

What is Alpha-1?

What you need
to know about Alpha-1
Antitrypsin Deficiency



ALPHA-1
FOUNDATION

What Is Alpha-1?

Alpha-1 Antitrypsin Deficiency (Alpha-1) is a hereditary condition that is passed on from parents to their children through genes. This condition may result in serious lung disease in adults and/or liver disease in infants, children and adults.

Alpha-1 occurs when there is a severe lack of a protein in the blood called alpha-1 antitrypsin (AAT) that is mainly produced by the liver. The main function of AAT is to protect the lungs from inflammation caused by infection and inhaled irritants such as tobacco smoke. The low level of AAT in the blood occurs because the AAT is abnormal and cannot be released from the liver at the normal rate. This leads to a buildup of abnormal AAT in the liver that can cause liver disease.

What are the most common symptoms of Alpha-1?

Symptoms related to the lungs:

- Shortness of breath
- Wheezing
- Chronic cough and sputum (phlegm) production
- Recurring chest colds

Symptoms related to the liver:

- Eyes and skin turning yellow (jaundice)
- Swelling of the abdomen (ascites)
- Vomiting blood or passing blood in the stool

What are some important facts about Alpha-1?

Alpha-1:

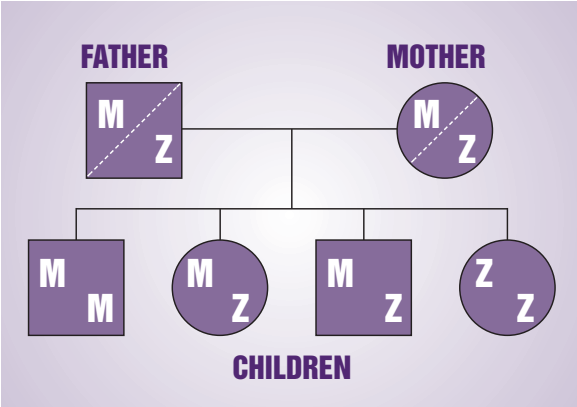
- Is a genetic disorder that leads to low or undetectable levels of AAT
- May cause lung disease in adults
- May cause liver damage that gets worse over time in adults, children and infants
- Often goes undetected for years
- Can be treated, but cannot be cured without a liver transplant
- Is easy to find through a blood test



How is Alpha-1 inherited?

Half of your genes are passed on from each parent. You can inherit Alpha-1 from Alphas (people with two abnormal AAT genes) or carriers (people with one normal and one abnormal AAT gene). Refer to the figure below to see what could happen for children if both parents are carriers. This figure shows the genes that result in normal AAT protein levels (M) and low to undetectable AAT protein levels (Z).

Risks associated with common genetic variants



Normal (MM) • Does not have the disorder; does not carry any abnormal AAT genes.

Carrier (MZ) • Mild to moderate AAT Deficiency — may get disease symptoms and does carry an abnormal AAT gene.

Carrier (MS) • It is unclear whether there is a risk for getting disease symptoms but does carry an abnormal AAT gene (most studies do not show an increased risk for disease).

Alpha-1 (SZ) or (ZZ) • Moderate (SZ) to severe (ZZ) deficiency — could get disease and does carry two abnormal AAT genes.

Alpha-1 (SS) • It is unclear whether there is a risk for getting disease symptoms but does carry two abnormal AAT genes (most studies do not show an increased risk for disease).

Who should be tested for Alpha-1?

Everyone with emphysema, chronic obstructive pulmonary disease (COPD), chronic bronchitis or asthma that is incompletely reversible after aggressive treatment:

- People with bronchiectasis
- Newborns, children and adults with unexplained liver disease
- Individuals with a family history of liver disease
- Blood relatives of a person diagnosed with Alpha-1
- Anyone with panniculitis, a skin disease



Testing for Alpha-1

Testing for Alpha-1 is fairly simple, quick, and highly accurate. It is done through a blood test or a mouth swab test. People at risk for Alpha-1 should be tested because:

- There are treatments and preventive measures that may help you take better control of your health.



- Since it is hereditary, Alpha-1 can be passed on to your children. It should be considered when making decisions about having children and should be discussed with your family members.

The test for Alpha-1 requires a physician's prescription and is usually covered by medical insurance. Confidential testing is available through the Alpha-1 Coded Testing (ACT) Study, through which anyone can have a free, confidential test and get their results. For more information on the ACT Study, call (877) 886-2383.

