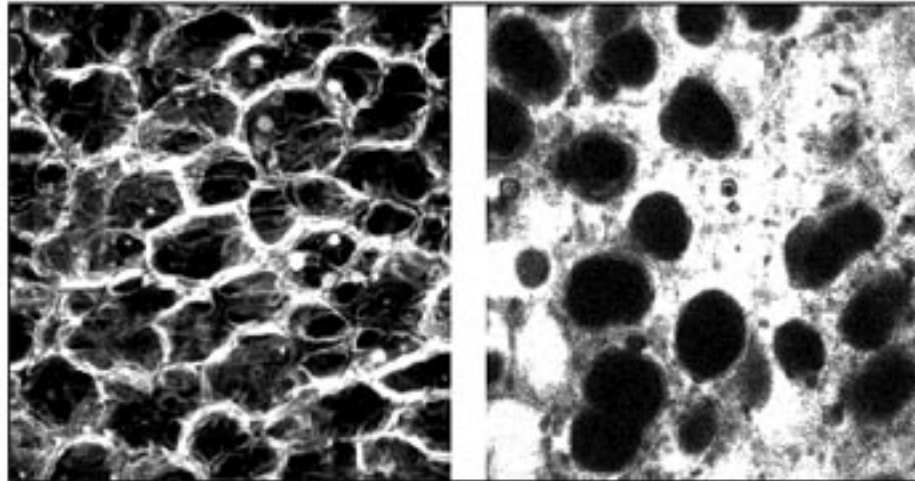


Respiratory Diseases



Laser confocal images of subpleural alveoli of a normal (left) and an edematous, or flooded (right) rat lung. The edema fluid appears white, the alveolar walls gray and airpockets are black. (Gajic and Lee, unpublished). (Reproduced from the web site of Rolf D Hubmayr, M.D., Mayo Clinic and Foundation, Rochester, MN, with permission.)

The respiratory system plays a vital role in delivering oxygen to the body — fuel for all the body's functions. It also removes carbon dioxide waste, eliminates toxic waste, regulates temperature, and stabilizes blood acid-alkaline balance (pH).

The lungs are the largest part of the respiratory system and have both "respiratory" and "non-respiratory" functions. The respiratory function involves gas exchange — the transfer of oxygen from the air into the blood and the removal of carbon dioxide from the blood. Non-respiratory lung functions are mechanical, biochemical, and physiological. The lungs provide a defense against bacterial, viral and other infectious agents; remove various metabolic waste products; control the flow of water, ions, and large proteins across its cellular structures; and manufacture a variety of essential hormones and chemical agents that have important biological roles.

Respiratory diseases can arise from a number of causes, including inhalation of toxic agents, accidents, and harmful lifestyles, such as smoking. Infections, genetic factors, and anything else that affects lung development, either directly or indirectly, can cause respiratory symptoms.

Alpha-1-antitrypsin deficiency

Alpha-1-antitrypsin (AAT) is a protein that protects the body from damage by its immune cells. Deficiency of this protein leaves the lung, and occasionally the liver, vulnerable to injury.

The lung is made of thin outpouchings called alveoli. These contain air, and oxygen travels across their walls into the bloodstream.

White blood cells release elastase, a powerful enzyme that can fight infections. But it can also attack normal tissues. If uncontrolled elastase is released around alveoli, it would destroy their walls and surrounding tissue, leaving areas of trapped air. This abnormal accumulation of air in the lungs is called emphysema and causes shortness of breath.

AAT inhibits elastase around normal tissue. Deficiency is caused by mutations in the SERPINA1 gene, located on chromosome 14. This gene has many different versions (alleles) that produce different amounts of AAT. The M allele produces normal levels of the AAT protein, the S allele produces moderately low levels, and the Z allele produces very low levels.

The alleles are expressed in a codominant manner — that means that a person with MZ has levels of AAT that are between the levels of those people who have alleles MM or ZZ.

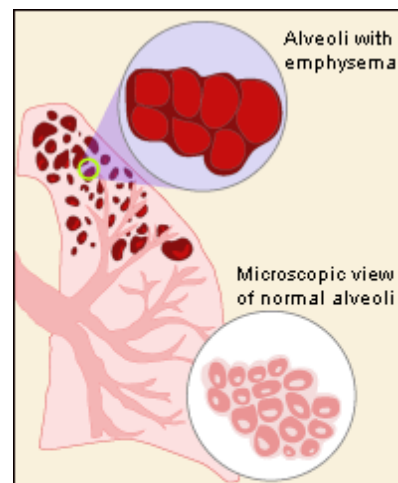
Individuals who have at least one normal allele (MZ or MS) or two copies of S (SS) usually produce enough AAT to protect the lungs but do have an increased risk of lung disease. The risk is particularly high if they smoke. Individuals who inherit the Z allele from each parent (ZZ) have very low AAT and are at a higher risk of developing emphysema and liver disease.

