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Ch 27 Pulmonary Disorders

Types of pulmonary diseases

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| **Respiratory system diseases** | **NCLEX-RN** | **Khan Academy**

COPD (Chronic Obstructive Pulmonary Disease), Chronic Bronchitis, Emphysema-NCLEX Part 1 *Lung Pleura - Clinical Anatomy and Physiology*

Chronic Obstructive Pulmonary Diseases (COPD) | Pulmonology Chronic Pulmonary Pathophysiology 2: COPD Pathology Chronic Pulmonary Pathophysiology 1: Types of Conditions

Bronchiectasis *Restrictive Lung Diseases; An Introduction | Pulmonary Medicine* **Chronic Pulmonary Pathophysiology 4: Other Obstructive Conditions**

Pulmonary Embolism (PE)

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~~Approach Well With Stage 4~~

~~Lung Cancer Pleural~~

~~Effusions Causes,~~

~~Diagnosis, Symptoms,~~

~~Treatment~~ **Life with lung**

cancer: A patient's story

Obstructive vs. Restrictive

Lung Disease || USMLE

Oncogenetics - Mechanism of

Cancer (tumor suppressor

genes and oncogenes) ??????

????? Lung Cancer - ??????

???? - S.M.V The Pleurae and

Pleural Fluid **weekly routine**

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Emphysema) Explained Clearly

—Pathophysiology \u0026

Diagnosis **Pleural Effusion**

Explained Clearly - Causes,

Pathophysiology, Symptoms,

Treatment, Chronic

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Obstructive Pulmonary
Disease: Pathophysiology
& Risk Factors (PART 1)
Pathophysiology of COPD |
Chronic Bronchitis &
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Edition. Dyspnea is a

sensation of breathlessness or subjective "shortness of

breath.". It. defi es strict defi nition with many

individual descriptions and varies widely. among

patients for comparable

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helps in distinguishing the
various pulmonary and extra

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y Vol 6 July 2018. Review

Pulmonary alveolar proteinosis in adults: pathophysiology and clinical approach. Anupam Kumar, Basem Abdelmalak, Yoshikazu Inoue, Daniel A Culver.

Pulmonary alveolar proteinosis (PAP) is a diffuse lung disease that results from the accumulation of lipoproteinaceous material in the alveoli and alveolar macrophages due to abnormal surfactant homeostasis.

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our understanding of the pathogenesis, natural history, and treatment of pulmonary alveolar proteinosis (PAP) has evolved considerably. PAP is characterised by the

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accumulation of surfactant in alveolar macrophages and alveoli, resulting in impaired gas exchange.

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of the pathophysiology of dyspnoea is essential. A structured clinical approach based on a thorough medical history and clinical examination is key to making the correct diagnosis. Special investigations play secondary and supplementary roles in the diagnosis; they are guided by the history and examination and

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Pathophysiology and a clinical approach

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Anticoagulant treatment for acute pulmonary embolism: a

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complement the clear,
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book has been revised and
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to assist problem-based
learning - including body
diagrams annotated with
signs, symptoms and diseases
and a separate index of
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