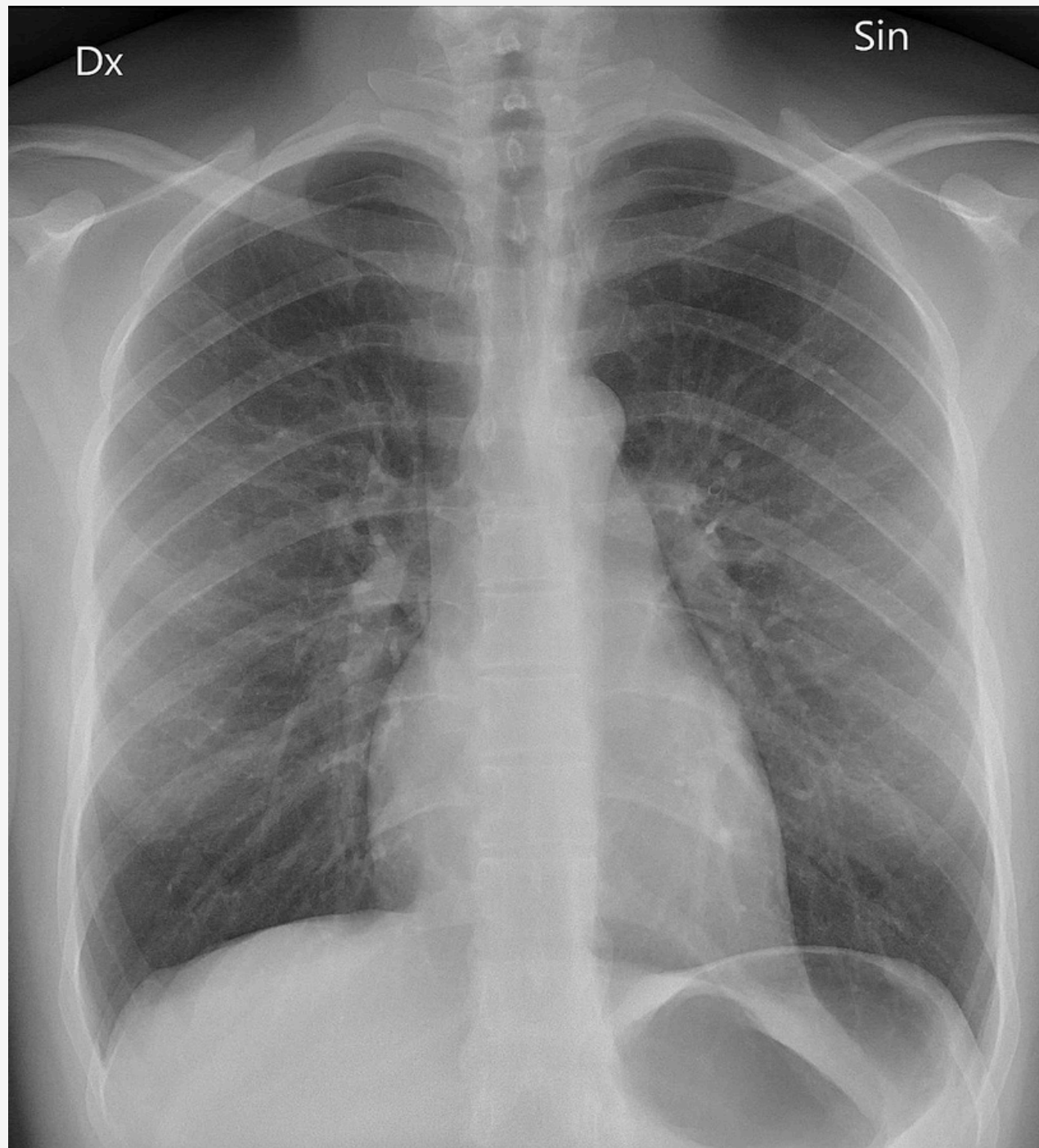


بسم الله الرحمن الرحيم

Chest 2

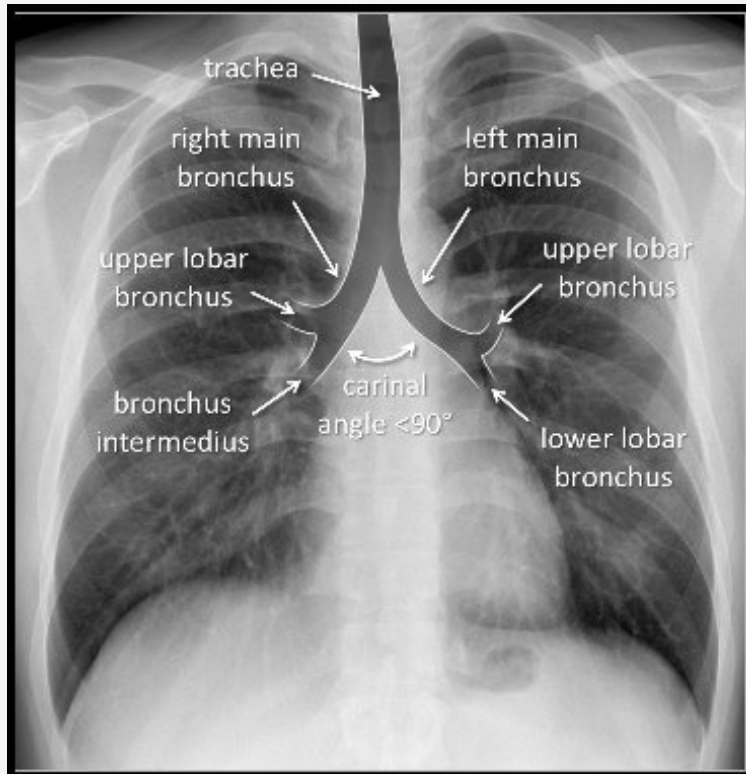
Dr.lina rawashdeh



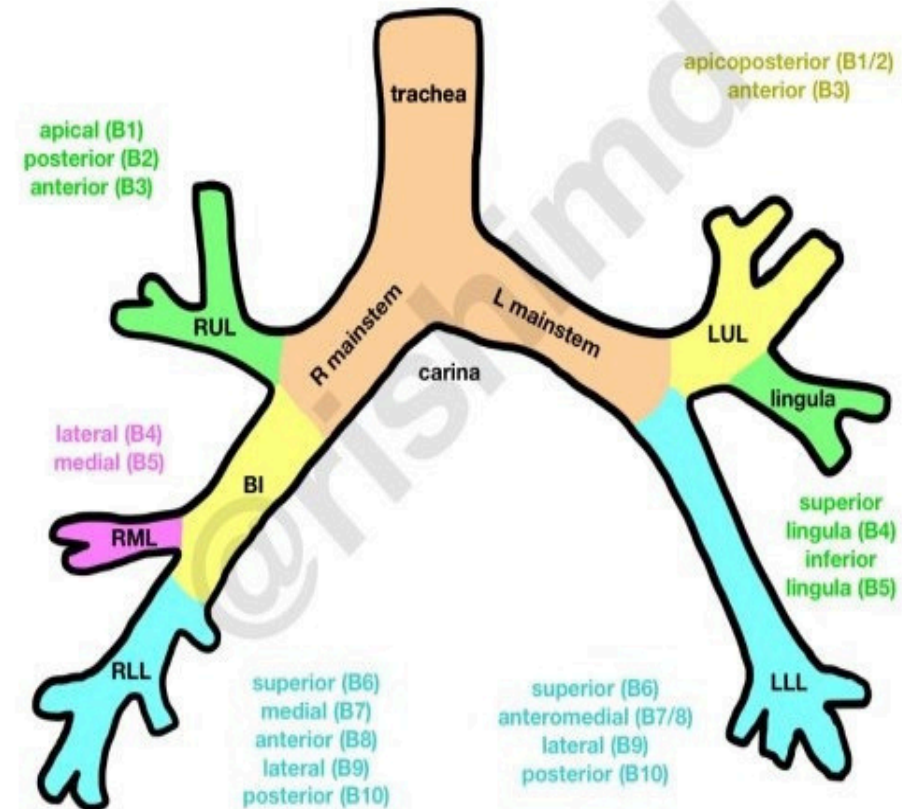


A – Airway and hilum

- Trachea carina, bronchi and hilar structures.



BRONCHOPULMONARY SEGMENTS



● The trachea extends from the inferior margin of the cricoid cartilage (C6) and branches into the right and left main bronchi at the carina, located at the T4 vertebral body level

- Reinforced anteriorly by approximately 20 semicircular, cartilaginous rings
- The trachea divides into the right and left mainstem bronchi which further divide into secondary bronchi (RUL, RML, RLL, LUL, and LLL)

Trachea

Inspect the trachea for evidence of deviation:

The trachea is normally located centrally or deviating very slightly to the right.

