lead to most GISTs are now understood, but it’s still not clear why these changes occur. There are no known lifestyle-related or environmental risk factors for GIST. Some of the gene changes that lead to GISTs might have causes that haven’t been found yet, but many of these changes may just be random events that sometimes happen inside cells that unfortunately lead to cancer.

A small number of families have GISTs that are caused by a gene mutation passed down from parent to child. (See Gastrointestinal Stromal Tumor Risk Factors.) But most gene mutations related to GISTs are not inherited. These changes occur for no apparent reason, and are called acquired or sporadic.

Changes in KIT or PDGFRA genes

In most people with GISTs, the cancer cells have a change in the KIT oncogene. This gene directs cells to make the KIT protein (also known as CD117), which causes the
cells to grow and divide.

Usually the \textit{KIT} gene is inactive in \textit{interstitial cells of Cajal (ICCs)}\(^1\), which are the cells in the walls of the GI tract from which GISTs develop. The \textit{KIT} gene is only active if there is a need for more ICCs. But in most GISTs the \textit{KIT} gene is mutated. It is always active, so the cells are always growing and dividing.

In about 5% to 10% of GISTs, the cancer cells have a mutation in the \textit{PDGFRA} gene, which causes the cells to make too much of the PDGFRA protein. This has the same effect on the cells as does KIT.

Most GISTs have changes in either the \textit{KIT} or the \textit{PDGFRA} gene, but not both.

\textbf{Other gene changes}

A small number of GISTs, especially those in children, do not have changes in either of these genes. Many of these tumors have changes in one of the \textit{SDH} genes. Researchers are still trying to determine what other gene changes can lead to these cancers.

As doctors have learned more about the gene and protein changes in GIST cells, they have been able to use this information to help diagnose and treat these cancers. (See \textit{Targeted Therapy for Gastrointestinal Stromal Tumors}\(^2\).)

\textbf{Hyperlinks}

1. \url{www.cancer.org/cancer/gastrointestinal-stromal-tumor/about/what-is-gist.html}
2. \url{www.cancer.org/cancer/gastrointestinal-stromal-tumor/treating/targeted-therapy.html}

\textbf{References}


Can Gastrointestinal Stromal Tumors Be Prevented?

The risk of many types of cancer can be reduced with certain lifestyle changes (such as getting to and staying at a healthy weight, or quitting smoking).

But the only known risk factors for gastrointestinal stromal tumors (GISTs) – older age and certain rare, inherited genetic syndromes – cannot be changed. There are no known lifestyle-related or environmental causes of GISTs, so at this time we do not know of any way to protect against these cancers.

References


Morgan J, Raut CP, Duensing A, Keedy VL. Epidemiology, classification, clinical presentation, prognostic features, and diagnostic work-up of gastrointestinal stromal...


Last Medical Review: December 1, 2019 Last Revised: December 1, 2019

Written by

The American Cancer Society medical and editorial content team (www.cancer.org/cancer/acs-medical-content-and-news-staff.html)

Our team is made up of doctors and oncology certified nurses with deep knowledge of cancer care as well as journalists, editors, and translators with extensive experience in medical writing.

American Cancer Society medical information is copyrighted material. For reprint requests, please see our Content Usage Policy (www.cancer.org/about-us/policies/content-usage.html).