

Respiratory block
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Pathology of Chronic obstructive airway diseases

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Objectives:

- **Chronic Bronchitis**

- a. Define chronic bronchitis.
- b. Describe the pathogenesis and the morphology of chronic bronchitis.
- c. Describe the mechanism of airway obstruction in a patient with chronic bronchitis. Understand that when severe obstruction is present in chronic bronchitis, significant emphysema is nearly always present

- **Emphysema**

- a. Define emphysema.
 - b. Describe the gross and microscopic changes in emphysema.
 - c. Discuss the typical clinical presentation and causes of death.
 - c. Describe the most likely mechanism of emphysema (the protease-antiprotease mechanism).
 - e. Describe the pathophysiologic mechanisms of emphysema
- Explain why emphysema and bronchitis are both considered to be examples of chronic obstructive pulmonary disease (COPD). Compare and contrast the major clinical and functional differences between predominant chronic bronchitis versus predominant emphysema in patients with COPD.
 - Define **Bronchiectasis** and its significance.

Obstructive and Restrictive Pulmonary Diseases

- Diffuse pulmonary diseases are divided into:

1. **Obstructive disease:**

characterized by limitation of airflow owing to partial or complete obstruction at any level from trachea to respiratory bronchioles.

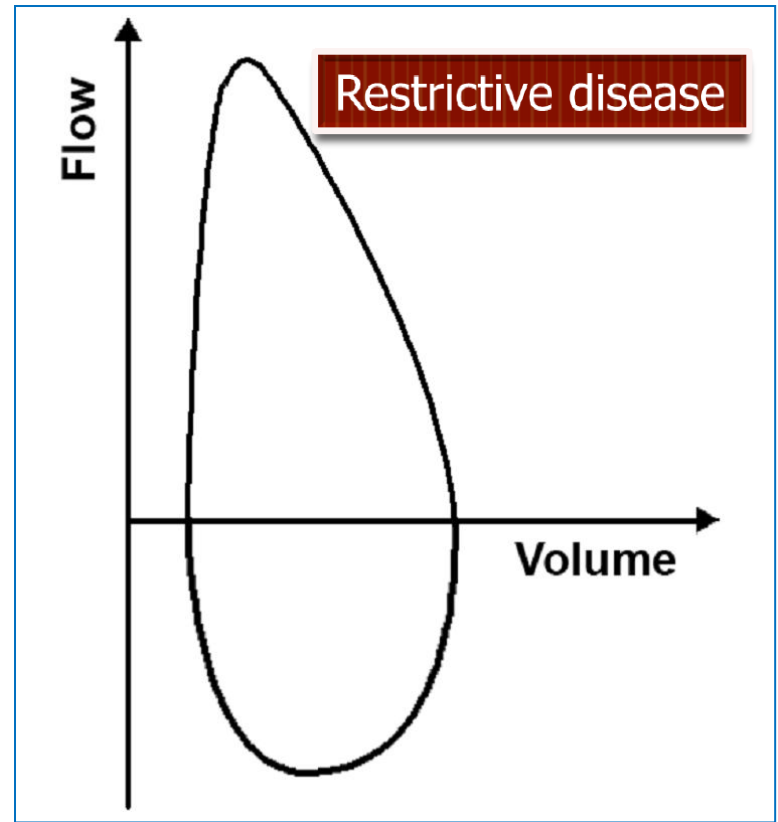
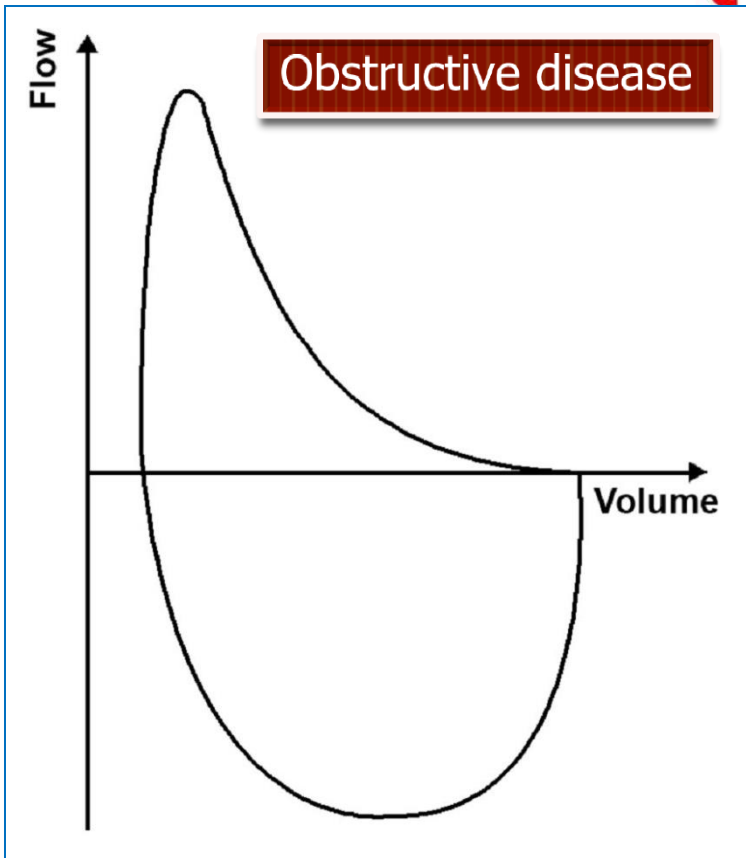
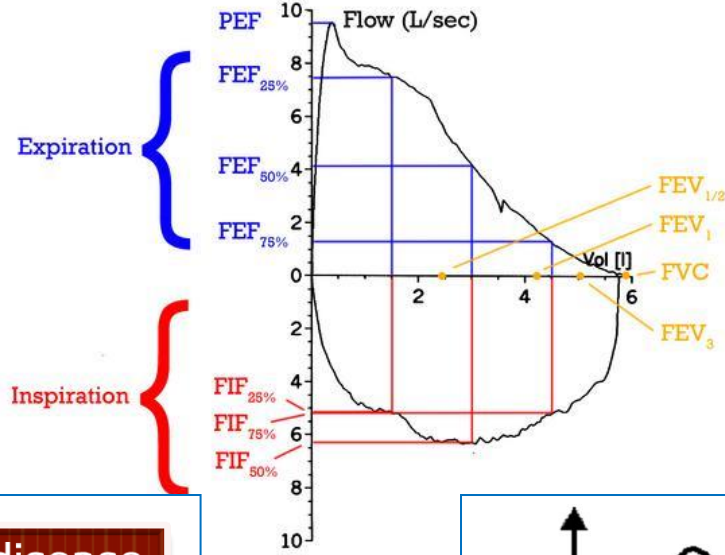
Pulmonary function test: limitation of maximal airflow rate during forced expiration (FEV1).

2. **Restrictive disease:**

characterized by reduced expansion of lung parenchyma with decreased total lung capacity while the expiratory flow rate is near normal.

Occur in:

1. Chest wall disorder.
2. Acute or chronic, interstitial and infiltrative diseases, e.g. ARDS and pneumoconiosis.



Main Categories of (diffuse) Obstructive Disease

- 1) Asthma
- 2) Chronic obstructive pulmonary/airway/lung disease(COPD/COAD/COLD). They are of two types:
 - a) Chronic bronchitis
 - b) Emphysema
- 3) Bronchiectasis

COPD:

Irreversible obstruction to airflow out of the lungs

Cigarette smoking is the principal cause of COPD

Greater than 10% of the population >45 years old has airflow obstruction.

Majority of patients with COPD have both emphysema (air space destruction) and chronic bronchitis

Chronic Bronchitis

Chronic Bronchitis

- Common among cigarette smokers and urban dwellers, age 40 to 65
- The diagnosis of chronic bronchitis is based on clinical features.
- **Persistent productive cough (with sputum) for at least 3 consecutive months in at least 2 consecutive years**
- Can occur in several forms:
 1. Simple chronic bronchitis.
 2. Chronic mucopurulent bronchitis.
 3. Chronic asthmatic bronchitis.
 4. Chronic obstructive bronchitis.

Some patients with a clinical definition of chronic bronchitis have hyperresponsive airways with intermittent bronchospasm..

Chronic bronchitis

Causative factor are:

- Cigarette smoking and pollutants. Most patients are smokers
- Infection
- Genetic factors e.g. cystic fibrosis
- Often, there are features of emphysema as well

Pathogenesis

Chronic irritation of inhaled substances or microbial infection leads to

- Hypersecretion of mucus that starts in the large airways with associated hypertrophy of the sub-mucosal glands.
- As chronic bronchitis persists the small bronchi and bronchioles also get affected.
- Inflammation and irreversible fibrosis may occur in chronically inflamed segmental bronchi and bronchioles.

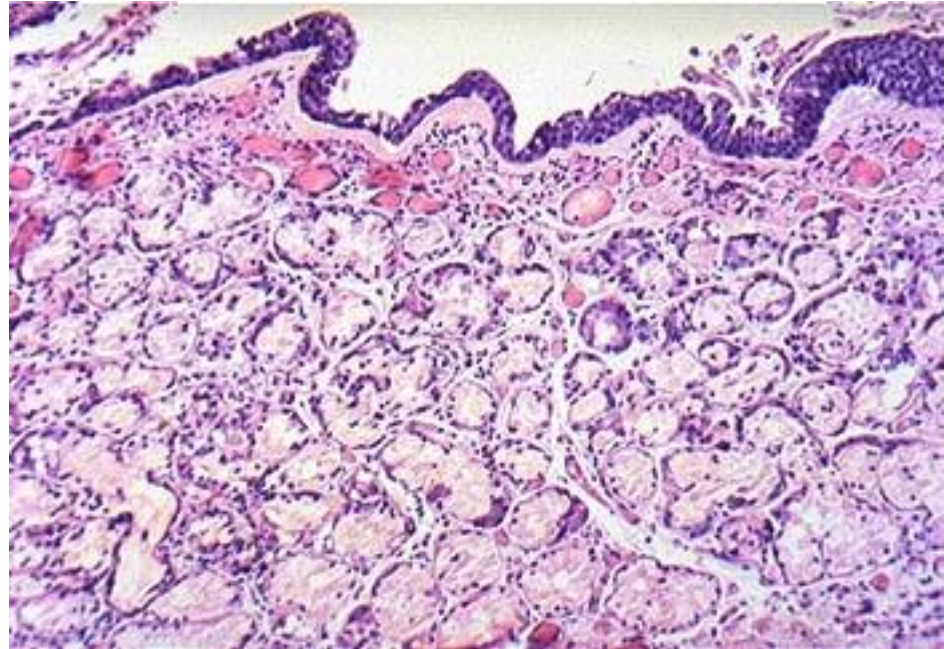
Chronic bronchitis morphology

- Chronic bronchitis does not have characteristic pathologic findings, but is defined clinically as a persistent productive cough for at least three consecutive months in at least two consecutive years.
- In bronchitis the airway mucosa is red and edematous



Chronic bronchitis: morphology

- **Inflammation of airways**, fibrosis and resultant narrowing of bronchioles.
- **Hypertrophy and hyperplasia of mucus producing cells** increased number of goblet cells,
- **Squamous metaplasia** which can progress to dysplasia and even invasive carcinoma.
- **Injury to cilia with** loss of ciliated epithelial cells
- squamous metaplasia.
- Coexistent emphysema.



How do these changes differ from the changes seen in a typical case of allergic asthma?

In typical allergic asthma, which also has mucous gland hyperplasia, the bronchial wall has an inflammatory infiltrate in which eosinophils are prominent. There is also hypertrophy and hyperplasia of smooth muscle cells in asthma.

How do the changes differ from those seen in bronchiectasis?

Infection-related destruction of the bronchial wall is the characteristic appearance of bronchiectasis

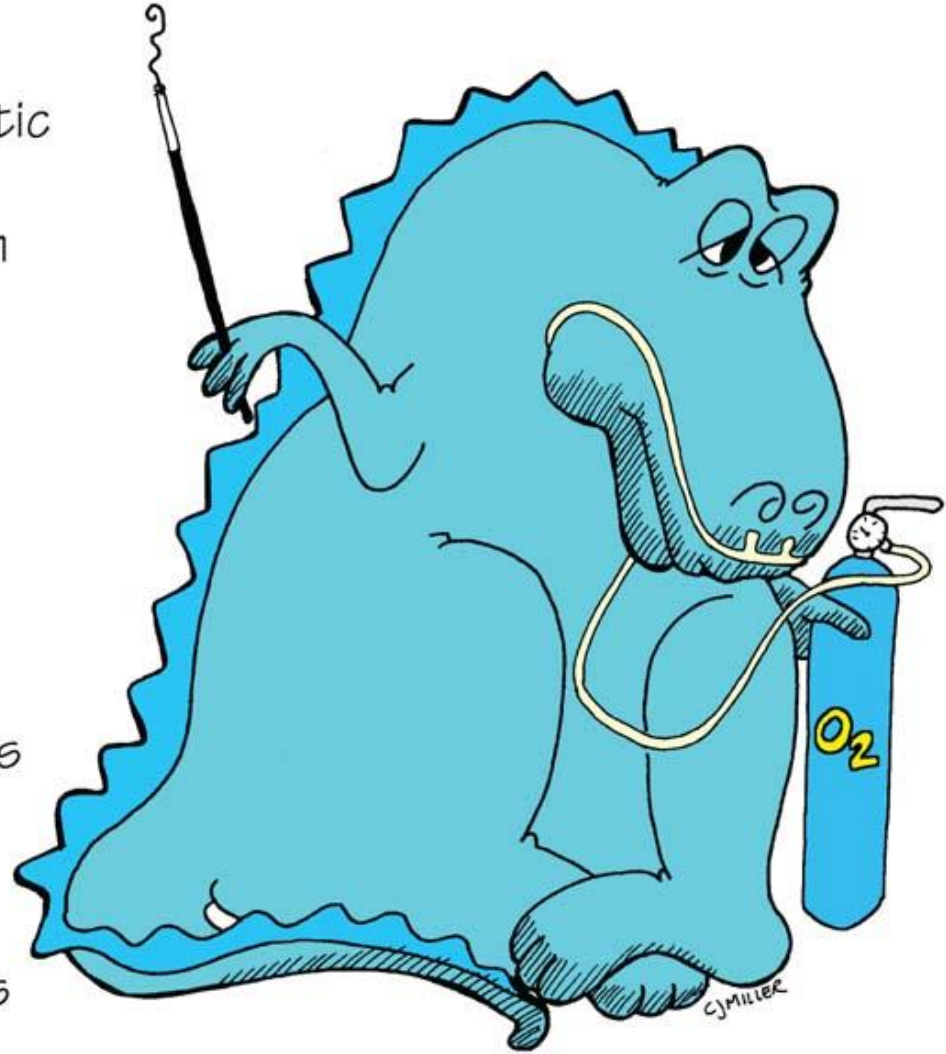
Chronic bronchitis: Clinical Course

- **Prominent cough and the production of sputum.**
- **Hypercapnia, hypoxemia and cyanosis. Patients with severe chronic bronchitis are termed blue bloaters.**
- **Patients can have:**
 - **increased sleepiness due to CO₂ narcosis**
 - **cyanosis due to very poor oxygenation**
 - **elevated red cell counts (secondary polycythemia) as a result of chronic hypoxemia**
 - **Cardiac failure (Cor pulmonale/right heart failure): diseases of the lung or pulmonary vasculature leads to pulmonary hypertension which leads to right ventricular dilation and hypertrophy (right heart failure).**