If the trachea appears **significantly deviated**, inspect for anything that could be pushing or pulling the trachea. Make sure to inspect for any paratracheal masses and/or lymphadenopathy.

- Causes of true and apparent tracheal deviation

- True tracheal deviation:

—
Pushing of the trachea: large pleural effusion or tension pneumothorax.

Pulling of the trachea: consolidation with associated lobar collapse.

- Apparent tracheal deviation:

Rotation of the patient can give the appearance of apparent tracheal deviation, so as mentioned above, inspect the clavicles to rule out the presence of rotation.

المراد من هذه النسخة هو أن تكون قادرة على التعرف على الـ opacity ← وهذا ما نلاحظه في الـ pleural effusion ← وبعيداً عن الـ reporting نلاحظه في الـ opacity.

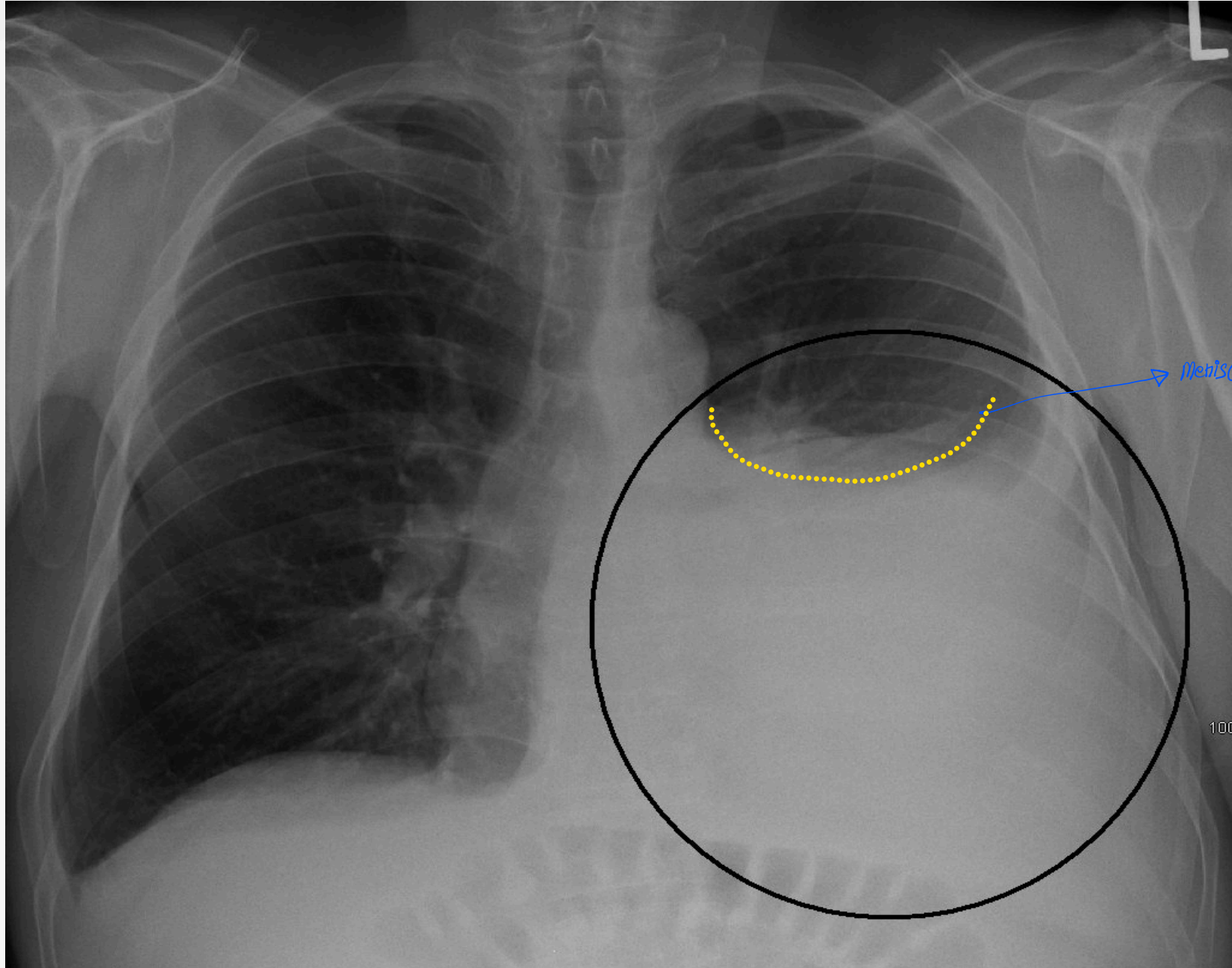
* Reporting :-

- PA CXR

- Lt lower opacity with blunting of Lt costophrenic angle.
- the picture is suggestive of **Lt Side pleural effusion** with decrease lung volume

Meniscus sign.