Informed consent

- Allows you to increase your awareness of Alpha-1 and of your personal risk
- Provides information for future health care decisions
- Allows you to take steps that may slow the progression of Alpha-1
- Assists you and your family in making decisions about work and lifestyle
- Assists those of child-bearing age in understanding risk to children

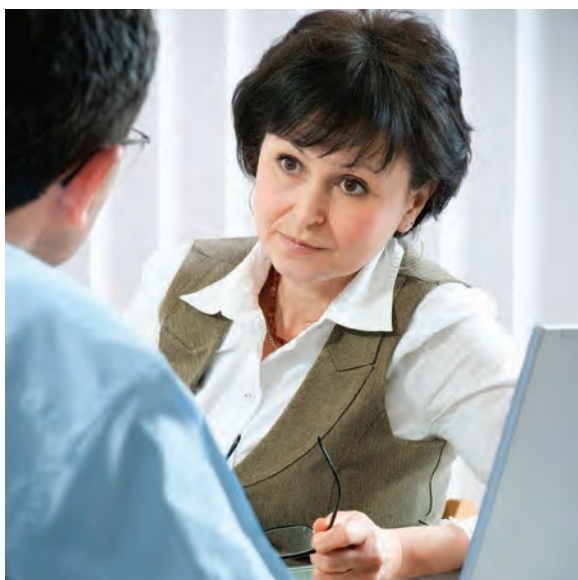
Potential harms

- May be personally unsettling
- May affect your ability to get health and life insurance
- May influence willingness of employers to hire you
- May create stress in your family
- May increase your personal health care costs

(For more information on Informed Consent, see www.alpha1.org/testing-for-alpha-1)

What should I do with the results?

- Contact your physician or primary healthcare provider.
- Create an exercise program (under medical supervision).
- Create a nutrition program (under medical supervision).
- Think about your health behavior (smoking, alcohol use, and excess weight).
- Ask your healthcare provider for a copy of the brochure ***“Guide for the Recently Diagnosed.”***
- Contact the resources listed in this brochure for more information.
- Avoid risk factors:
 - Stop smoking and avoid second-hand smoke as much as possible.
 - Avoid being around dust and fumes.
- Decide whom to inform in your family, and urge anyone who might be affected to get tested also.



RESOURCES

Alpha-1 Foundation

Toll Free: (877) 228-7321 • www.alpha1.org

The not-for-profit Foundation provides resources, education, and information on testing and diagnosis for healthcare providers and people affected by Alpha-1. It funds cutting edge research to find treatments and a cure and supports worldwide detection of Alpha-1.

AlphaNet

Toll Free: (800) 577-2638 • www.alphanet.org

AlphaNet assists patients and families with support, education and strategies to manage their health. It also sponsors clinical trials for Alpha-1 therapies and produces ***The Big Fat Reference Guide to Alpha-1***, a complete guide to understanding, managing and living with Alpha-1, and includes key terms, testing, genetics, and treatment options. It is available through the website or number listed above.

The Alpha-1 Research Registry

Toll Free: (877) 886-2383

www.alphaoneregistry.org

The Research Registry is a confidential database of Alphas and carriers. The Registry gives patients the opportunity to provide information to help advance research on the disorder through questionnaires and clinical trials. It also provides access to experts on Alpha-1 care. People enrolled in the Registry have the ongoing opportunity to participate directly in clinical trials of new therapeutic approaches, in addition to other research opportunities.

Alpha-1 Foundation Patient Information Line

Toll Free: (800) 245-6809

The Patient Information Line is available free of charge to anyone affected by Alpha-1 and provides support to newly diagnosed Alphas, Alphas seeking basic information, or help such as requesting a peer guide, physician referral, requests for literature, etc.

American Association for the Study of Liver Diseases

(703) 299-9766 • www.aasld.org

This is the leading organization of scientists and healthcare professionals committed to preventing and curing liver disease.

Alpha-1 Foundation Genetic Counseling Program

Toll Free: (800) 785-3177

www.alpha1.org/genetic-counseling

The Genetic Counseling Program offers free phone-based confidential information and resources to Alphas, family members and medical professionals. It provides information on the genetics of Alpha-1 and testing options.

Alpha-1 Foundation Support Network

Toll Free: (855) 351-6610

www.alpha1.org/support-groups

The Support Network is a collective of more than 80 Support Groups including four Virtual Support Groups. These groups are committed to improving the quality of life of people and their families affected by Alpha-1 and to extending the mission of the Alpha-1 Foundation.

Alpha-1 Kids

Toll Free: (877) 346-3212

www.alpha1.org/alpha-1-kids

Alpha-1 Kids provides support and information for parents and children with Alpha-1.

American Liver Foundation

Toll Free: (800) 465-4837

www.liverfoundation.org

The Foundation provides information on prevention, treatment and potential cures for liver diseases.

American Lung Association (ALA)

Toll Free: (800) 548-8252 • www.lung.org

The ALA focuses on the prevention of lung disease through educational programs, research and advocacy.

Children's Liver Association for Support Services

Toll Free: (877) 679-8256 • www.classkids.org

This group serves the emotional, educational and financial needs of families and children with liver disease.

Childhood Liver Disease Research Network

www.childrennetwork.org

The Consortium provides support and information for children and families with rare cholestatic liver diseases. For information on liver issues related to AAT, visit this website and click on the Alpha-1 link.



The Alpha-1 Foundation is committed to finding a cure for Alpha-1 Antitrypsin Deficiency and to improving the lives of people affected by Alpha-1 worldwide.

www.alpha1.org
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