CHRONIC BRONCHITIS

“BLUE BLOATER”

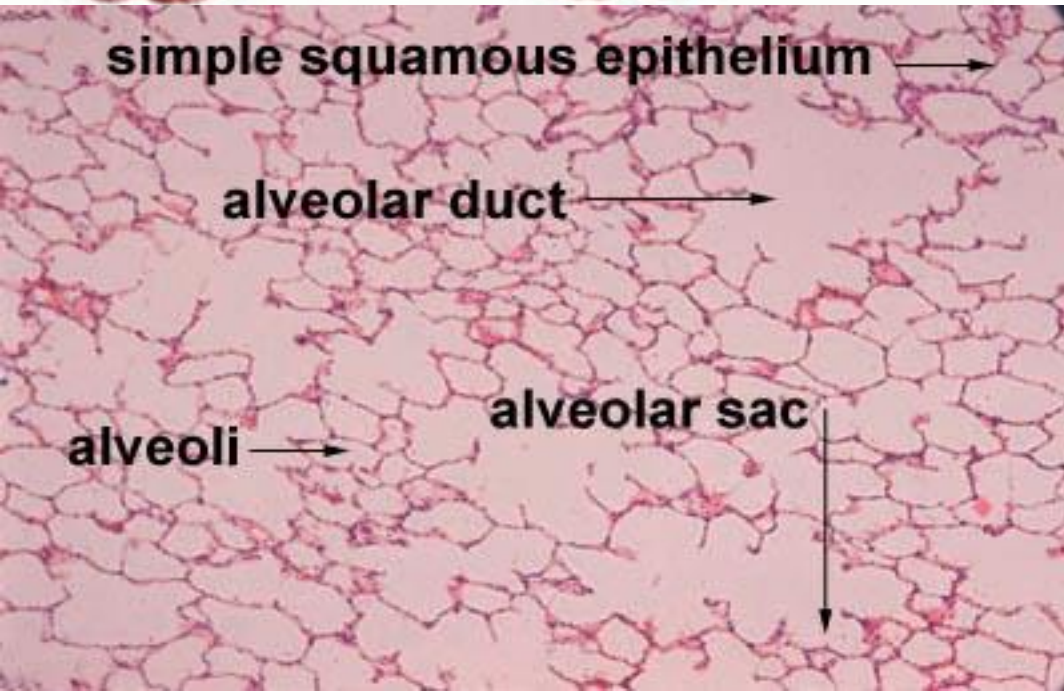
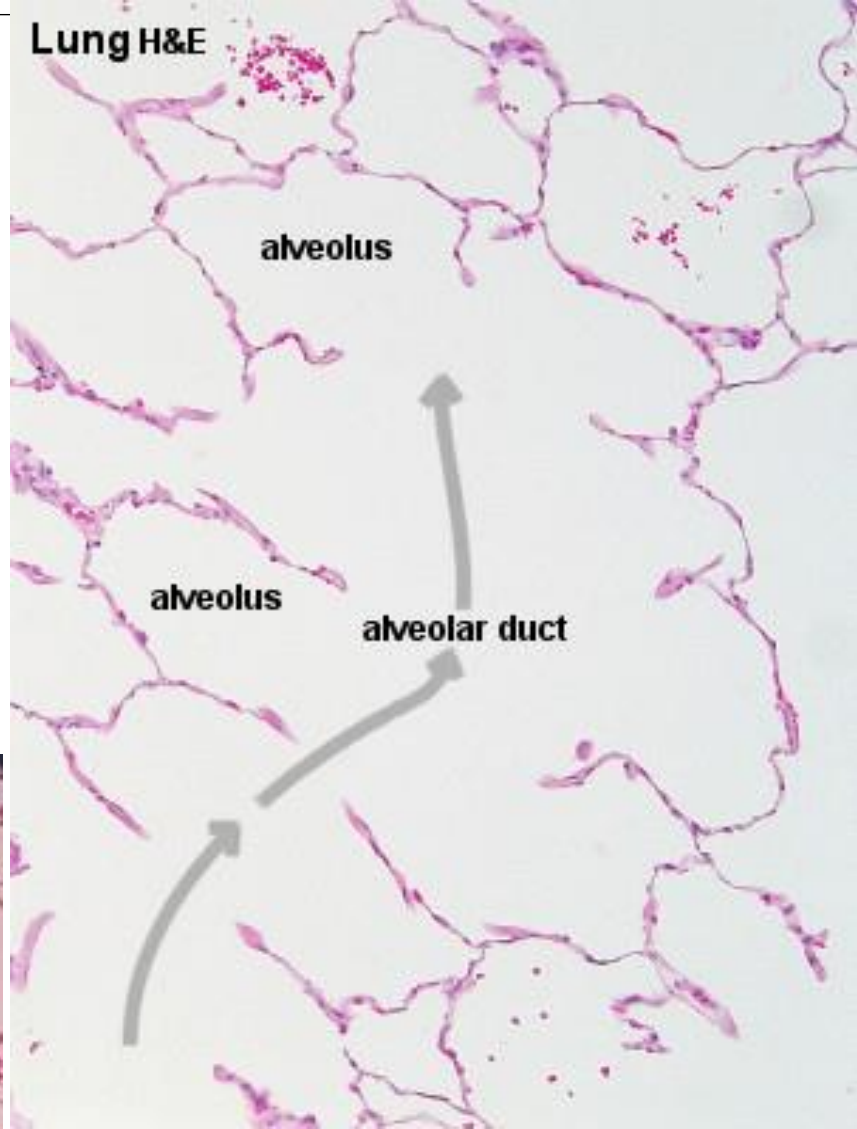
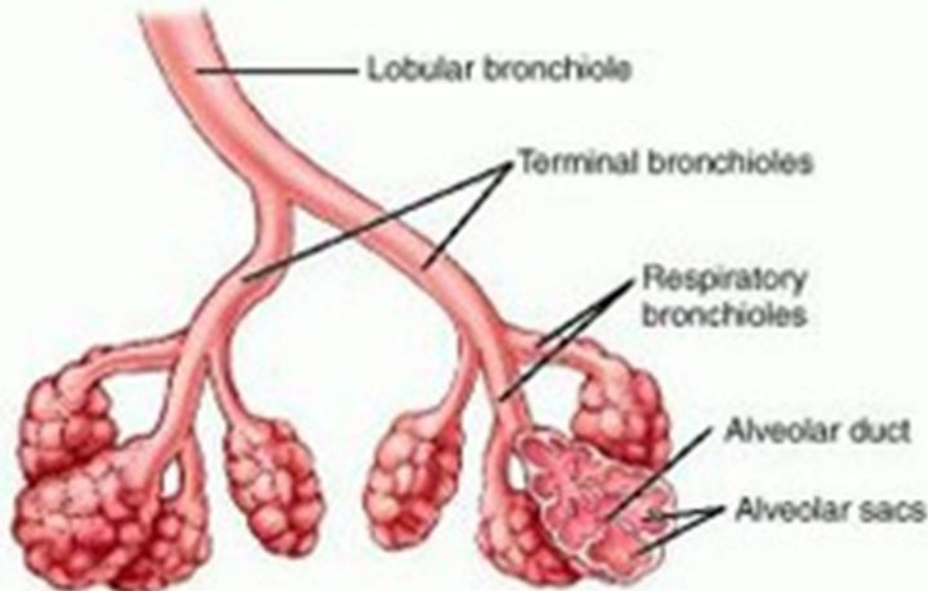
- * Airway Flow Problem
- * Color Dusky to Cyanotic
- * Recurrent Cough & ↑ Sputum Production
- * Hypoxia
- * Hypercapnia (↑ pCO₂)
- * Respiratory Acidosis
- * ↑ Hgb
- * ↑ Resp Rate
- * Exertional Dyspnea
- * ↑ Incidence in Smokers
- * Digital Clubbing
- * Cardiac Enlargement
- * Use of Accessory Muscles to Breathe
- * Leads to Right-Sided Heart Failure: Bilateral Pedal Edema, ↑ JVD



Emphysema

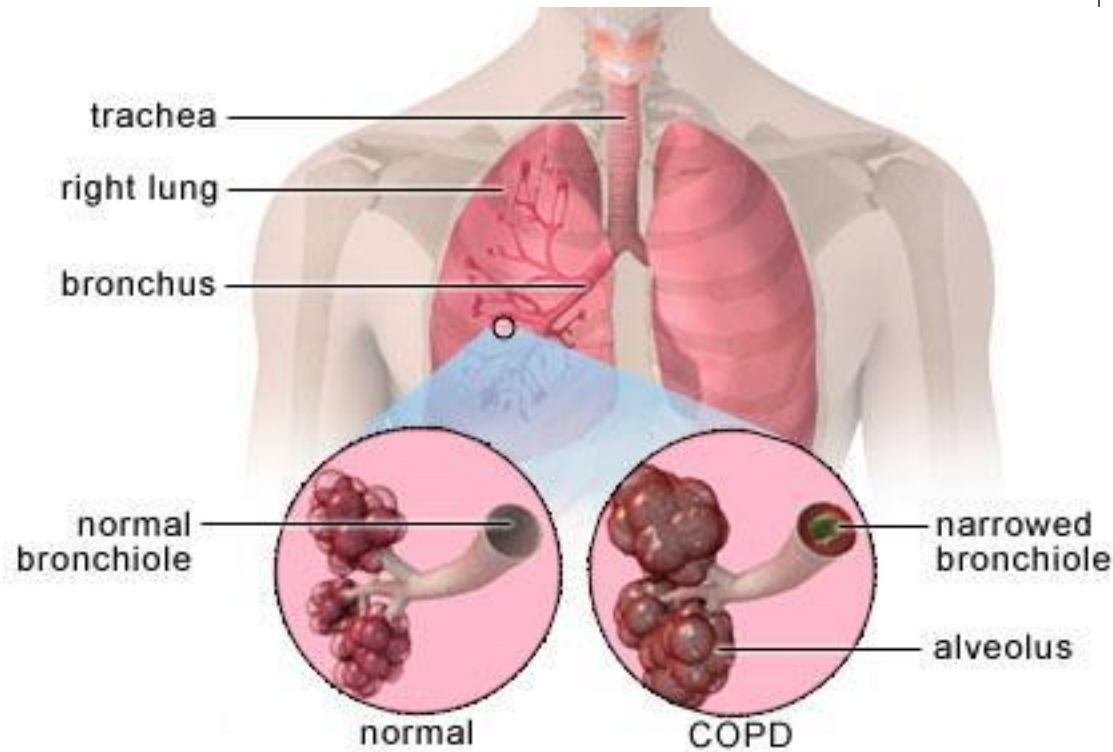
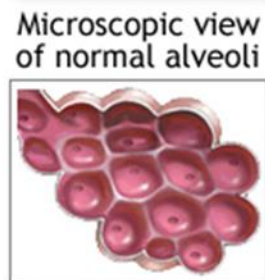
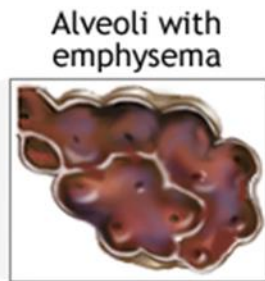
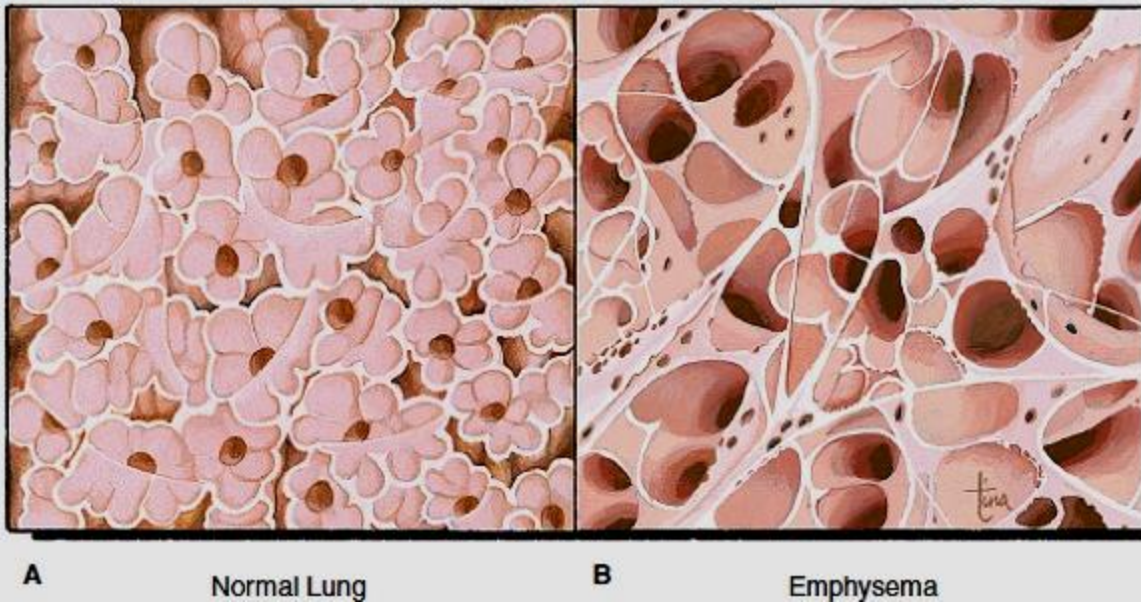
**Permanent enlargement of all or part of the
respiratory unit**