* و الـ normal point
 Rt lung
 Rt Costo. Angle
 hilum
 bone cage
 No cardiomegaly
 ↓
 و الـ normal point
 ↓
 و الـ normal point
 ↓
 و الـ normal point

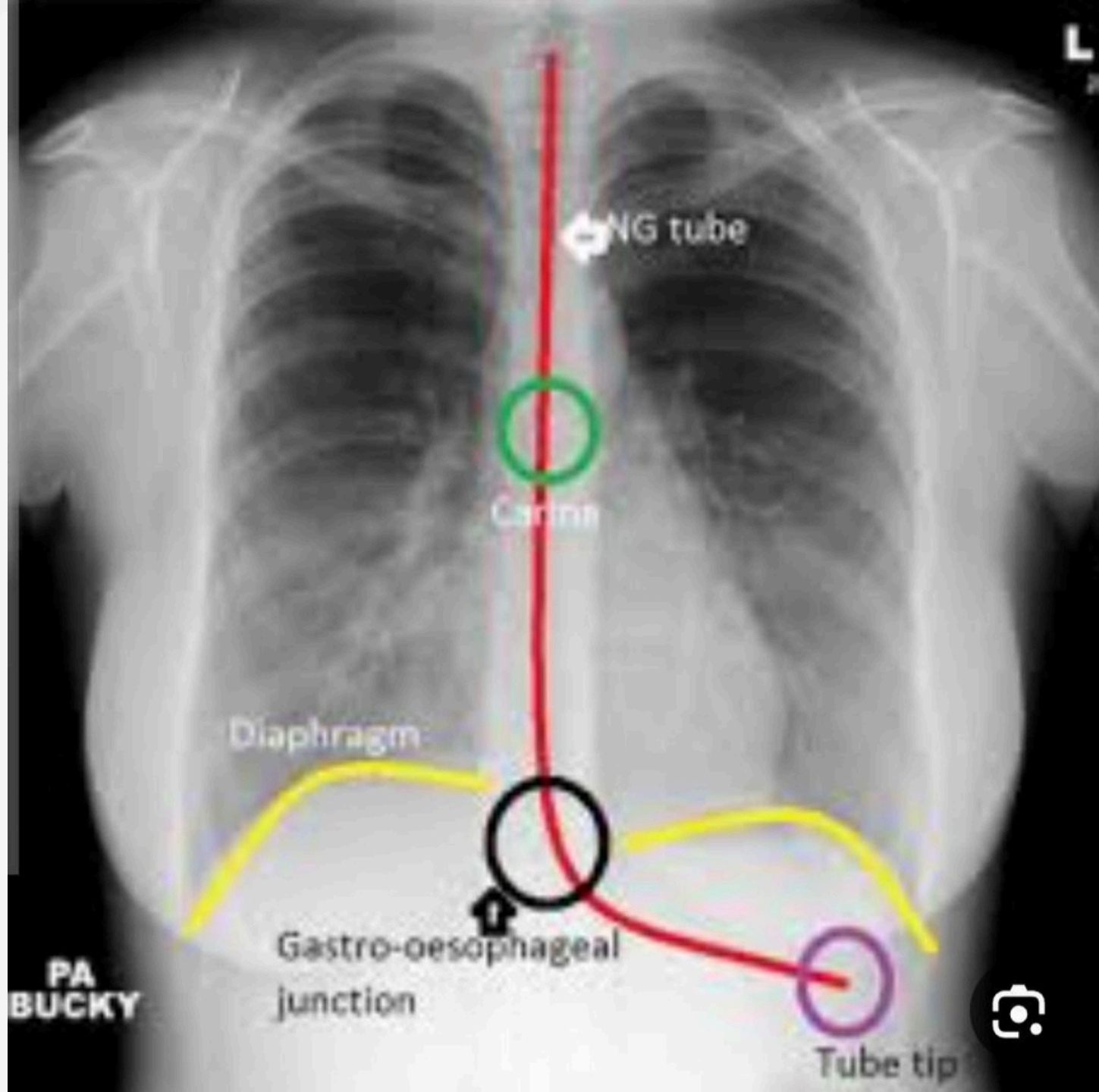


Carina and bronchi

- The carina is cartilage situated at the point at which the trachea divides into the left and right main bronchus.