There are over 70 known mutations that occur at the SERPINA1 gene. A common mutation that creates the Z allele involves a switch in amino acids — glutamic acid is replaced by lysine at position 342 (Glu342Lys). The resulting AAT protein cannot

fold properly. This hinders its secretion from the liver (which makes AAT) into the bloodstream (which transports AAT to the lungs). The accumulation of AAT complexes can damage the liver, whereas the lack of AAT fails to stop the destruction of lung tissue.

Treatment of AAT deficiency includes the standard treatment of emphysema (bronchodilators, early use of antibiotics in infections) as well as the more experimental therapies of correcting AAT levels by replacing the protein. Gene therapy to replace the defective SERPINA1 gene with a functional copy is currently being investigated.

However, the most important part of treatment of AAT deficiency is to avoid smoking. Affected individuals are far less likely to develop emphysema if they do not smoke. Not only is smoking a lung irritant, which attracts white blood cells (and therefore neutrophil elastase) to the lungs, it also prevents any AAT that is present in the lungs from working properly.



In AAT deficiency, the walls of lung tissue (alveoli) are broken down. The result is a trapping of air known as emphysema. The damaged lung tissue can no longer transfer oxygen to the bloodstream.

Important Links

Gene sequence

Genome view see gene locations

LocusLink [www.ncbi.nlm.nih.gov/LocusLink/LocRpt.cgi?l=5265] collection of gene-related information

BLink [www.ncbi.nlm.nih.gov/sutils/blink.cgi?pid=21361198&all=1] related sequences in different organisms

The literature

Research articles online full text

Books online books section

OMIM [<http://www.ncbi.nlm.nih.gov/entrez/dispomim.cgi?id=107400>] catalog of human genes and disorders

Websites

Patient information [<http://www.nhlbi.nih.gov/health/public/lung/other/antitryp.htm>] from National Heart, Lung and Blood Institute, NIH

MEDLINEplus [<http://www.nlm.nih.gov/medlineplus/ency/article/000120.htm>] Medical encyclopedia from the National Library of Medicine, NIH

Alpha-1 Association [<http://www.alpha1.org/>] Information and patient support

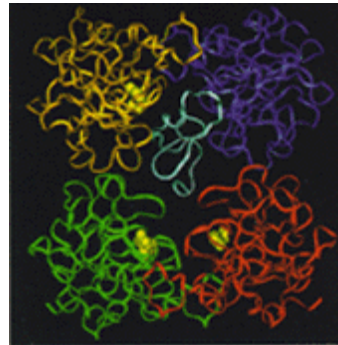
Asthma

Asthma affects more than 5% of the population of the US, including children. It is a chronic inflammatory disorder of the airways characterized by coughing, shortness of breath, and chest tightness. A variety of "triggers" may initiate or worsen an asthma attack, including viral respiratory infections, exercise, and exposure to irritants such as tobacco smoke. The physiological symptoms of asthma are a narrowing of the airways caused by edema (fluid in the intracellular tissue space) and the influx of inflammatory cells into the walls of the airways.

Asthma is a what is known as a "complex" heritable disease. This means that there are a number of genes that contribute toward a person's susceptibility to a disease, and in the case of asthma, chromosomes 5, 6, 11, 14, and 12 have all been implicated. The relative roles of these genes in asthma predisposition are not clear, but one of the most promising sites for investigation is on chromosome 5. Although a gene for asthma from this site has not yet been specifically identified, it is known that this region is rich in genes coding for key molecules in the inflammatory response seen in asthma, including cytokines, growth factors, and growth factor receptors.

The search for specific asthma genes is ongoing. Assisting in this international human effort are model organisms such as mice, which have similar

chromosomal architecture to our chromosome 5 site on their chromosomes 11, 13, and 18. Further study of the genes in these areas (and others) of the human genome will implicate specific genes involved in asthma and perhaps also suggest related biological pathways that play a role in the pathogenesis of asthma.



Tryptase is an enzyme found specifically in mast cells, a type of white blood cells important for fighting infection. It may have a role in causing asthma and other inflammatory disorders. [Reproduced from Pereira, P.J.B. et al. (1998) *Nature* 392, 30-311, with permission.]

