

RESTRICTIVE vs. obstructive

lung disease

GENERAL PATHOLOGICAL CHARACTERISTICS

- lung expansion is restricted
- primarily due to decreased lung or chest wall compliance or muscular dysfunction
- primarily affects deep **inspiration**
- difficulty getting air **in**

- increased airway obstruction/resistance
- increased work of breathing
 - ↳ hyperventilation
- decreased **expiratory** flow
- difficulty getting air **out**

CLINICAL MANIFESTATIONS

- decreased lung volumes/capacities
- dyspnea
- tachypnea
- increased work of breathing → fatigue
- impaired gas exchange

- hyperinflated lungs & flattened diaphragm
- dyspnea on exertion
- wheezing
- increased work of breathing → paradoxical
- chronic cough & expectoration of mucus

VOLUMES, CAPACITIES, & SPIROMETRY

- ↓ IRV, TV, ERV
- ↓ VC, RV
- ↓ TLC
- ↓ FEV₁
- ↓ FVC
- ↑ FEV₁/FVC (%)

- ↓ ERV, VC
- ↑ TV, RV
- ↑ TLC
- ↓ FEV₁
- ↓ FVC
- ↓ FEV₁/FVC (%)

I:E ratio → 1:1

I:E ratio → 1:3 or 1:4

CAUSES

PULMONARY

- interstitial pulmonary fibrosis
- pneumonia
- TB
- tumors
- atelectasis
- pleural effusion
- pneumothorax
- pulmonary emboli
- pulmonary edema
- acute respiratory distress syndrome

EXTRAPULMONARY

- chest wall trauma/surgery
- NM disorders
- pectus carinatum
- scoliosis
- ankylosing spondylitis
- diaphragmatic paralysis
- obesity

- inflammation of airway
- airway thickening over time
- increased mucus
- constriction of the bronchiole walls (bronchospasms)

PATHOLOGIES



INTERSTITIAL PULMONARY FIBROSIS

- affects alveoli, capillaries, & connective tissue
- may be idiopathic, viral, genetic, or immune system related
- **pathology**: chronic inflammation injures tissue → normal tissue is now fibrous → the spaces in alveoli are thickening → loss of space in alveoli
- **symptoms**: fatigue, dyspnea on exertion, tachypnea, cough
- **signs**: impaired PFTs & blood gases (hypoxemia)
- dx with CT scans, chest x-rays, lung biopsy
- **treatment**:
 - ↳ meds → O₂, glucocorticosteroids, anti-inflammatories
 - ↳ mechanical ventilation
 - ↳ lung transplant
 - ↳ physical therapy: want to decrease the progression
 - breathing techniques
 - OOB mobilization
 - walking program w/ O₂
 - strength training
 - inspiratory muscle training

ATELECTASIS

- alveolar collapse causes lung to collapse
- **causes**: airway obstruction, restrictive conditions
- **kinds**: compression, contraction, resorption
- **treatment**:
 - ↳ OOB mobilization
 - ↳ incentive spirometry (often)
 - ↳ splinted coughing
 - ↳ breathing techniques

PNEUMONIA

- inflammatory process of lung parenchyma
- **causes**: bacteria, virus, fungus, aspiration
- **pathology**: usually localized in a segment/lobe
 - ↳ alveolar edema & congestion
 - ↳ alveolar consolidation (from inflammation & mucus)
 - ↳ decreased lung compliance
- **signs & symptoms**: dyspnea, cough, pleuritic chest pain, fever, hypoxemia
- **treatment**: meds, incentive spirometry, early mobilization, airway clearance & breathing techniques

CHRONIC BRONCHITIS (COPD)

- productive cough condition that lasts > 3 months for at least 2 consecutive years
- **causes**: (response to chronic irritation) smoking, pollution, occupational exposure
- **signs & symptoms**: productive cough, exertional dyspnea, barrel-shaped chest, wheezes, LE edema, blue bloaters, cor pulmonale, prone to pulmonary infections, ↑ symptoms with irritants (cold / damp weather, etc.)
- **pathology**: hypertrophy of mucus glands → ↑ mucus secretion → mucosal edema → bronchospasm & air trapping → cilia unable to clear secretions → chronic infections

EMPHYSEMA (COPD)

- enlargement & destruction of alveoli
- **causes**: disruption of elastic fiber network of lungs
 - ↳ 95% from chronic bronchitis, smoking, or air pollutants
 - ↳ 5% from genetic disorder (α -1 anti-trypsin)
- **types**: * defined by anatomical location
 - ↳ **centriobular emphysema**: inflammation causes destruction in the terminal bronchioles
 - most common type
 - alveolar sac intact but enlarged
 - ↳ **panlobular emphysema**: destroys air spaces of entire alveoli
 - bronchioles might be okay / not as inflamed
 - ↳ **paraseptal emphysema**: destroys alveoli along lung periphery
 - can cause spontaneous pneumothorax
 - around alveoli, NOT inside
- **signs & symptoms**: progressive & exertional dyspnea, cough (variable productivity), barrel-shaped chest, hypertrophied accessory muscles, pursed-lipped breathing, ↓ FEV₁ & ↑ TLC, pink puffers, thin, cachectic, ↑ symptoms w/ acute respiratory infection, digital clubbing, postures that stabilize UE's.