normal

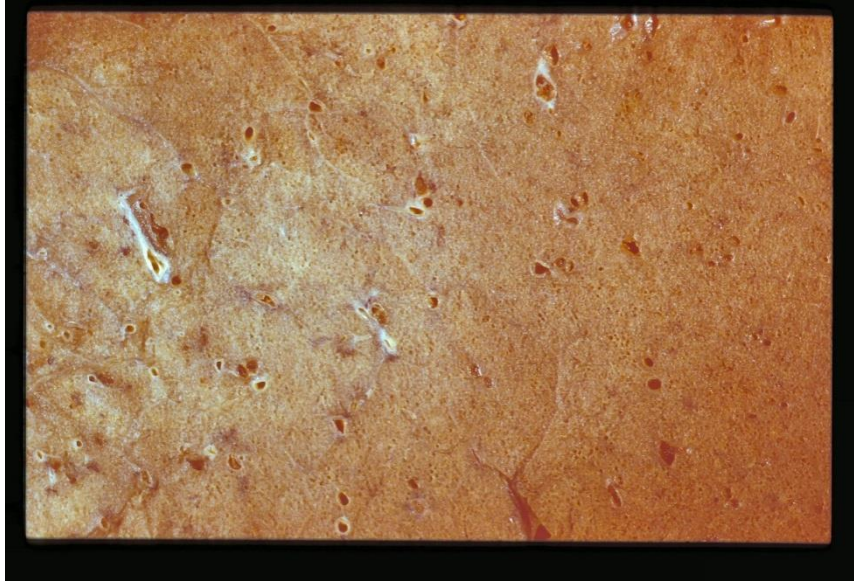


Emphysema

- Is abnormal permanent enlargement of the airspaces distal to the terminal bronchioles accompanied by destruction of their walls, without obvious fibrosis.
- Element of chronic bronchitis coexists
- Types of emphysema:
 1. Centriacinar
 2. Panacinar
 3. Distal acinar /paraseptal
 4. Irregular
- “Dilatation” is due to destruction and loss of alveolar walls (tissue destruction)
- Appears as “holes” in the lung tissue
- Emphysema Impairs Respiratory Function:
 - Diminished alveolar surface area for gas exchange (decreased Tco)
 - Loss of elastic recoil and support of small airways leading to tendency to collapse with obstruction

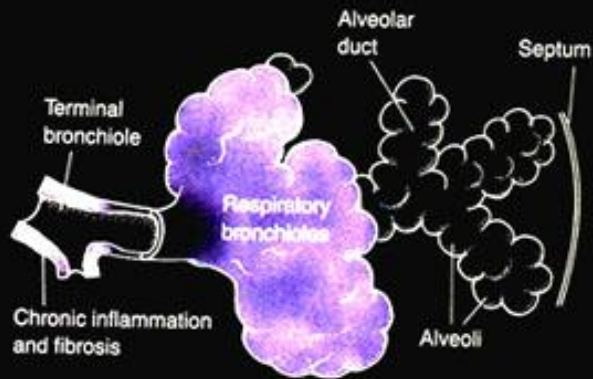
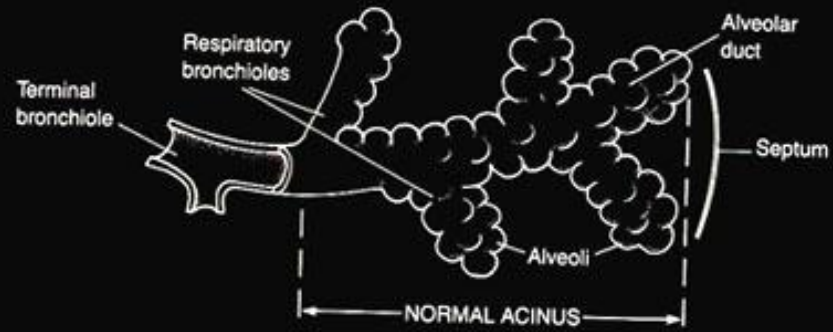


Normal

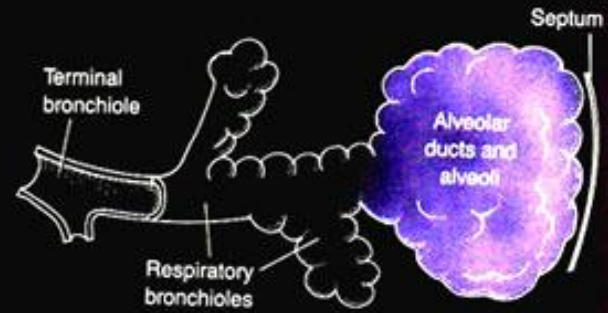


Emphysema

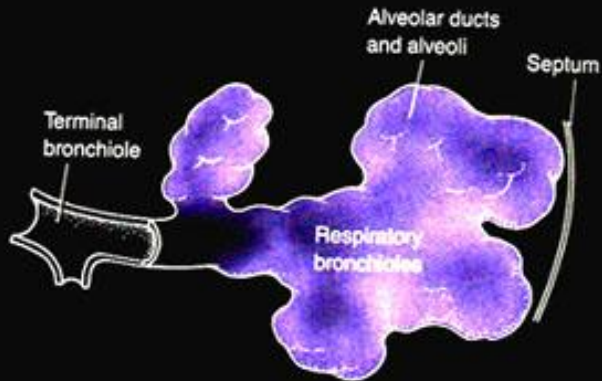




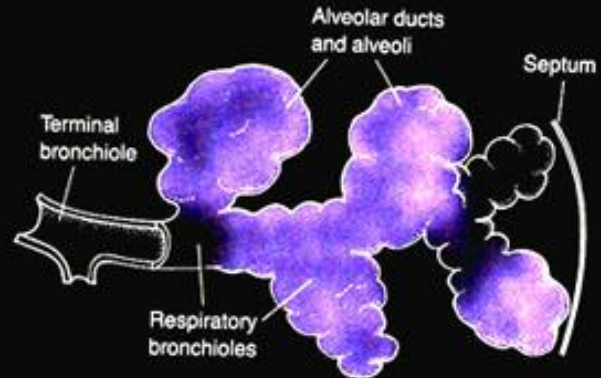
CENTRILOBULAR EMPHYSEMA



PARASEPTAL EMPHYSEMA



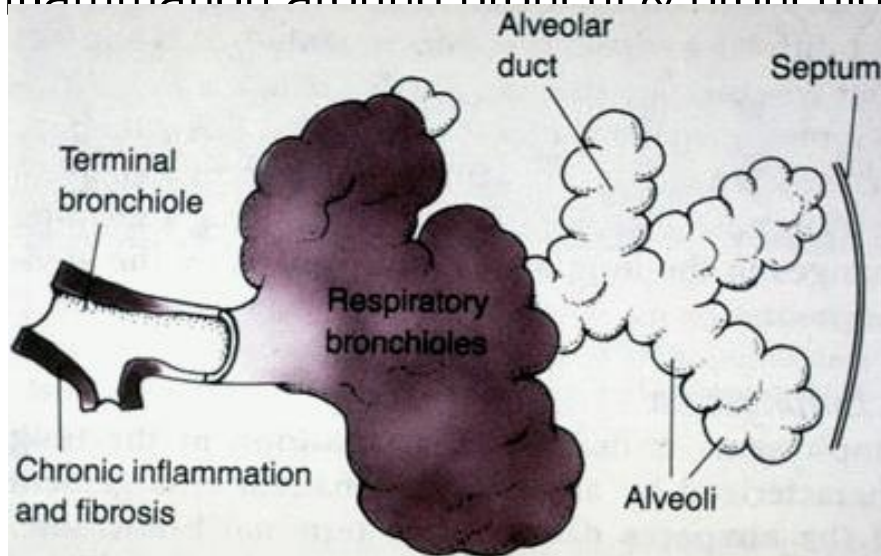
PANACINAR EMPHYSEMA



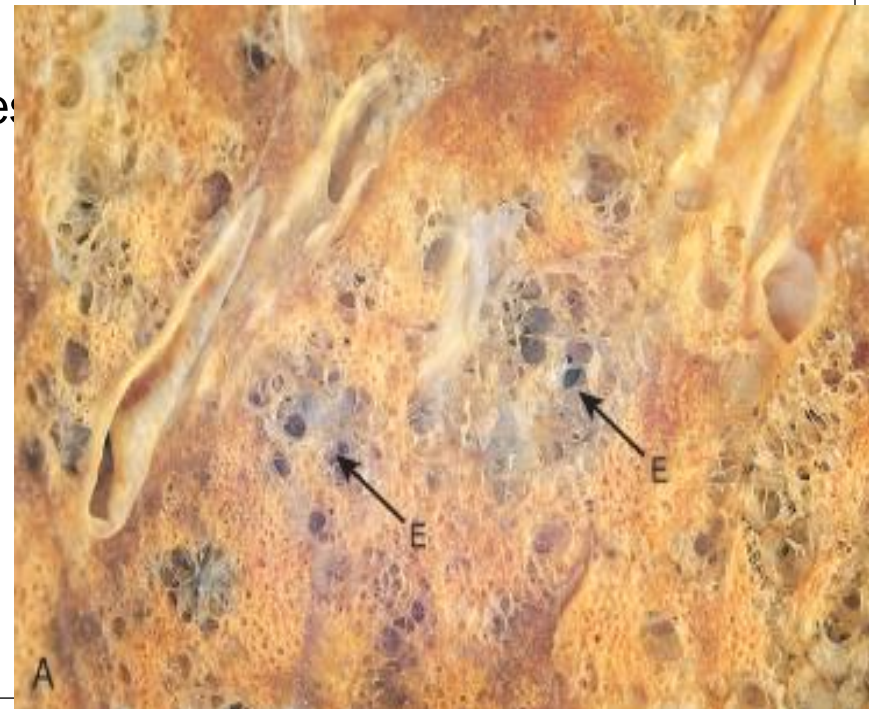
IRREGULAR EMPHYSEMA

Centriacinar (centrilobular) emphysema

- Occur in heavy smoker in association with chronic bronchitis
- The central or proximal parts of the acini are affected, while distal alveoli are spared
- More common and severe in upper lobes (apical segments)
- The walls of the emphysematous space contain black pigment.
- Inflammation around bronchi & bronchioles

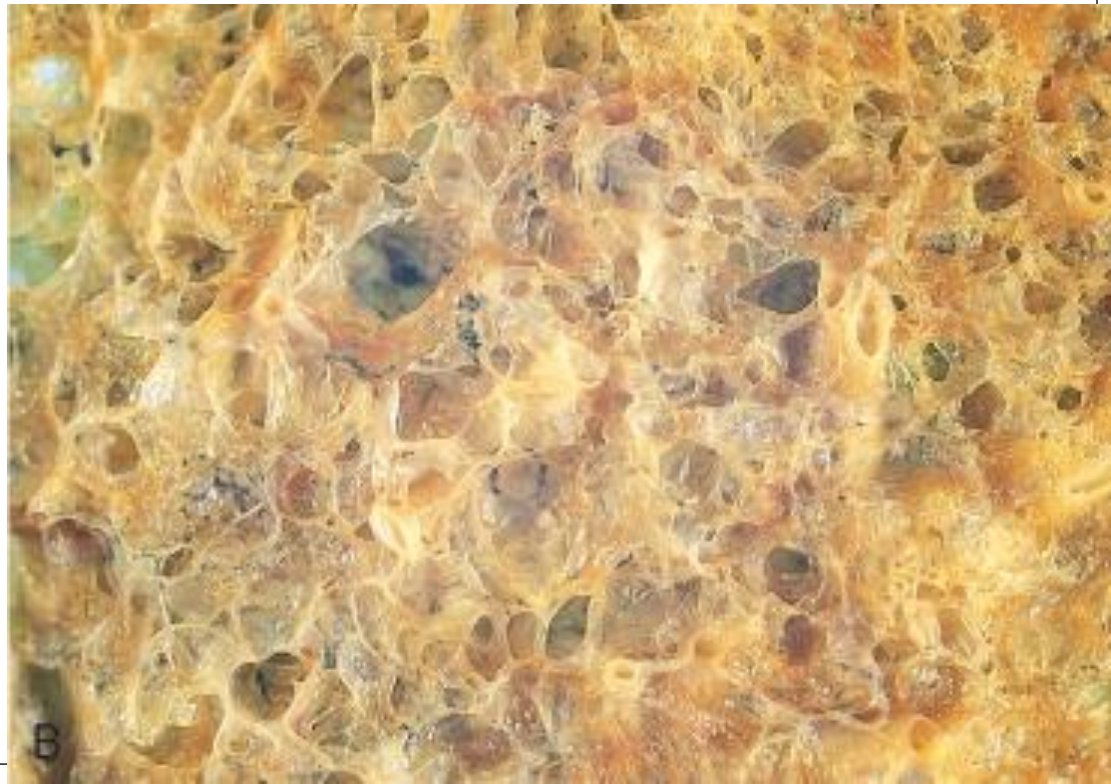
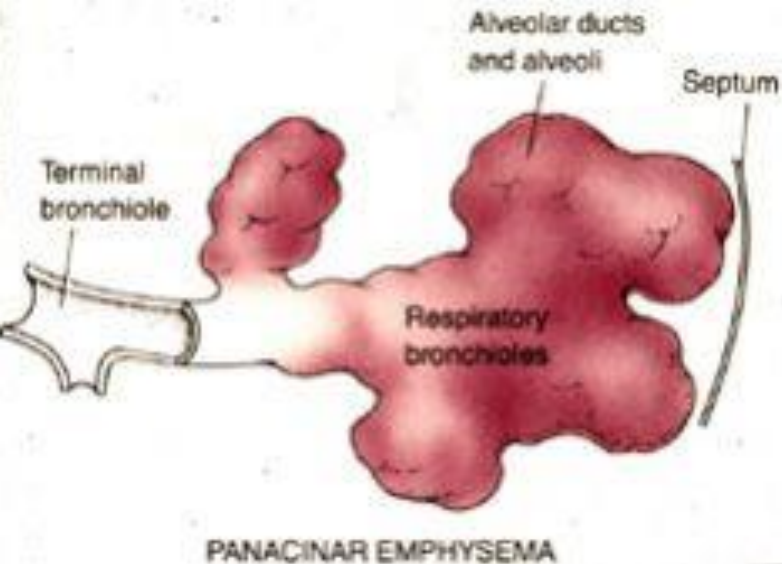