Important Links

Gene sequence

Genome view see gene locations

LocusLink [www.ncbi.nlm.nih.gov/LocusLink/list.cgi?Q=asthma&ORG=Hs&V=0] collection of gene-related information

BLink [www.ncbi.nlm.nih.gov/sutils/blink.cgi?pid=4501969&org=1] related sequences in different organisms

The literature

Research articles online full text

Books online books section

OMIM [www.ncbi.nlm.nih.gov/entrez/dispomim.cgi?id=600807] catalog of human genes and disorders

Websites

Global initiative for asthma [www.ginasthma.com:80/] a project conducted in collaboration with the National Heart, Blood and Lung Institute, NIH, and the World Health Organization

National Heart, Blood and Lung Institute [www.nhlbi.nih.gov/health/public/lung/index.htm#asthma] information on asthma

MEDLINEplus [www.nlm.nih.gov/medlineplus/asthma.html] links on asthma compiled by the National Library of Medicine

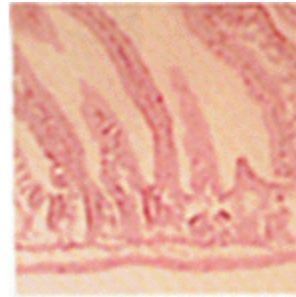
Cystic fibrosis

Cystic fibrosis (CF) is the most common fatal genetic disease in the US today. It causes the body to produce a thick, sticky mucus that clogs the lungs, leading to infection, and blocks the pancreas, stopping digestive enzymes from reaching the intestines where they are required to digest food.

CF is caused by a defective gene, which codes for a sodium and chloride (salt) transporter found on the surface of the epithelial cells that line the lungs and other organs. Several hundred mutations have been found in this gene, all of which result in defective transport of sodium and chloride by epithelial cells. The severity of the disease symptoms of CF is directly related to the characteristic effects of the particular mutation(s) that have been inherited by the sufferer.

CF research has accelerated sharply since the discovery of CFTR in 1989. In 1990, scientists successfully cloned the normal gene and added it to CF cells in the laboratory, which corrected the defective sodium chloride transport mechanism. This technique—gene therapy—was then tried on a limited

number of CF patients. However this treatment may not be as successful as originally hoped. Further research will be required before gene therapy, and other experimental treatments, prove useful in combating CF.



Building mouse models of human disease. Expression of a human cystic fibrosis (CFTR) gene in the gut of a mouse. A human anti-sense probe was used to show human CFTR expressed in the mouse duodenum. [Reproduced with permission from Manson, A.L. et al. (1997) *EMBO J.* 16, 4238-4249.]

Important Links

Gene sequence

Genome view see gene locations

LocusLink [www.ncbi.nlm.nih.gov/LocusLink/list.cgi?Q=cystic%20fibrosis&ORG=Hs&V=0] collection of gene-related information

BLink [www.ncbi.nlm.nih.gov/sutils/blink.cgi?pid=6995996&org=1] related sequences in different organisms

The literature

Research articles online full text

Books online books section

OMIM catalog of human genes and disorders

Websites

Fact sheet [www.nhlbi.nih.gov/health/public/lung/other/cystfib.htm] from the National Heart, Lung and Blood Institute, NIH

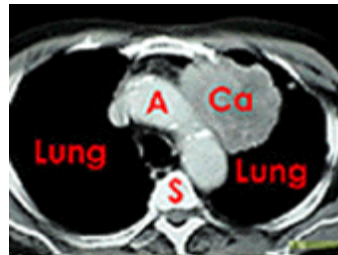
The Cystic Fibrosis Foundation [www.cff.org/] information and links

Lung carcinoma, small cell

In the US, lung cancer is the most common cause of cancer deaths among both men and women. In fact, North Americans have the highest rates of lung cancer in the world. In 1997, some 178,100 new cases were diagnosed, and roughly 160,400 deaths occurred from the disease. Sadly, the 5-year survival rate for persons with lung cancer is only 14%. Since the 1940s, the increase in lung cancer mortality by gender has followed historic patterns of smoking, with a 20-year time lag. About 90% of male lung cancer deaths and 80% of female lung cancer deaths are attributable to cigarette smoking. Although smoking is by far the major risk factor for lung cancer, certain industrial substances, such as asbestos, and environmental factors can contribute.

Small cell lung carcinoma is distinctive from other kinds of lung cancer (metastases are already present at the time of discovery) and accounts for approximately 110,000 cancer diagnoses annually. A deletion of part of chromosome 3 was first observed in 1982 in small cell lung carcinoma cell lines.

As with other cancers, mutations in a variety of molecules (oncogenes and tumor-suppressor genes) that control cell growth and division are observed, and no one mutation is likely to result in cancerous growth. Basic research into the function of these molecules—how and when they play their role—should help the fight against lung, and other, cancers and give clues to find appropriate therapies.



CT scan showing lung cancer.
[Image credit: Pat Connelly,
Miami Valley Hospital, Dayton,
OH, USA.]

Important Links

Gene sequence

Genome view see gene locations

LocusLink [www.ncbi.nlm.nih.gov/LocusLink/list.cgi?Q=lung+cancer&ORG=Hs&V=0] collection of gene-related information

BLink [www.ncbi.nlm.nih.gov/sutils/blink.cgi?pid=4826696&org=1] related sequences in different organisms

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Websites

CancerNet [cancernet.nci.nih.gov/] from the National Cancer Institute, NIH

American Cancer Society [www.cancer.org] research and patient support

Oncolink [oncolink.upenn.edu/] comprehensive cancer information from the University of Pennsylvania