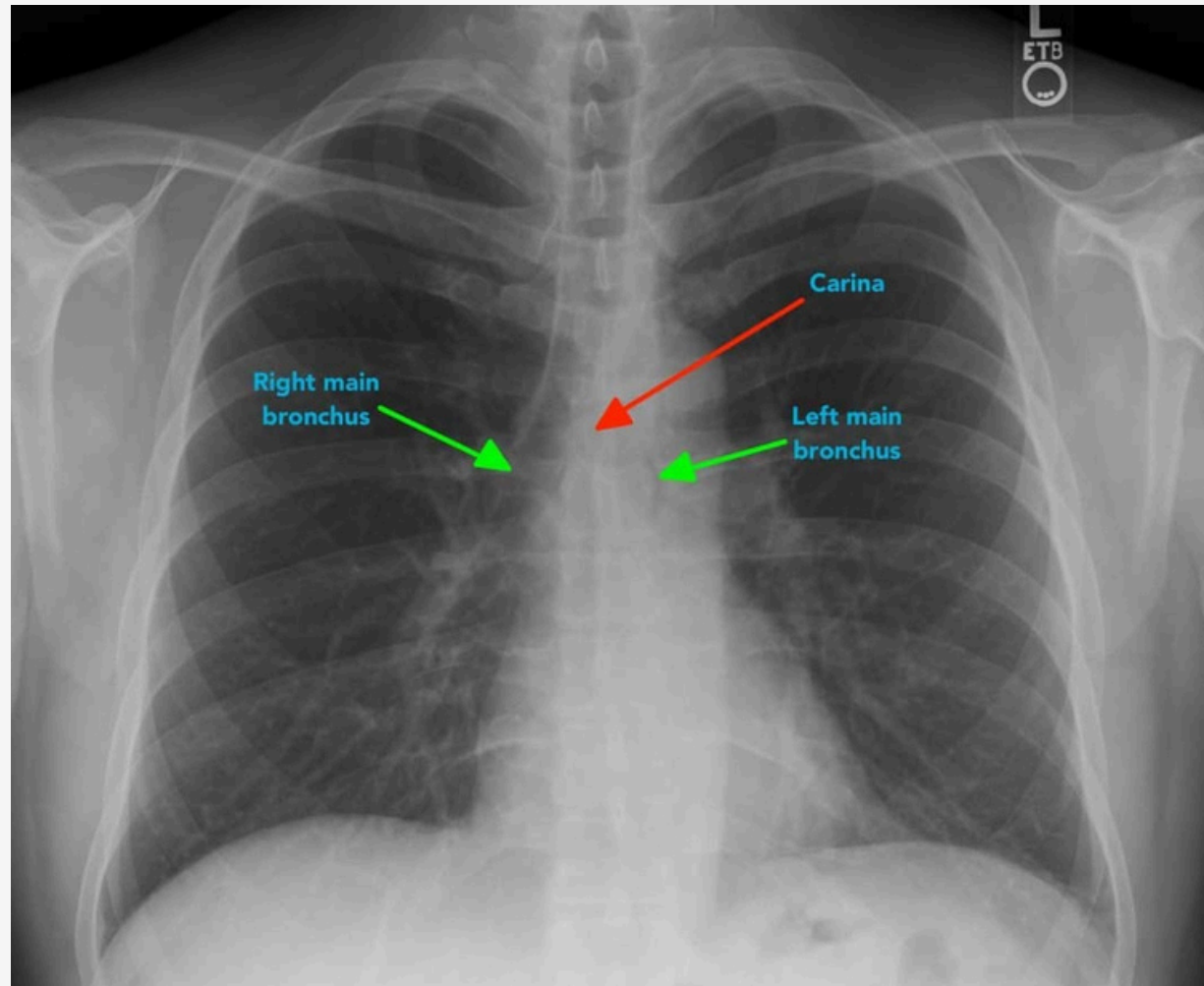
On appropriately exposed chest X-ray, this division should be clearly visible.

- The carina is an important landmark when assessing nasogastric (NG) tube placement, as the NG tube should bisect the carina if it is correctly placed in the gastrointestinal tract.



- The **right** main bronchus is generally **wider, shorter** and more **vertical** than the left main bronchus.
- As a result of this difference in size and orientation, it is **more common** for **inhaled foreign objects to become lodged in the right main bronchus.**

Depending on the quality of the chest X-ray you may be able to see the main bronchi branching into further subdivisions of bronchi.



Bronchiectasis

→ Cuz by
→ Cystic Fibrosis
→ viral Inf
→ PCD → primary ciliary dyskinesia
→ pneumonia

- progressive, irreversible dilation of cartilage-containing bronchi. *with thick wall.*
- Three etiologies of bronchiectasis have been described, with a final common pathway of mucus plugging, superimposed bacterial colonization, and inflammatory response.

Bronchial wall injury, typically from infection or inflammation.

Bronchial lumen obstruction.

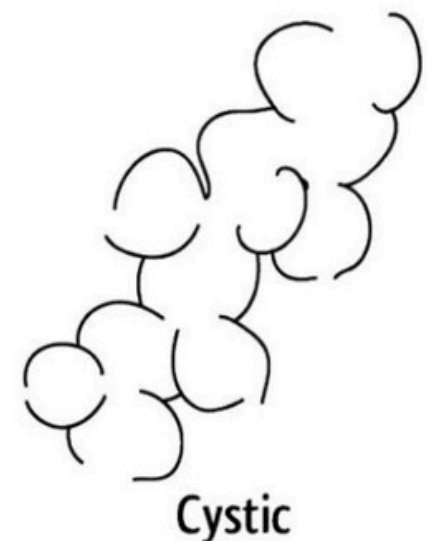
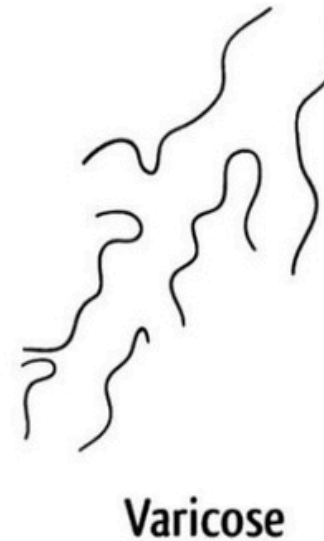
→ Usual Interstitial Fibrosis [UIF] → make Traction bronchiectasis changes.
obstruction → Irreversible dilatation
Infection → Cystic Fibrosis

- Morphologic classification of bronchiectasis is most useful as a rough gauge of severity.

1. **Cylindrical bronchiectasis** (least severe): mild bronchial dilation.

2. **Varicose bronchiectasis** (moderately severe): Bronchi may become beaded and irregular. *→ Tubular*

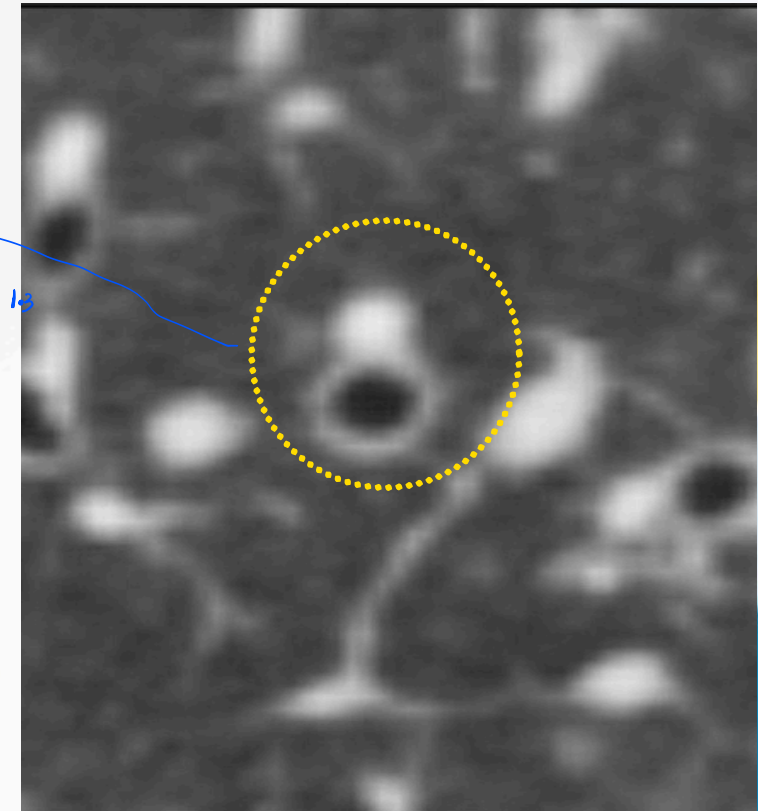
3. **Cystic bronchiectasis** (most severe) : *With Cystic Fibrosis*
Bronchi are markedly enlarged and ballooned, with formation of multiple cysts that may not connect to the airways.