CENTRILOBULAR EMPHYSEMA



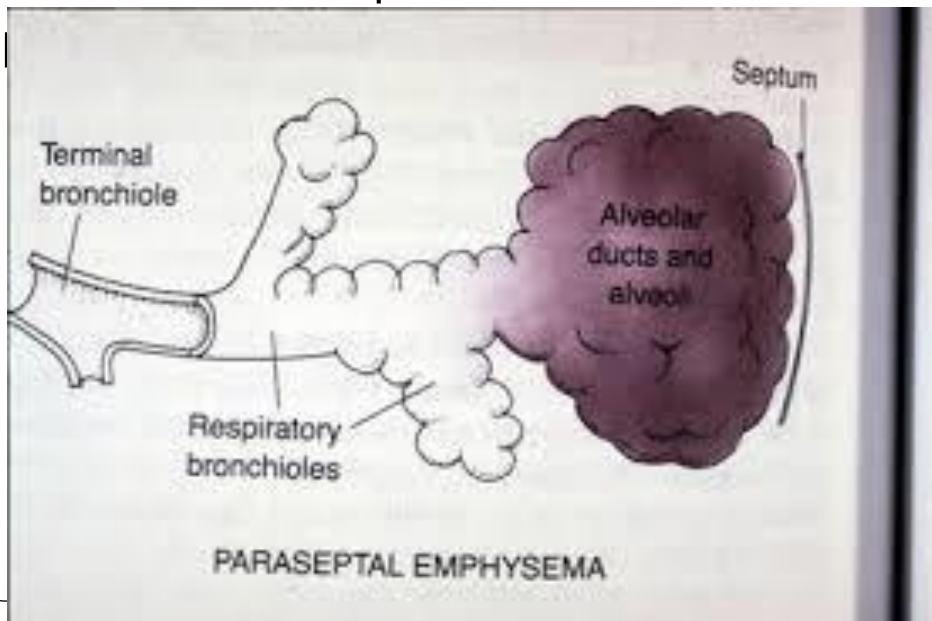
Panacinar (panlobular) emphysema

- Cause :Occurs in α_1 -anti-trypsin deficiency.
- **Uniform injury:** Acini are uniformly enlarged from the level of the respiratory bronchiole to the terminal blind alveoli.
- More commonly in the lower lung zones



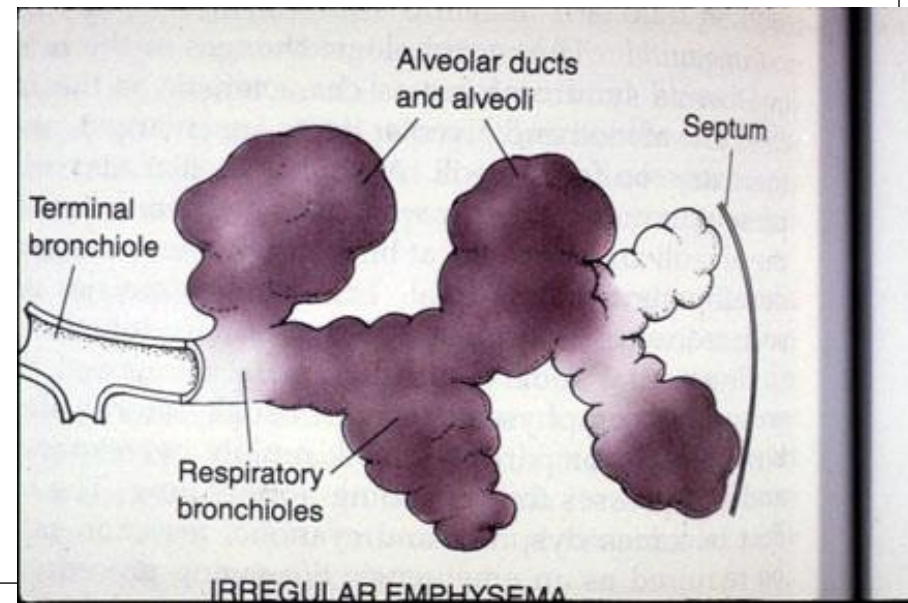
Distal acinar (paraseptal) emphysema

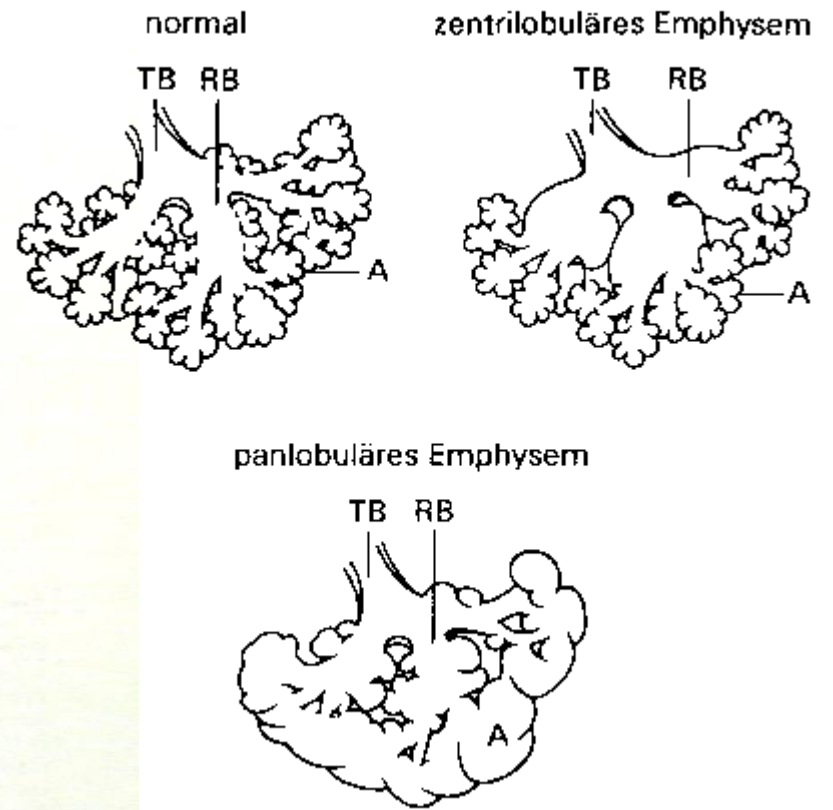
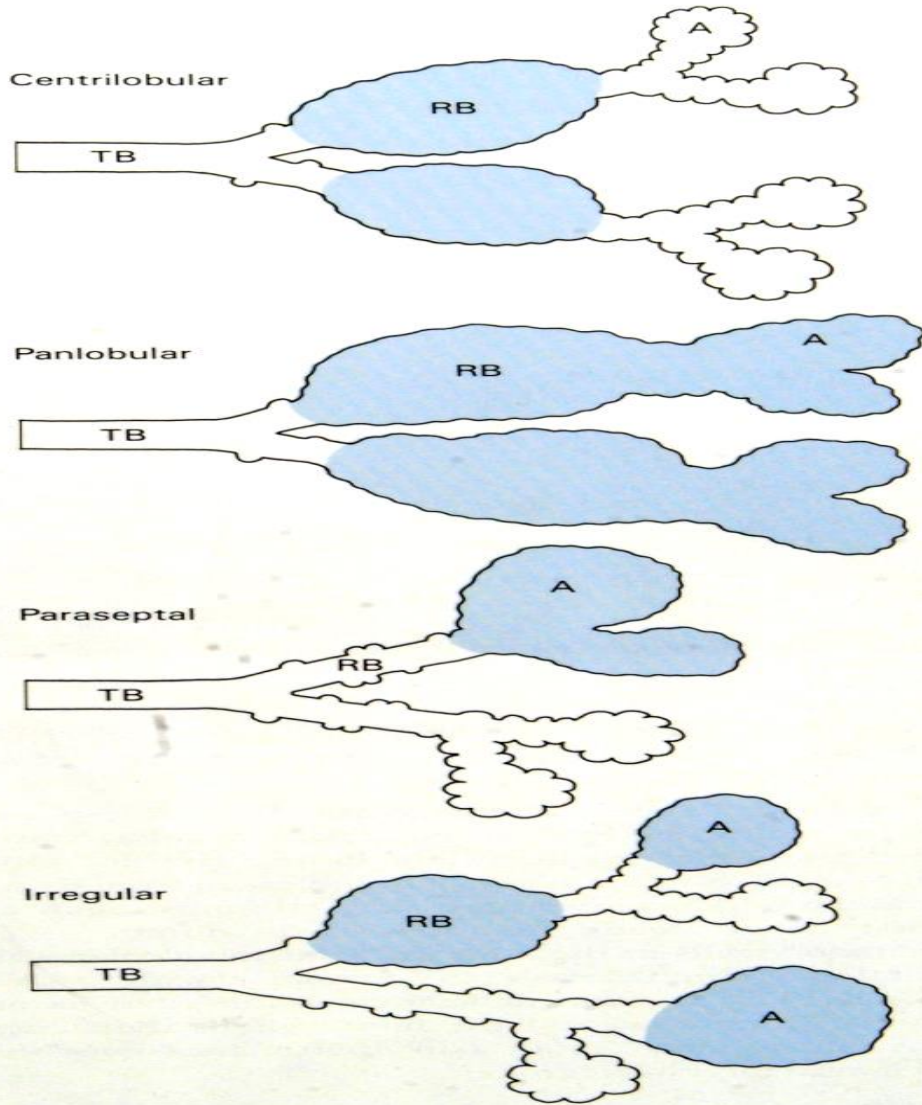
- The proximal portion of the acinus is normal but the distal part is dominantly involved.
- Occurs adjacent to areas of fibrosis, scarring or atelectasis.
- More severe in the upper half of the lungs.
- Sometimes forming multiple cyst-like structures with spontaneous



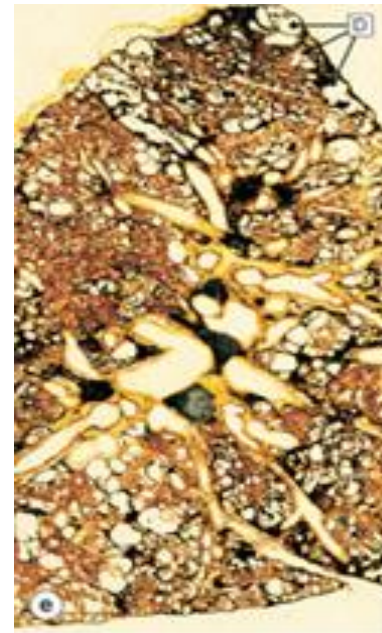
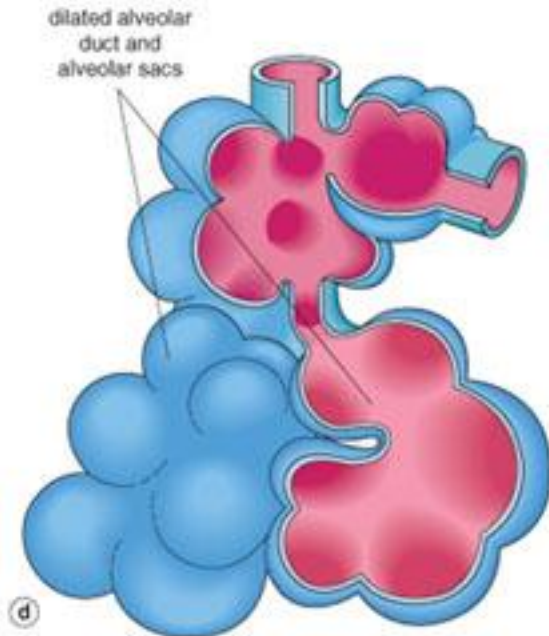
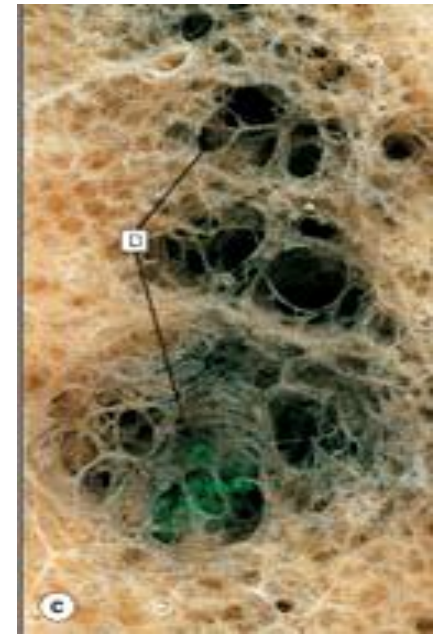
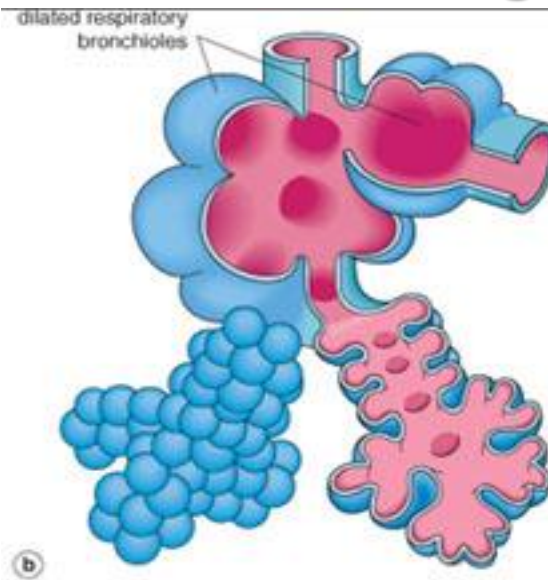
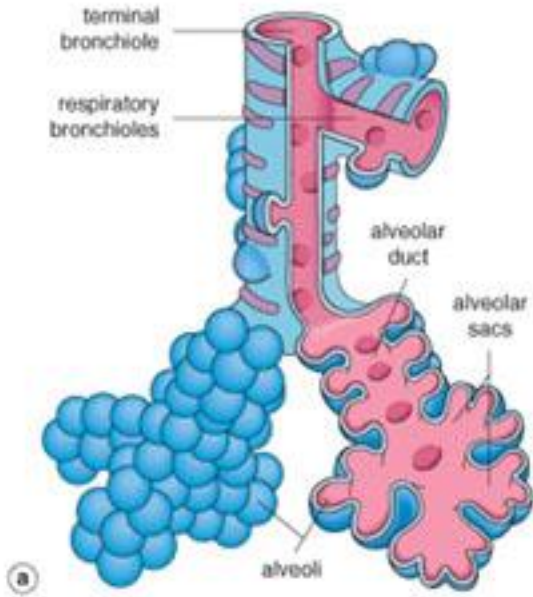
Irregular Emphysema