ASTHMA

- airway hyper-reactivity to various external/ internal stimuli
- recurrent episodes of intermittent, reversible airway obstruction

PNEUMOTHORAX

- accumulation of air/gas in pleural space
 - ↳ can lead to atelectasis (but atelectasis CAN'T cause a pneumothorax)
- **causes**: defect in visceral/parietal pleura (see types)
- causes the affected lung to collapse
- **types**:
 - ↳ **primary spontaneous**: no underlying cause
 - usually young, tall, thin men
 - ↳ **secondary spontaneous**: due to COPD, blebs, or bullae
 - most common type
 - ↳ **traumatic**: due to central line (iatrogenic), gm shot, knife wound, rib fracture
 - ↳ **tension**: air enters pleural space but can't escape
 - medical emergency
- **signs & symptoms**: acute dyspnea, pleuritic pain, hypoxemia, absent/diminished breath sounds, hyper-resonant breath sounds, tracheal deviation
- **treatment**: O₂, chest tube drainage, incentive spirometry

PULMONARY EDEMA

- pulmonary congestion that restricts air-flow
- **causes**: L-sided heart failure, ARD's, high altitude sickness
- **signs & symptoms**: severe dyspnea, anxiety, tachypnea, crackles, coughing, hypoxemia
- **treatment**: diuretics, oxygen, upright positioning

PULMONARY EMBOLUS

- embolus causing occlusion of blood flow through the pulmonary A
- **causes**: R side of the heart, DVT, non-thrombotic material
- **characteristics**: ↓ pulmonary blood flow, ↑ pulmonary vascular resistance
- **signs & symptoms**: rapid onset dyspnea, pleuritic chest pain, anxiety, restlessness, apprehension, tachypnea, hypoxemia, ↓ PFT's
- **diagnostic tests**: V/Q scan, pulmonary angiography, D-dimer blood test (most common)
- **treatments**: prevention of DVT's, meds, surgery

ACUTE RESPIRATORY DISTRESS SYNDROME

- acute/rapid onset of respiratory failure
- **causes**: acute extensive lung inflammation (trauma, sepsis, drug overdose, blood transfusions, pneumonia)
- **characteristics**: hypoxemia, pulmonary edema, atelectasis, ↓ surfactant, ↓ compliance
- **treatment**: ICU monitoring, mechanical ventilation, meds, positioning

- inflammation on bronchiole walls
- **signs & symptoms**: recurrent, paroxysmal coughing attacks, chest tightness, difficulty breathing, & wheezing, symptom-free between attacks, ↓ FEV₁, ↑ TLC
- not all asthma are obstructive (can also be restrictive)

BRONCHIECTASIS

- permanent abnormal dilation & distortion of one or more medium sized bronchi/bronchioles
- **causes**: destruction of elastic & muscular components of the bronchiole wall, usually because of another pathology
- **pathology**: necrotizing infection → abnormal bronchial dilation & spasms → destruction/fragmentation of the bronchial wall
 - ↳ usually in L lower lobe
- **signs & symptoms**: productive cough, purulent sputum, hemoptysis, crackles, wheezes, dullness to percussion, ↓ breath sounds, digital clubbing, dyspnea, ↓ FEV₁, ↑ RV, hypoxemia, hypercapnia, acidosis, pulmonary HTN, cor pulmonale, R CHF

CYSTIC FIBROSIS

- genetic disorder → dysfunction of exocrine glands
 - ↳ autosomal recessive disorder
 - ↳ multi-organ involvement → "salty disease"
 - pancreas → steatorrhea
 - liver → obstructive jaundice, gallstones
 - GI tract → fecal impaction, intussusception
 - reproductive organs → difficulty w/ pregnancy, vas deferens obstruction
- most common fatal genetic disease of caucasians
- **pathology**: thick secretions plug airways → bacterial lung infections → small airway inflammation → larger airways → hyperplasia of mucus-secreting cells
- **signs & symptoms**: thick, tenacious mucus, increased sputum, chronic cough, increased RR & SOB, crackles, wheezes, digital clubbing, hyperinflation of lungs, ↑ RV, ↓ FEV₁, V/Q mismatching, pulmonary HTN → cor pulmonale, hypercapnia, respiratory acidosis

PT Management for All obstructive diseases:

- postural drainage & ACBT
- breathing control training
- effective Huff/cough
- improve thoracic mobility & exercise tolerance
- patient education (self-management, relaxation)