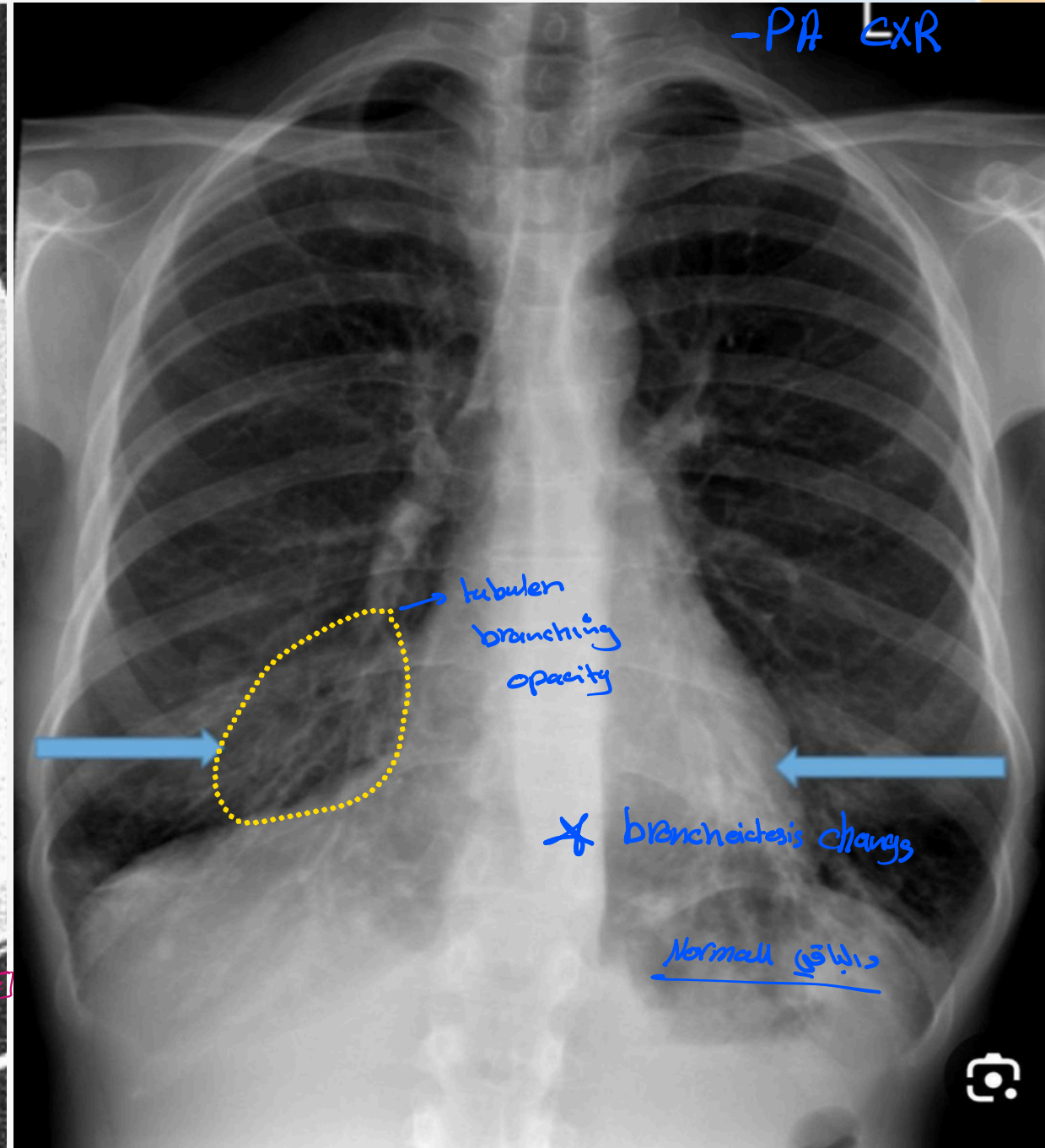
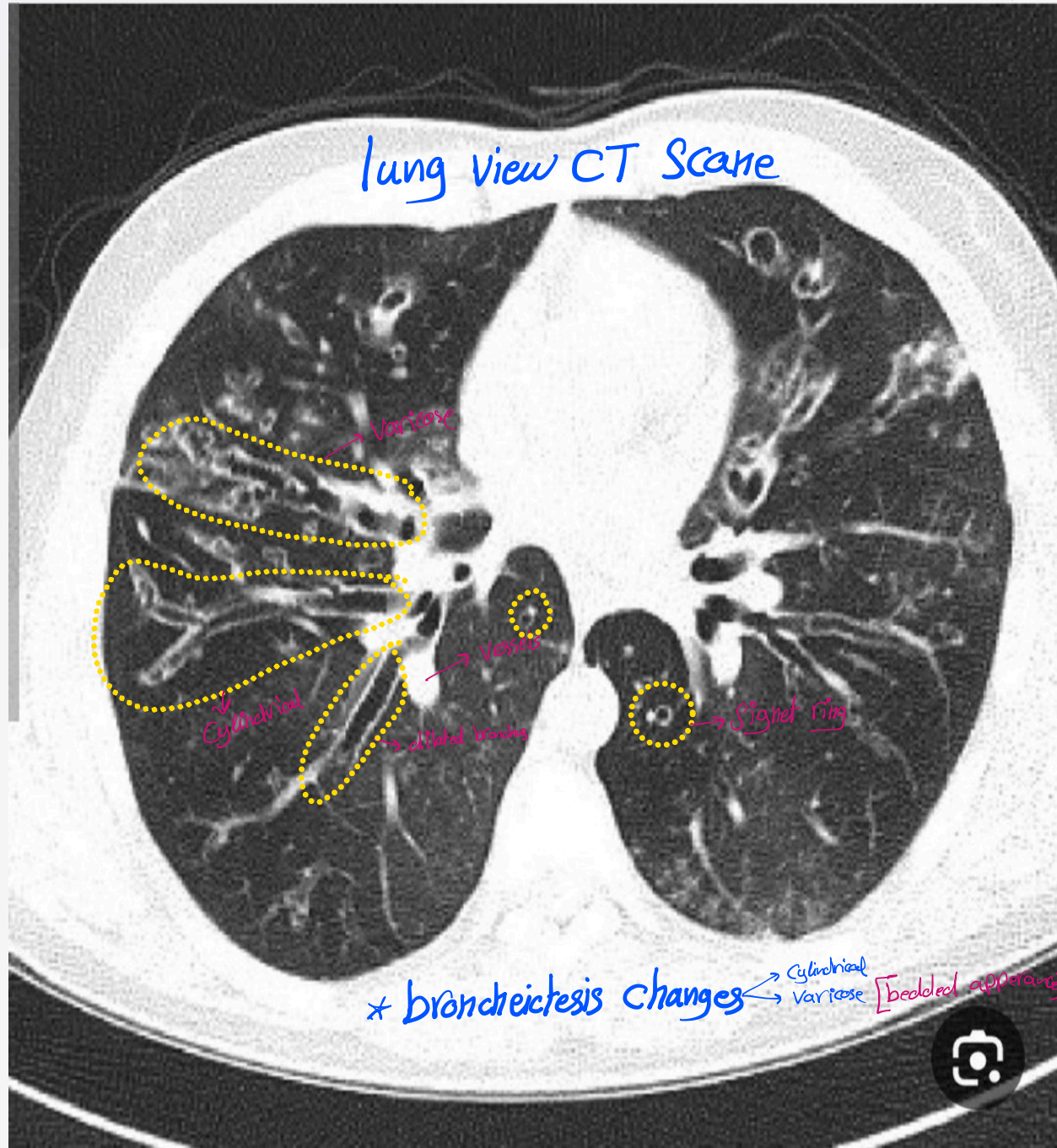
- Radiographic findings depend on severity. In mild cases only tram tracks may be visible, representing thickened bronchial walls causing parallel radiopaque lines resembling tram tracks. In more severe cases there can be extensive cystic change