- The acinus is irregularly involved, associated with scarring.
- Most common form found in autopsy.
- Asymptomatic.
- usually a complication of various inflammatory processes including chronic pulmonary tuberculosis



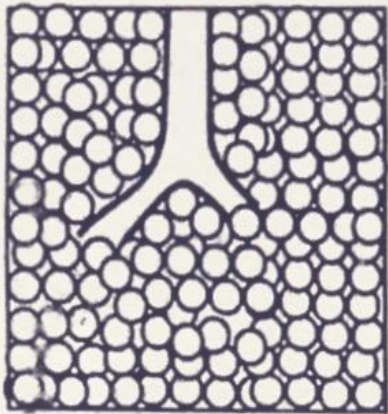


Diseases of Lung

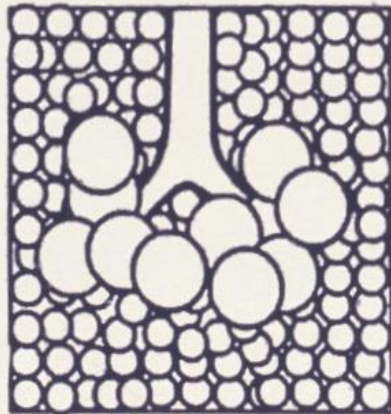


Generalized emphysema. (a) Normal distal lung acinus. (b) Centriacinar emphysema. (c) Centriacinar emphysema. (d) Panacinar emphysema. (e) Panacinar emphysema (Gough-Wentworth section).

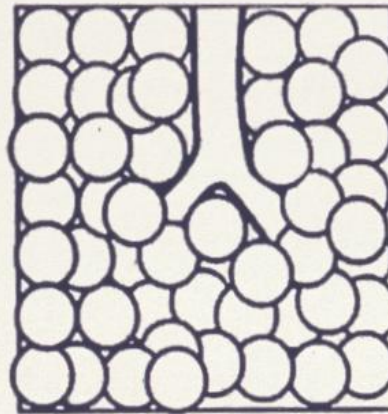
Loss of surface area (emphysema)



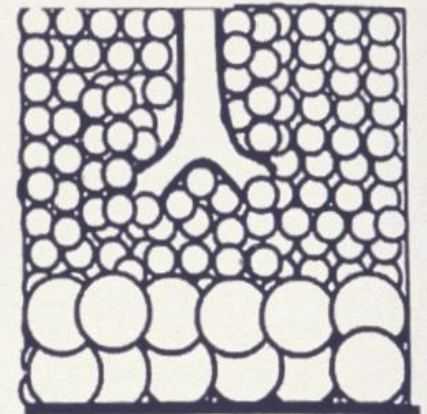
Normal lung



Centriacinar emphysema



Panacinar emphysema



Paraseptal emphysema

Why is emphysema considered to be an obstructive airway disease?

Is there any mechanical obstruction ?

- Because emphysema affects the peripheral airways, it is not, anatomically speaking, an obstructive disease, and there is no mechanical obstruction. However, it is functionally an obstructive disease, because destruction of the wall of the air spaces prevents the elastic recoil that is necessary to push air out of the lungs. Thus, in effect, there is limitation of airflow, just as there would be if there were mechanical obstruction.

Pathogenesis of Emphysema

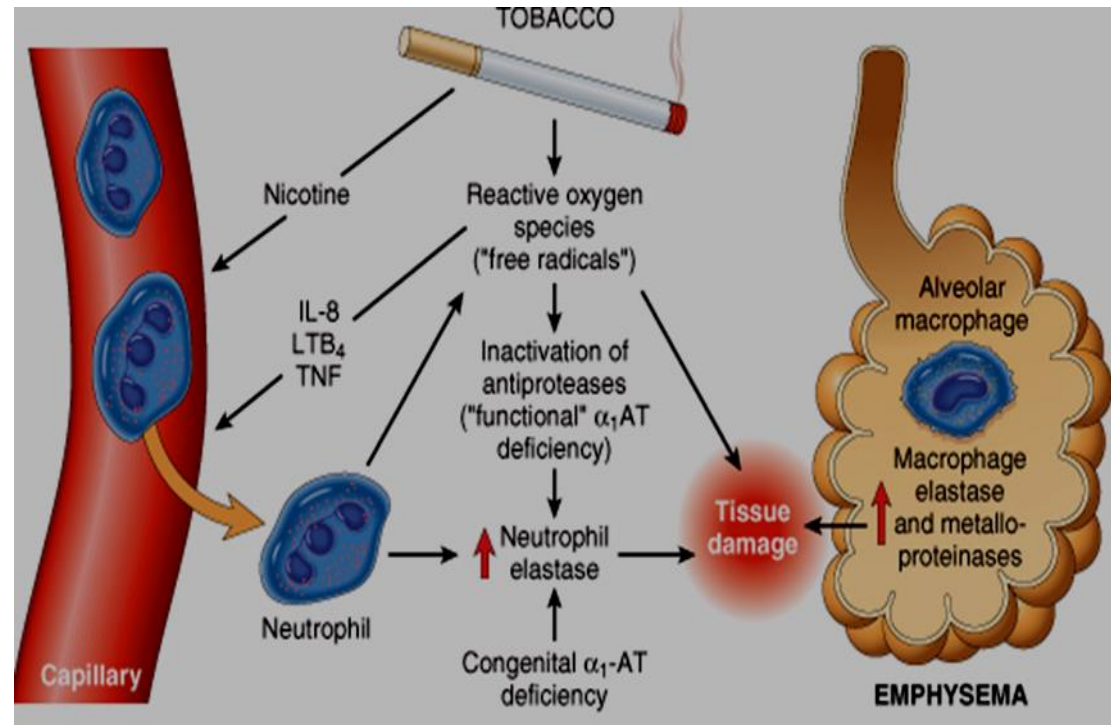
- Is not completely understood
- Elastic tissue of the alveolar wall is broken down by action of proteolytic enzymes like **protease (e.g.elastase)**.
- **Protease** is produced by neutrophils and macrophages.
- ***Alpha 1 antitrypsin* is an anti-protease** (anti-elastase) and it counter acts the protease. It is a major inhibitor of proteases secreted by neutrophils during inflammation. α_1 -antitrypsin is normally present in the serum, in tissue fluids and in macrophages.
- Normally there is a balance between protease and anti protease activity.

Pathogenesis of Emphysema

- Any condition that increases the neutrophils or macrophages in the lung will lead to release of protease enzyme which causes damage to the elastic tissue of the alveolar wall.
- Therefore one of the key mechanisms in emphysema is alveolar wall destruction which occurs due to excess proteases (elastase) activity coupled with low anti-protease level and inflammation (protease-antiprotease hypothesis)
- Elements of chronic bronchitis may co-exist
- Hereditary alpha 1-antitrypsin deficiency leads to panacinar emphysema.

Pathogenesis of Emphysema

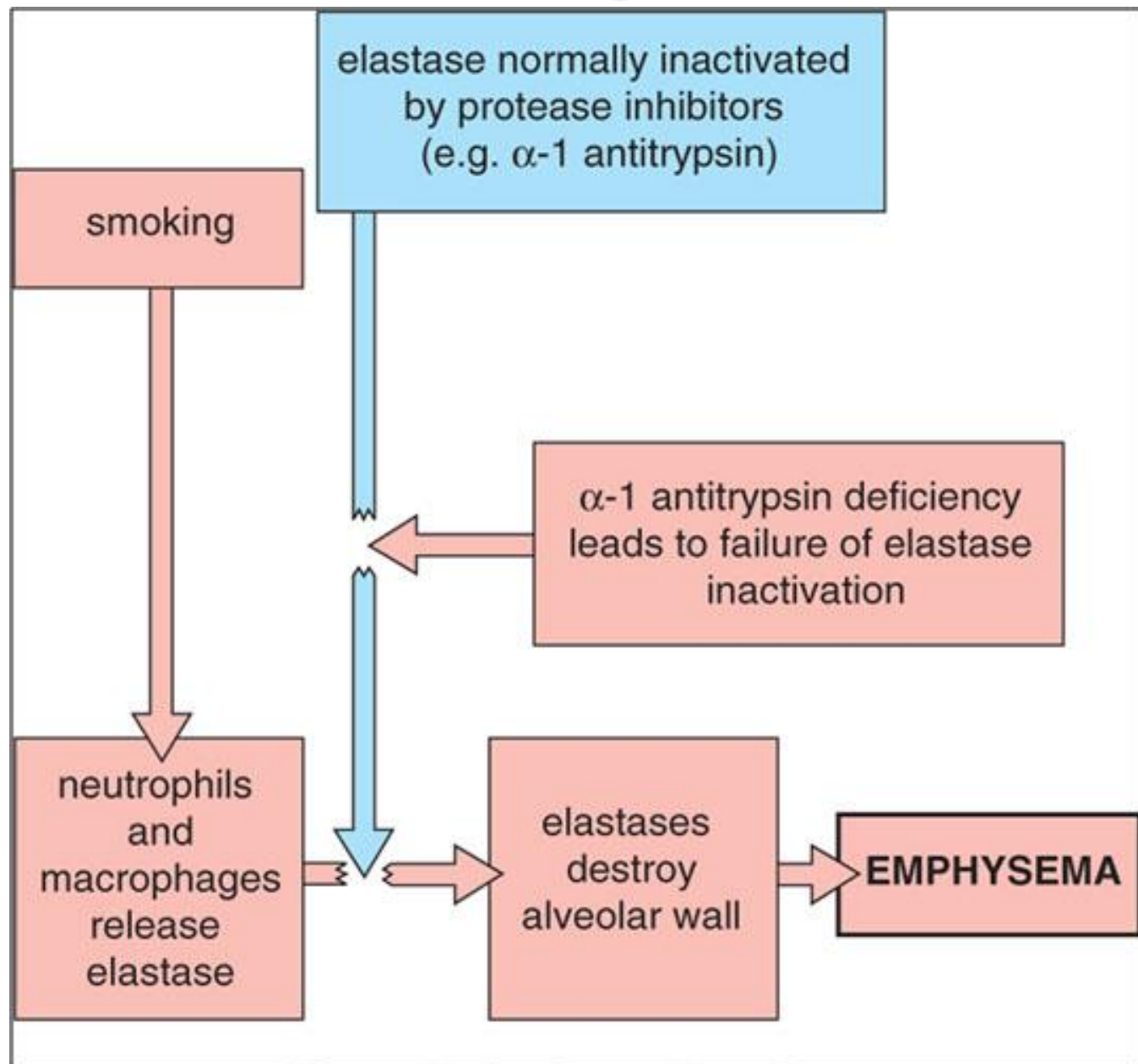
- Smokers have increases number of neutrophils and macrophages in their alveoli
- Smoking stimulates release of elastase and enhances elastase activity in macrophages.
- Smoking Inhibits alpha 1 antitrypsin.
- Tobacco smoke contains reactive oxygen species with inactivation of anti-proteases.
- The protease-antiprotease hypothesis explains the effect of cigarette smoking in the production of centriacinar emphysema



Diseases of Lung



I don't want emphysema so I'm never going to start smoking!

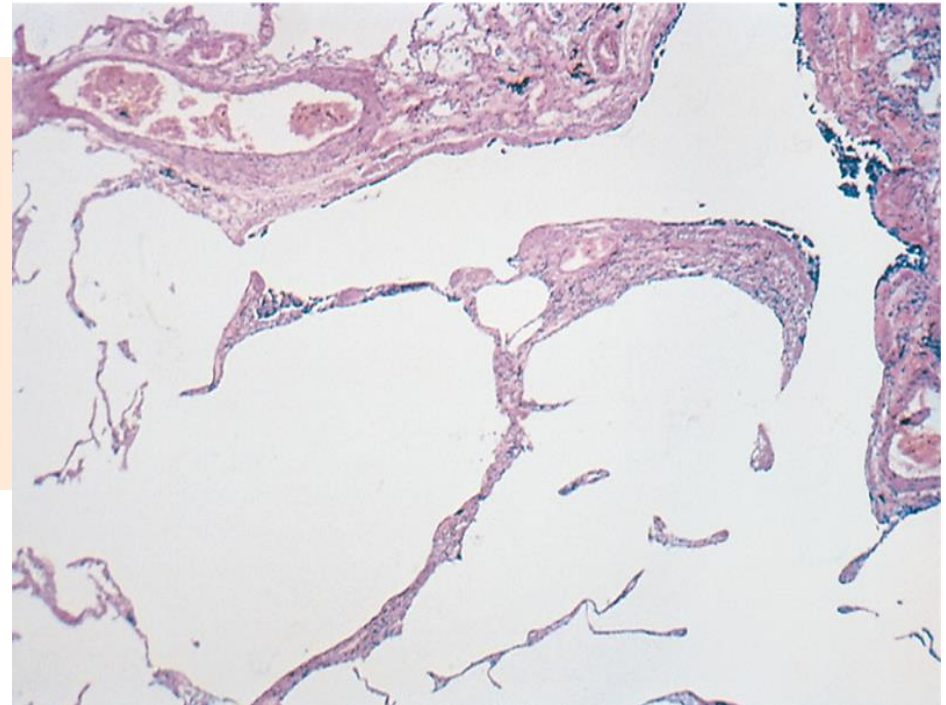


Pathogenesis of emphysema.

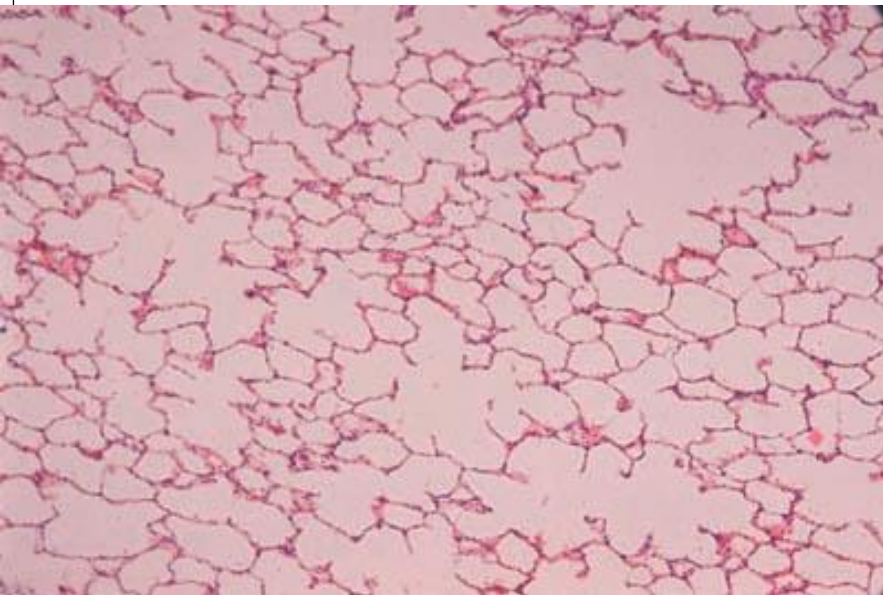
Emphysema: Morphology

- The lungs are pale, voluminous.
- Histologically, thinning and destruction of alveolar walls creating large airspaces.

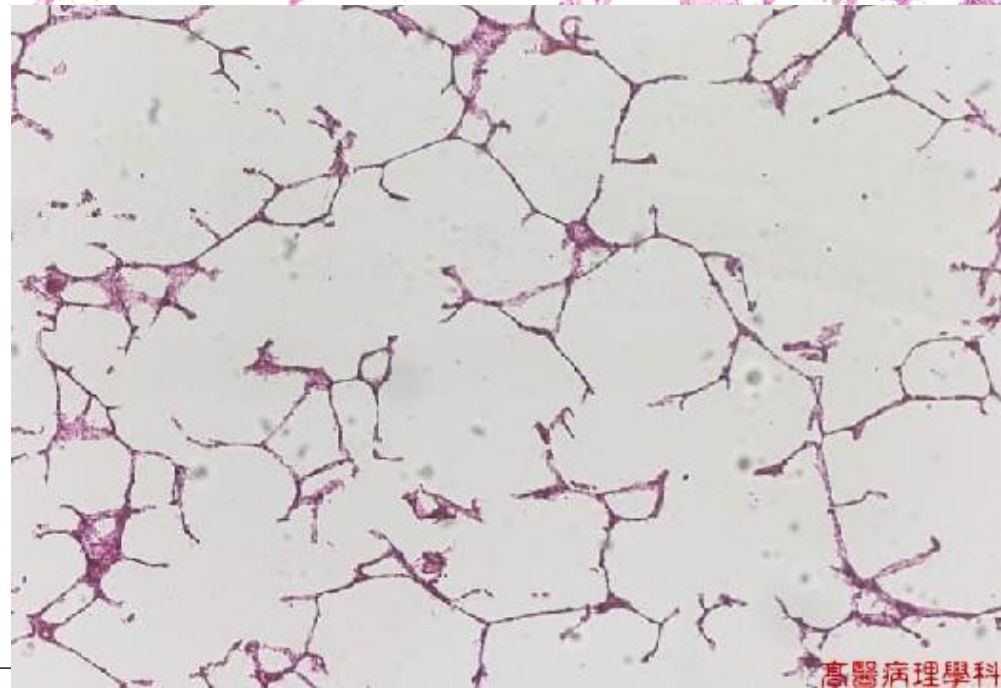
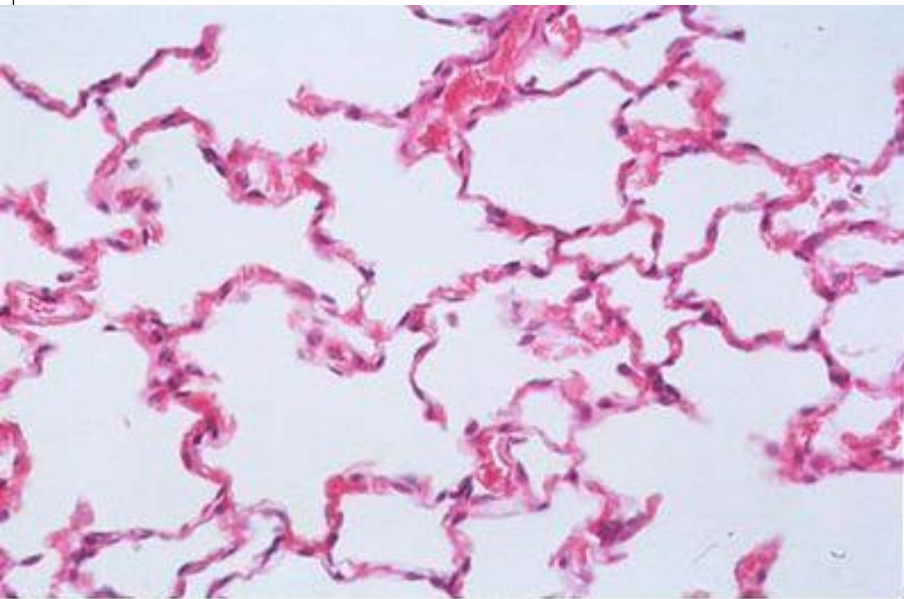
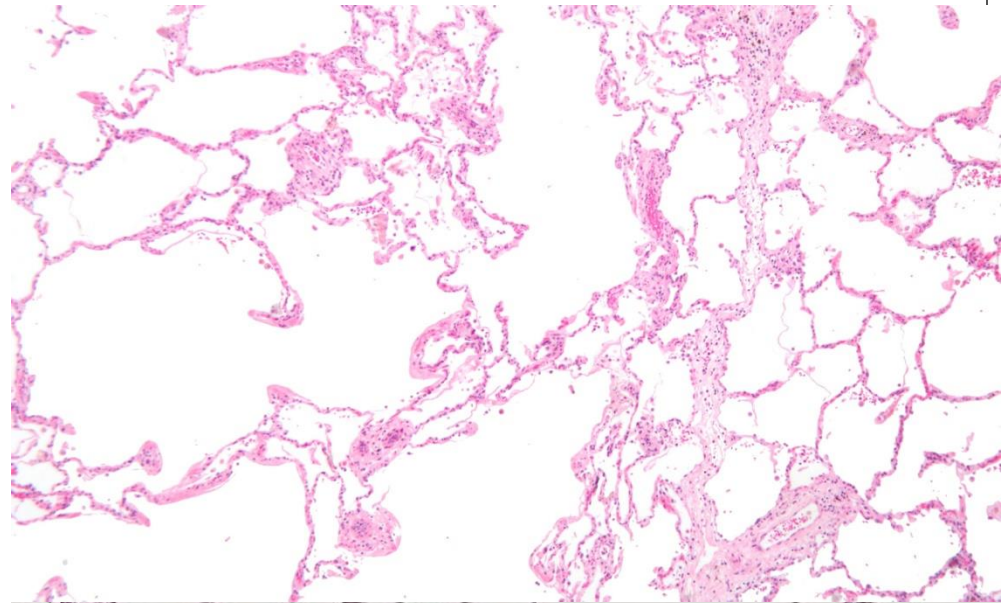
- Loss of elastic tissue.
- Reduced radial traction on the small airways.
- Alveolar capillaries is diminished.
- Accompanying bronchitis and bronchiolitis.



- Normal



- Emphysema



Lung 400x

Emphysema:

Clinical course

- Cough and wheezing
- Weight loss
- Barrell chest
(↑ anteroposterior diameter of chest)
- Pulmonary function tests reveal reduced FEV1
- Advanced: hypoxia, cyanosis, respiratory acidosis
- Patients are known as pink puffers

Complications

- Coexistent chronic bronchitis
- Interstitial emphysema in which air escapes into the interstitial tissues of the chest from a tear in the airways.
- may also be complicated by rupture of a surface bleb with resultant Pneumothorax
- Death from emphysema is related to::
 1. Pulmonary failure with respiratory acidosis, hypoxia and coma.
 2. Cor pulmonale : (Right-sided heart failure induced by pulmonary disease)

EMPHYSEMA

“PINK PUFFER”



- * ↑CO₂ Retention (Pink)
- * Minimal Cyanosis
- * Purse Lip Breathing
- * Dyspnea
- * Hyperresonance on Chest Percussion
- * Orthopneic
- * Barrel Chest
- * Exertional Dyspnea
- * Prolonged Expiratory Time
- * Speaks in Short Jerky Sentences
- * Anxious
- * Use of Accessory Muscles to Breathe
- * Thin Appearance

Emphysema:

Dilated air spaces beyond respiratory arteriols

Types

- Centriacinar: Smoking
- Panacinar: deficiency of $\alpha 1$ AT
- Paraseptal
- Irregular: scar

Clinical features

- Cough and wheezing. Respiratory acidosis
- Weight loss.
- Pulmonary function tests reveal reduced FEV1.

Complications

- Pneumothorax
- Death from emphysema is related to:
 - Pulmonary failure with respiratory acidosis, hypoxia and coma.
 - Right-sided heart failure (Cor pulmonale)

Emphysema and Chronic Bronchitis

	Predominant Bronchitis	Predominant Emphysema
Appearance	“Blue bloaters”	“Pink Puffers”
Age	40-45	50-75
Dyspnea	Mild, late	Severe, early
Cough	Early, copious sputum	Late, scanty sputum
Infection	Common	Occasional
Cor pulmonale	Common	Rare, terminal
Airway resistance	Increased	Normal or slightly increased
Elastic recoil	Normal	Low
Chest radiography	Prominent vessels, large heart	Hyperinflation, small heart
PaCO ₂	Increased	Normal to decreased
Cyanosis	Present	Absent

CHRONIC BRONCHITIS

CLINICAL DIAGNOSIS: DAILY PRODUCTIVE COUGH FOR THREE MONTHS OR MORE, IN AT LEAST TWO CONSECUTIVE YEARS

OVERWEIGHT AND CYANOTIC



ELEVATED HEMOGLOBIN



PERIPHERAL EDEMA

RHONCHI AND WHEEZING



EMPHYSEMA

PATHOLOGIC DIAGNOSIS: PERMANENT ENLARGEMENT AND DESTRUCTION OF AIRSPACES DISTAL TO THE TERMINAL BRONCHIOLE

OLDER AND THIN

SEVERE DYSPNEA

QUIET CHEST



X-RAY:
HYPERINFLATION
WITH FLATTENED
DIAPHRAGMS



Chronic bronchitis vs. Emphysema

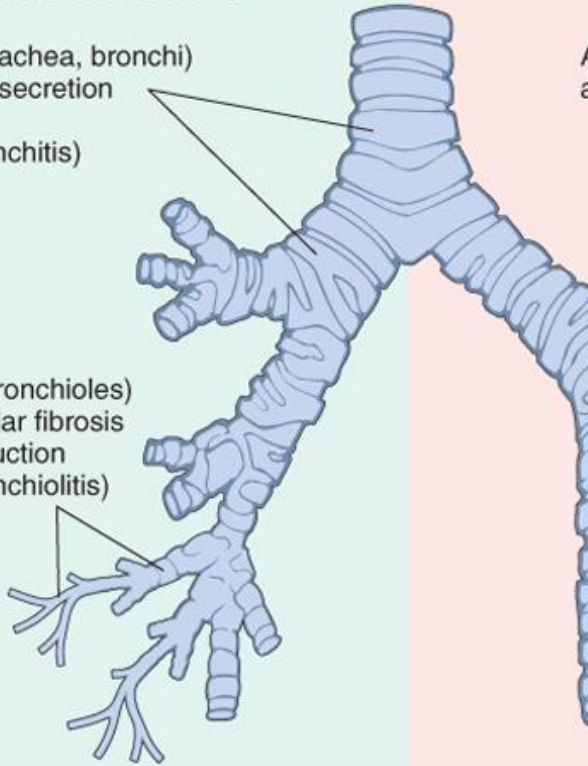
PURE CHRONIC BRONCHITIS

Large airways (trachea, bronchi)

- Mucus hypersecretion
- Inflammation
- (Chronic bronchitis)

Small airways (bronchioles)

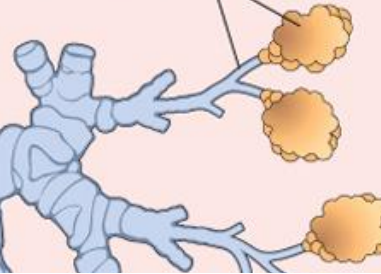
- Peribronchiolar fibrosis
- Airway obstruction
- (Chronic bronchiolitis)