- CT findings include the signet ring sign

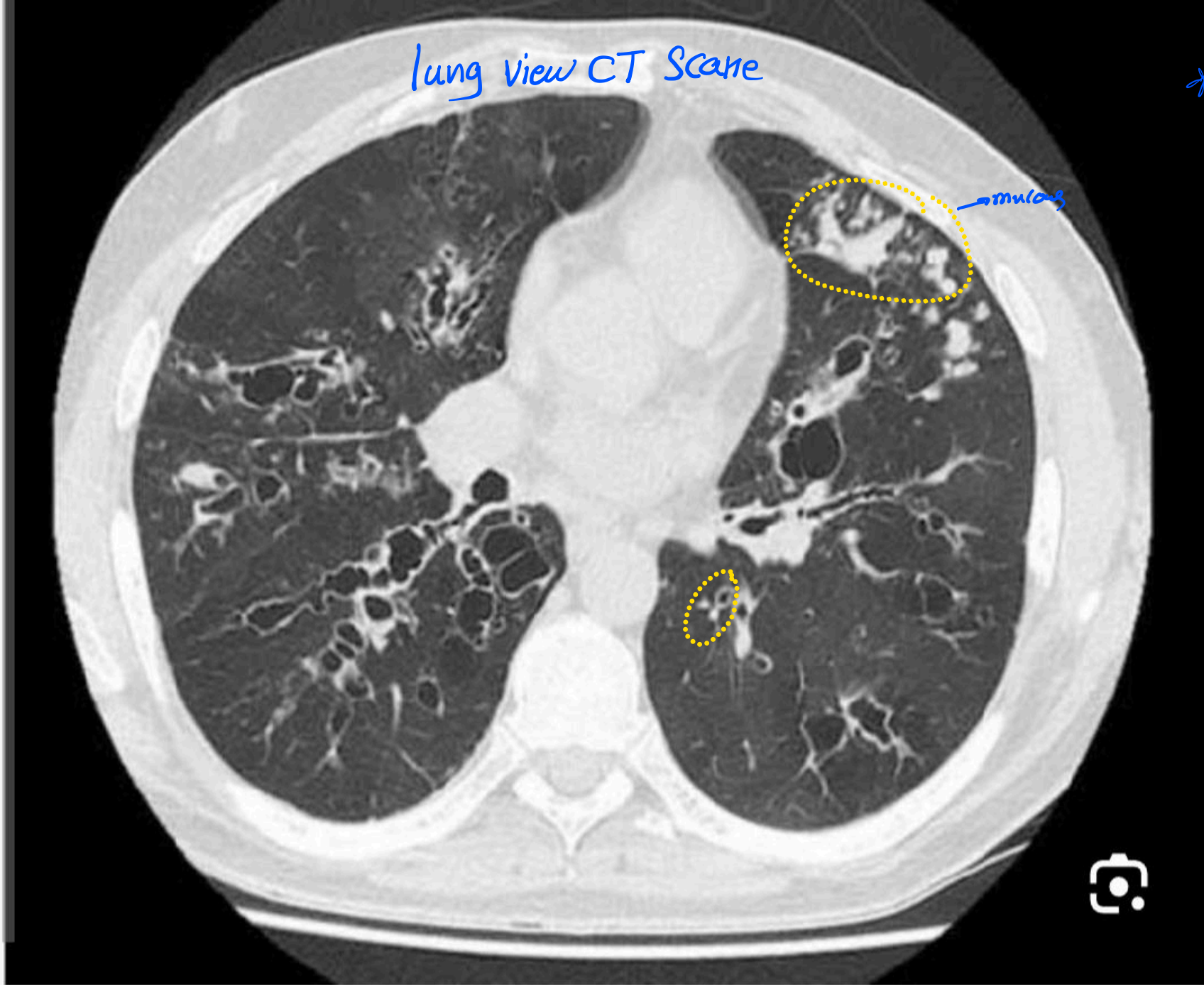
↳ Normally the vessels & bronchus have same size
but here bronchus > vessels mostly in 1



* Cystic Fibrosis affect mostly pancreas & upper lobe lung



lung view CT Scan



→ multiple

* Bronchiectasis change
with → Cystic
→ Beaded appearance [varicose]

* signet ring sign



-lung window

multiple variable
size cysts.

-PA CXR of multiple cystic opacity
in Rt lung lobe.