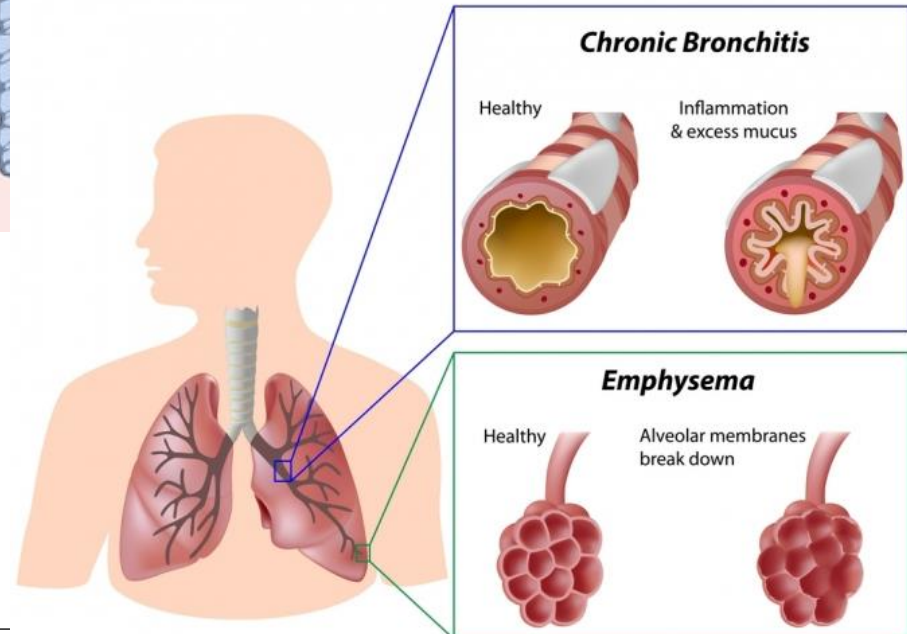
PURE EMPHYSEMA

Acinus (respiratory bronchiole, alveolar ducts, and alveoli)

- Loss of elastic recoil
- (Emphysema)

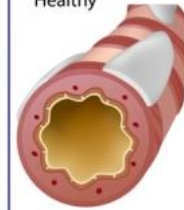


Chronic Obstructive Pulmonary Disease (COPD)



Chronic Bronchitis

Healthy



Inflammation & excess mucus



Emphysema

Healthy



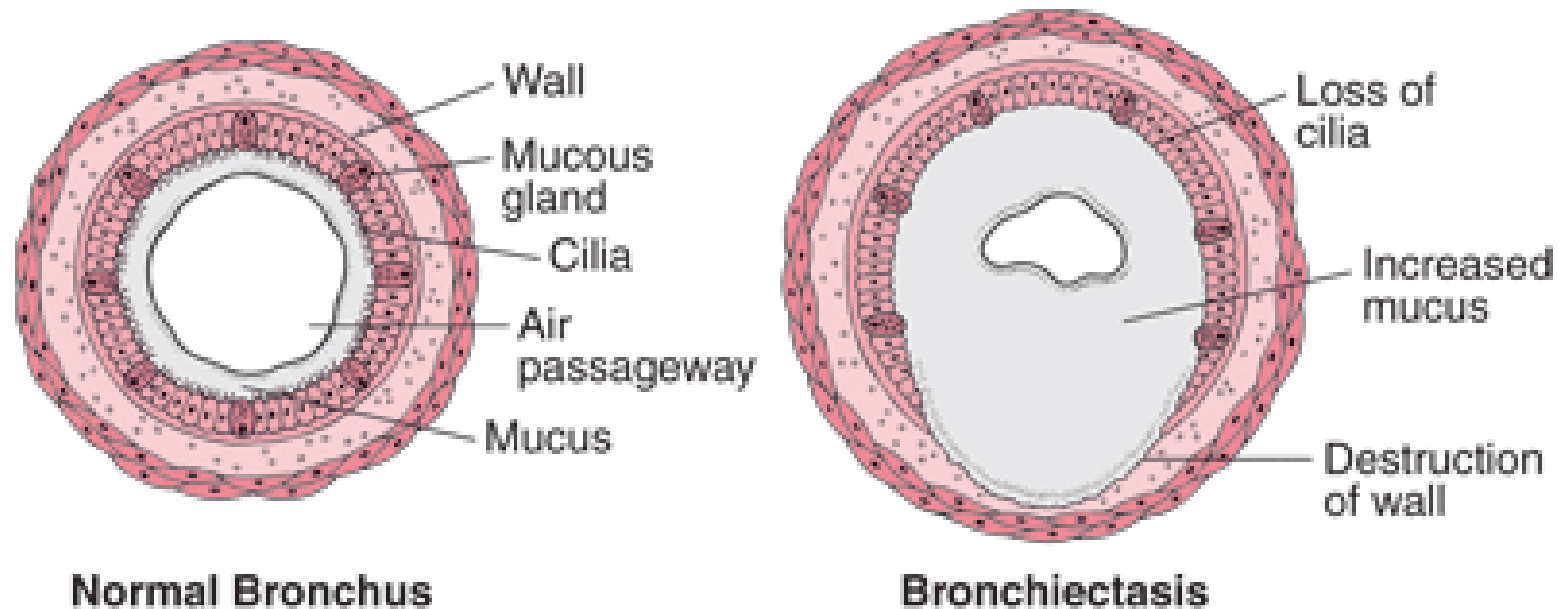
Alveolar membranes break down



Bronchiectasis

Bronchiectasis

- is chronic necrotizing infection and inflammation of the bronchi and bronchioles leading to abnormal permanent dilation of these airways. It represents the end stage of a variety of pathologic processes that cause destruction of the bronchial wall.
- most often involves the lower lobes of both lungs.
- is characterized by fever and cough with production of copious purulent foul smelling sputum,
- and recurrent pulmonary infection that may lead to lung abscess.



Bronchiectasis is a result of chronic inflammation compounded by an inability to clear mucoid secretions. Conditions commonly associated with Bronchiectasis are as follows:

1. Bronchial obstruction

Localized:

- tumor, foreign bodies or mucous impaction

Generalized:

- bronchial asthma
- chronic bronchitis

2. Congenital or hereditary conditions:

- Congenital bronchiectasis
- Cystic fibrosis.
- Intralobar sequestration of the lung.
- Immunodeficiency status.
- Immotile cilia and kartagner syndrome

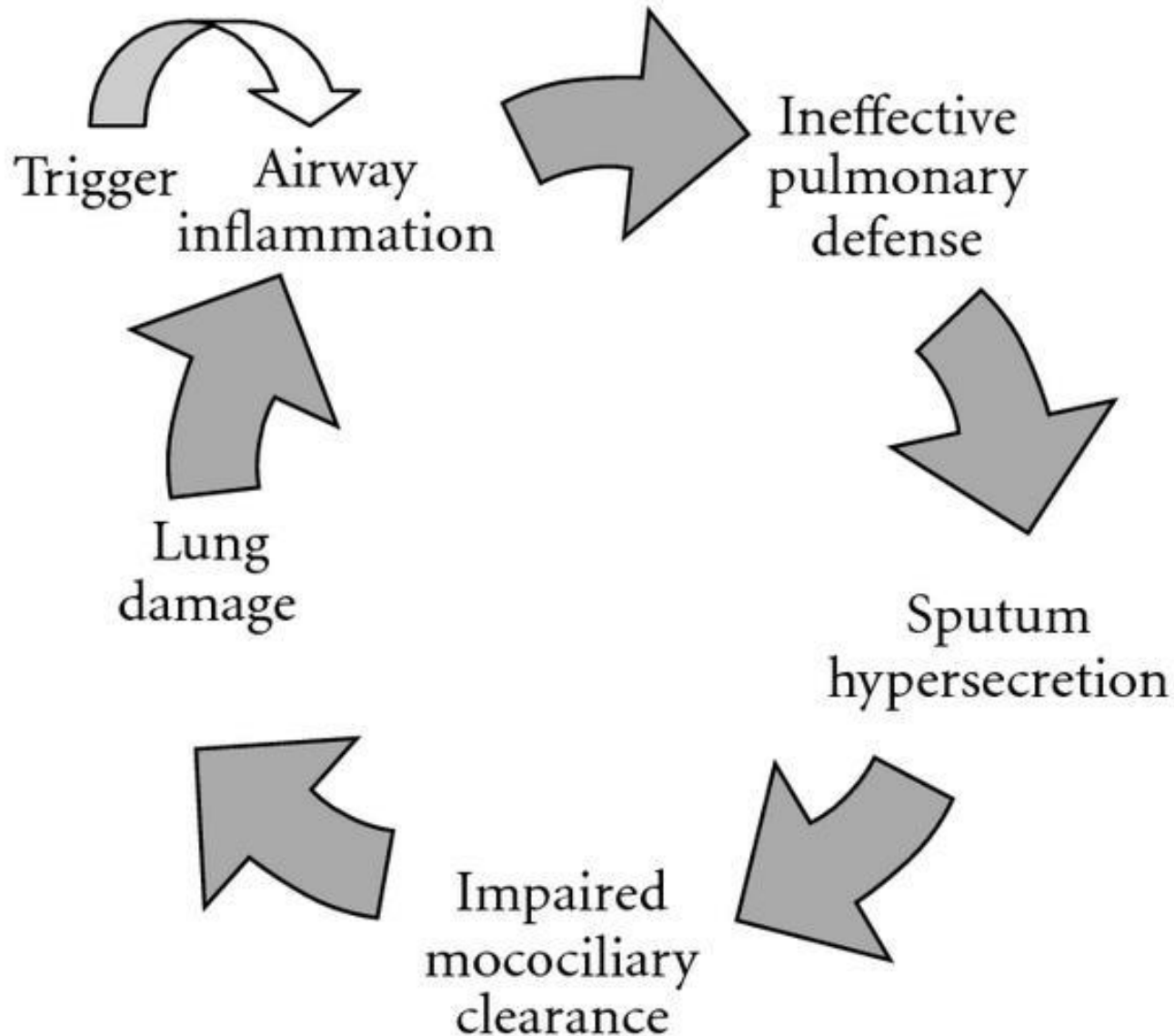
3. Chronic or severe infection / necrotizing pneumonia

Caused by TB, staphylococci or mixed infection.

Pathogenesis

- Any of the previously mentioned conditions can cause damage to the airways resulting in **impaired mucociliary clearance, mucus stasis and accumulation** which in turn further makes the airways susceptible to microbial colonization. The persistence of the pathology with superadded infection leads to a "vicious circle" of inflammation and tissue damage.
- Inflammation results in progressive destruction of the normal lung architecture, in particular the elastic fibres of bronchi.
- Neutrophils are thought to play a central role in the pathogenesis of tissue damage that occurs in bronchiectasis.

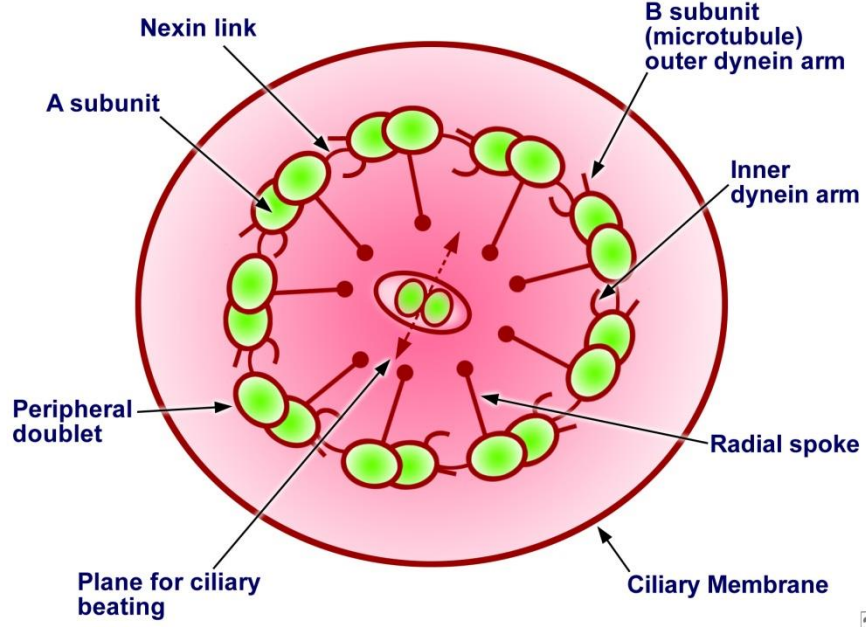
Pathogenesis of bronchiectasis



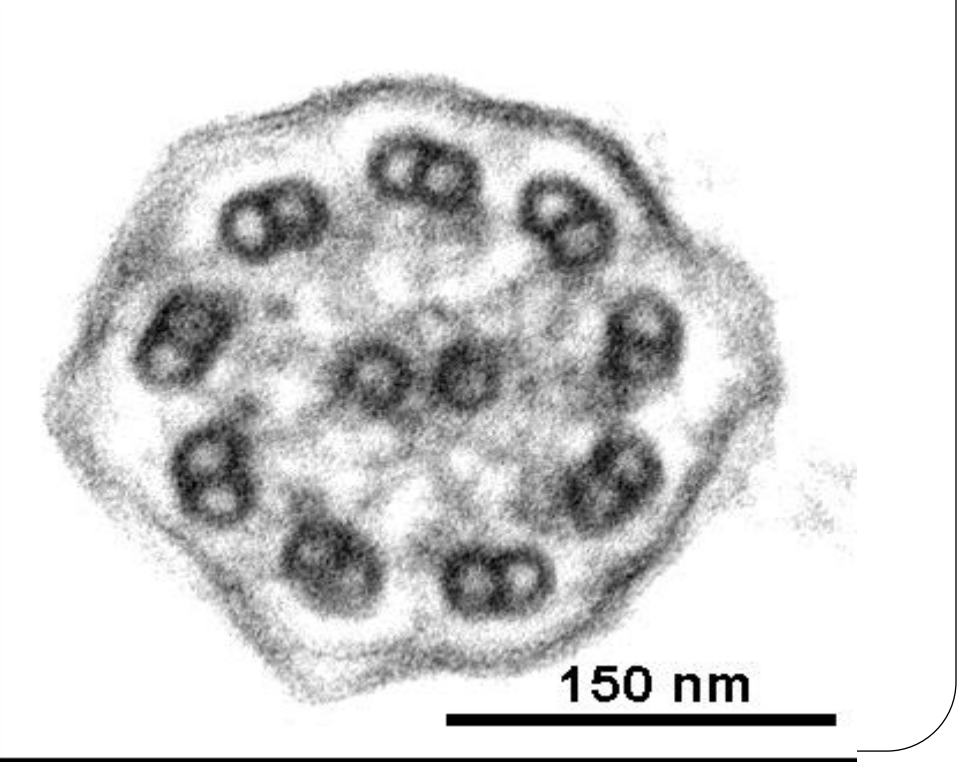
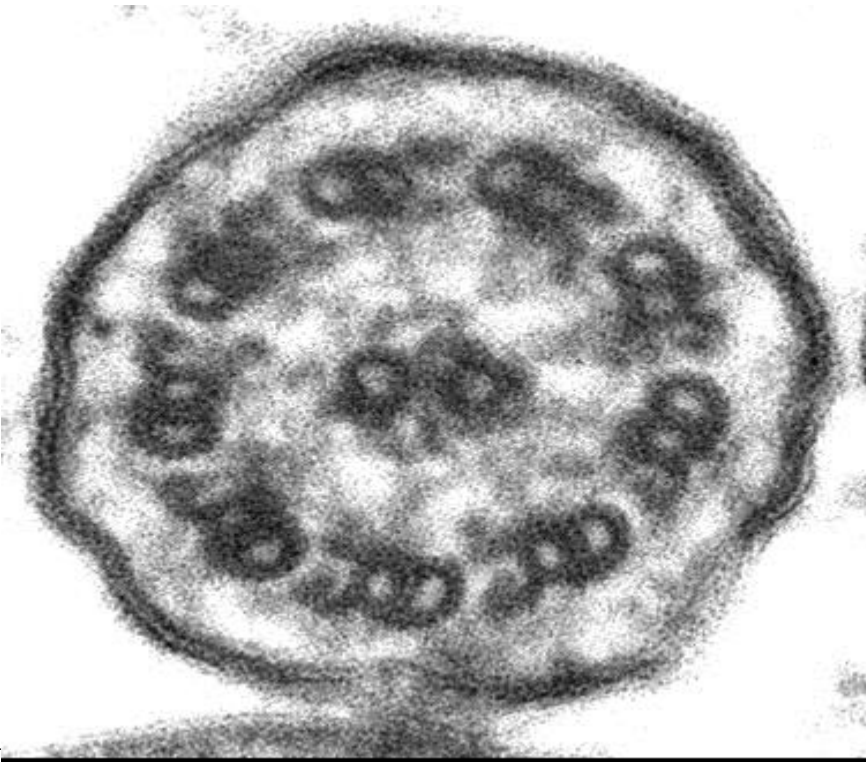
Kartagener Syndrome/ immotile cilia syndrome

- It is a genetic condition resulting in the failure to clear sputum (Primary ciliary dyskinesia) caused by a defect in the motility of respiratory, auditory, and sperm cilia.
- Inherited as autosomal recessive trait.
- Patient develop bronchiactasis, sinusitis and situs invertus sometimes with hearing loss and male sterility.
- Lack of ciliary activity interferers with bronchial clearance of mucus.

absence of outer and inner dynein arms in a patient with primary ciliary dyskinesia.



Dynein, a type of ATPase, provides energy for microtubule sliding and the longitudinal displacement of adjacent microtubular doublets, resulting in ciliary bending.

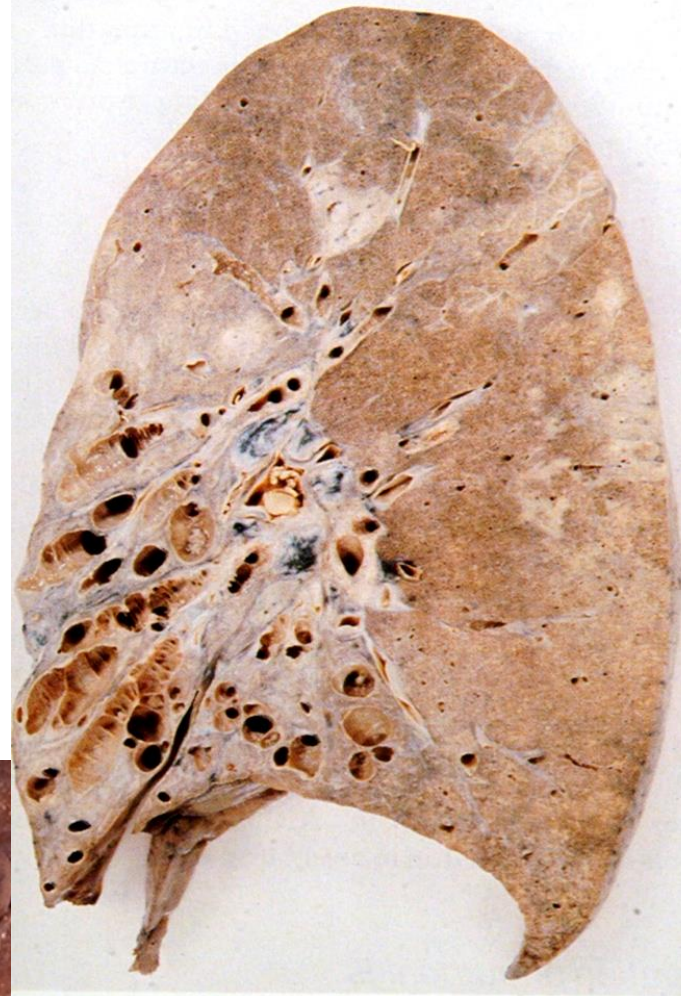


Cystic fibrosis

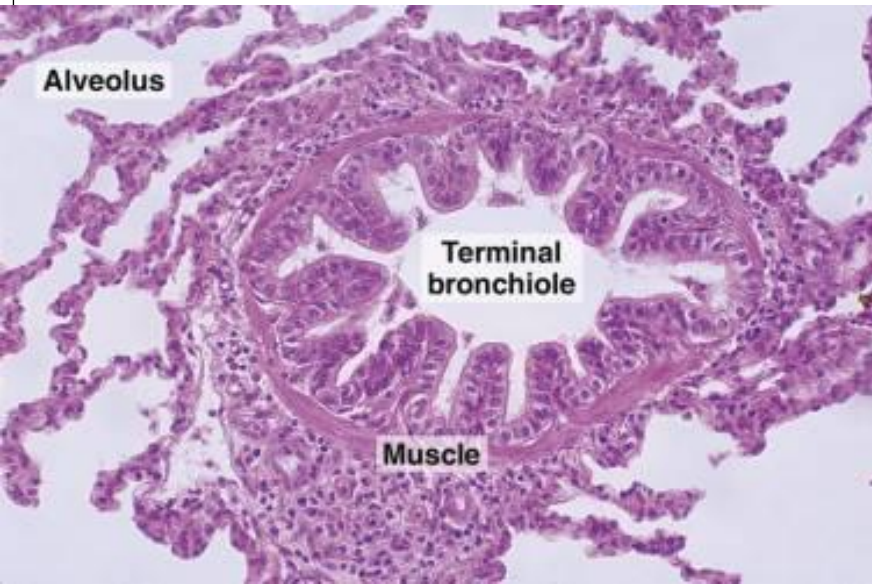
- Cystic fibrosis is an inherited disease that causes thick, sticky viscous mucus to build up in the lungs and digestive tract. It is one of the most common chronic lung diseases in children and young adults, and may result in early death.
- It may lead to bronchiectasis.

Morphology of Bronchiectasis

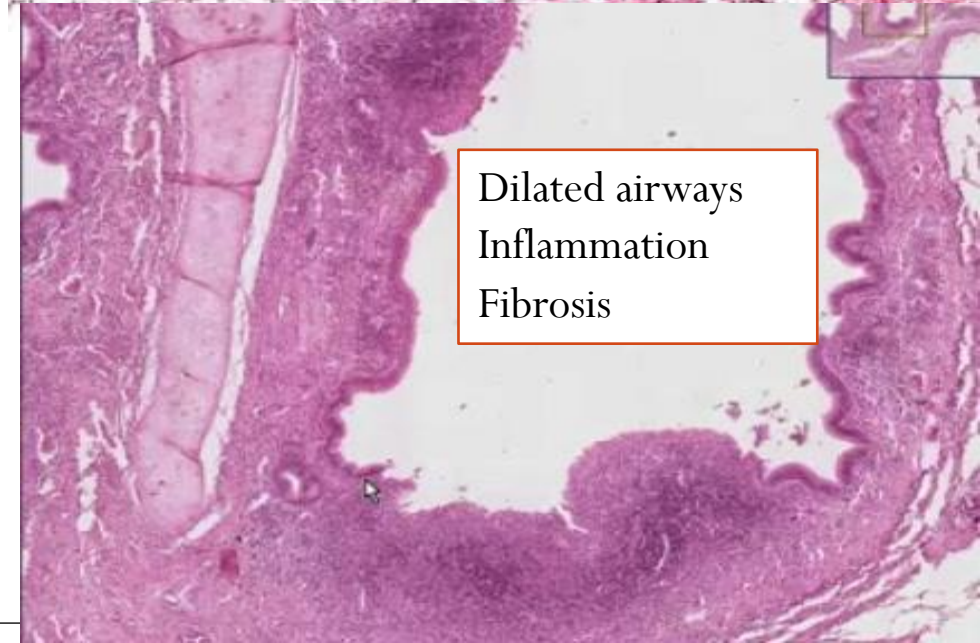
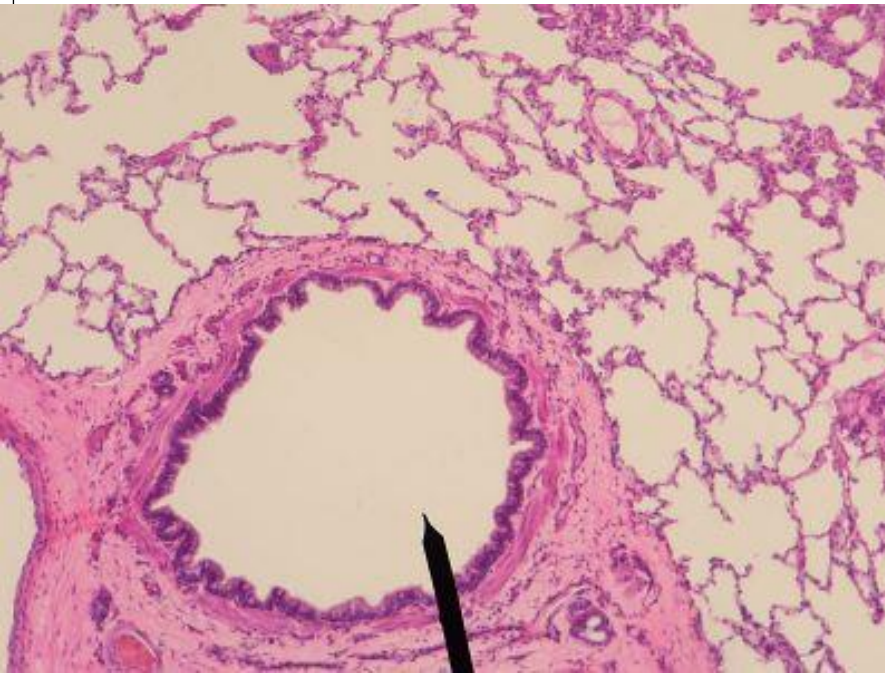
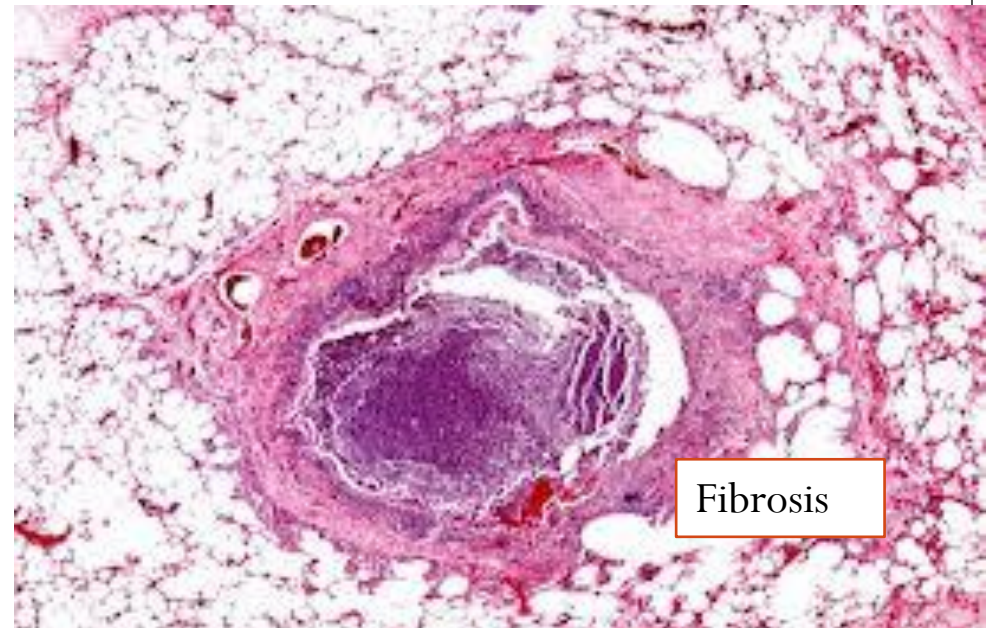
- Usually affects lower lobes bilaterally (vertical airways).
- Dilated airways up to four times of normal, reaching the pleura.
- Acute and chronic inflammation (neutrophils, lymphocytes, histiocytes and plasma cells)
- Necrosis and ulceration in the wall of the bronchi and bronchioles with loss of cilia, squamous metaplasia and fibrosis.



- Normal



- bronchiectasis



Bronchiectasis

- **Clinical course:**
 - Severe persistent cough with sputum (mucopurulent, fetid sputum) sometimes with blood.
 - Clubbing of fingers.
 - If severe, obstructive pulmonary function develops.
 - Other complications: metastatic brain abscess and amyloidosis

Bronchiectasis:

Dilatation of bronchi and bronchioles secondary to chronic inflammation

Causes

- Infection/ Necrotizing pneumonia
- Obstruction
- Congenital (Cystic fibrosis, Kartagener's Syndrome)

Clinical features

- Severe persistent cough with sputum (mucopurulent sputum) sometime with blood.
- Clubbing of fingers.

complications

- If severe, obstructive pulmonary disease develops.
- Lung Abscess
- Rare complications: metastatic brain abscess and amyloidosis.