Cystic fibrosis

- Autosomal recessive genetic disease that affects the exocrine function of the lungs, liver, pancreas, small bowel, sweat glands, and the male genital system. This is resulting in progressive disability and multisystem failure
- Although the entire lung is affected, there is a predilection for:
 1. central (perihilar) distribution
 2. upper lobes
 3. apical segment of lower lobes

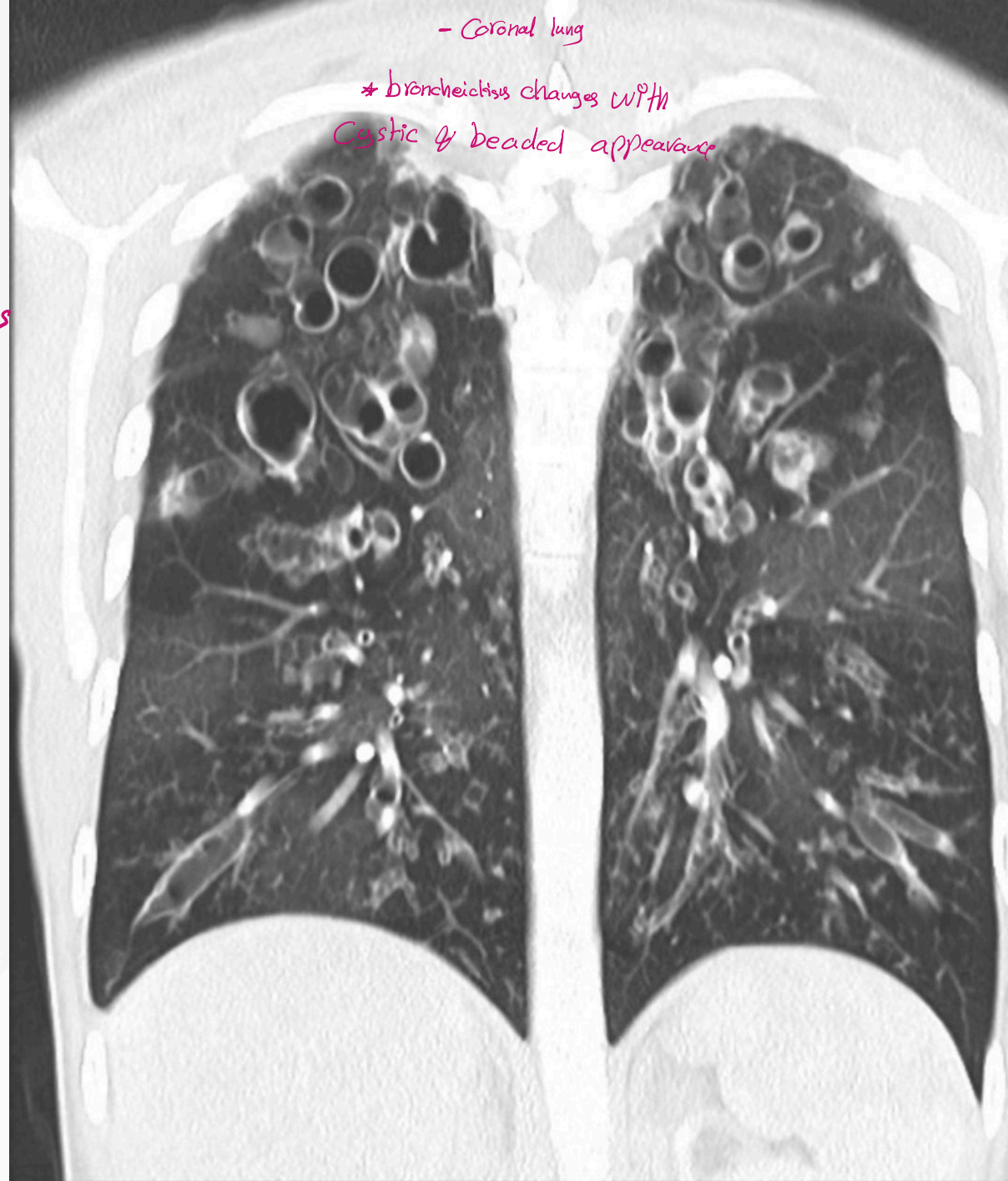
Infertility

R

* PA CXR shows
Tubular branching
of cystic
opacity with
Tram - Trachs
appearance

- Coronal lung

* bronchiectasis changes with
Cystic & beaded appearance



- Chest radiographs are insensitive to the early changes of cystic fibrosis, with changes seen on HRCT in 65% of patients with CF and normal chest radiographs

- Later changes include:

1. Bronchiectasis

2. Hyperinflation

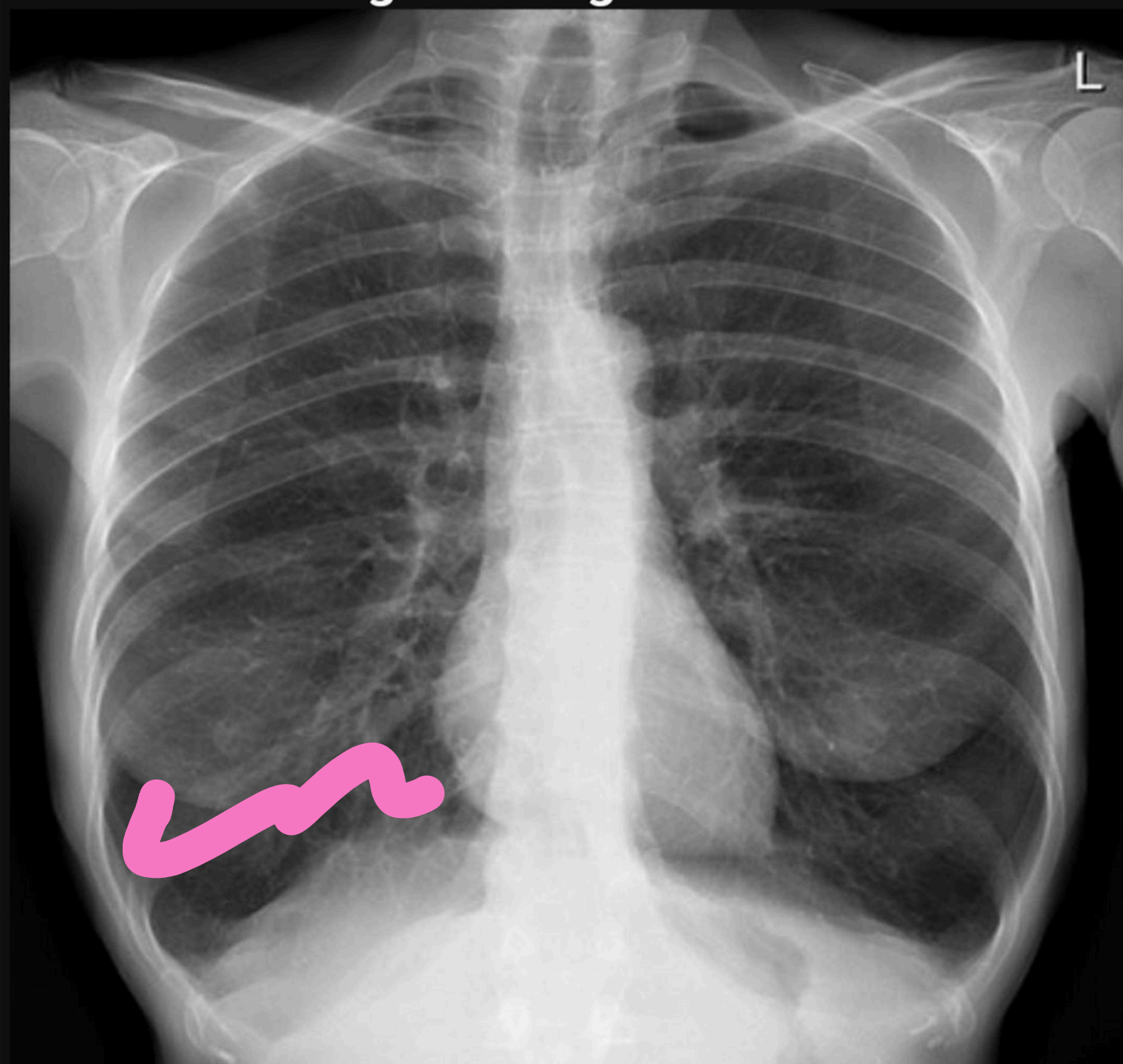
3. lobar collapse

4. pulmonary arterial enlargement due to pulmonary arterial hypertension in patients with long-standing disease



Hyperinflation

- Lung hyperinflation:
- is a common feature of patients with chronic obstructive pulmonary disease (COPD).
- It is also linked to aging and other chronic diseases that cause airflow obstruction.



● Plain radiograph

- Observable features include:

1. flattened hemidiaphragmatic contour

2. air trapping: when comparing two radiographs acquired in maximal inspiration and maximal expiration, the vertical movement of the diaphragm is less than 3 cm

3. appearance of the ribs:

- more than 6 anterior or 10 posterior ribs above the diaphragm level on the midclavicular line

- horizontalisation of ribs

4. increased anteroposterior diameter of the chest, also called barrel chest

5. hyperlucent lungs (i.e. less bronchovascular markings per unit area)

6. presence of air below the heart



* Reporting :-

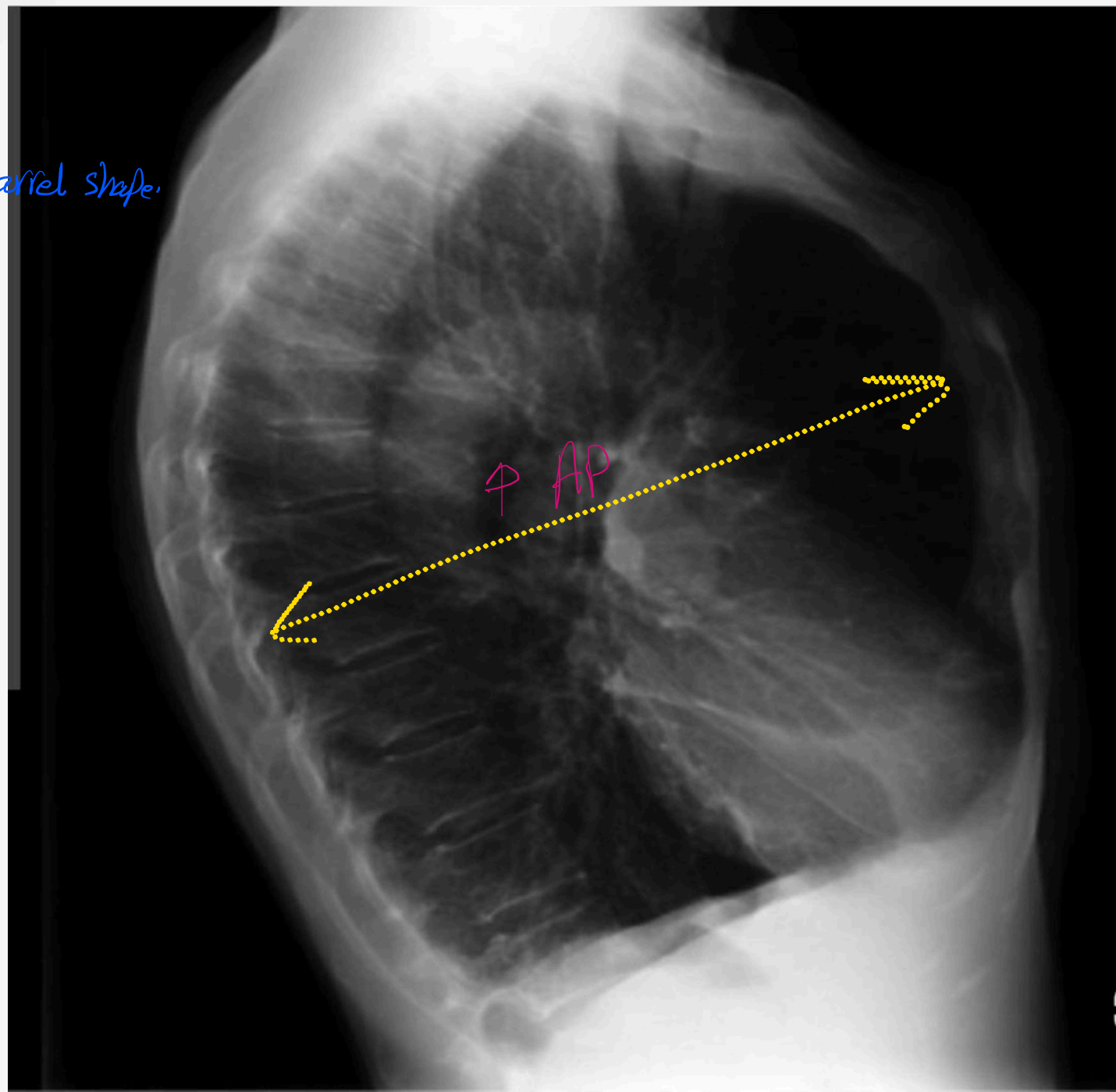
- PA cXR
- Flattening in diaphragm
- ↑ lucency in both lung fields
- Air below Heart
- ribs # posteriorly >10
- the picture suggestive of COPD

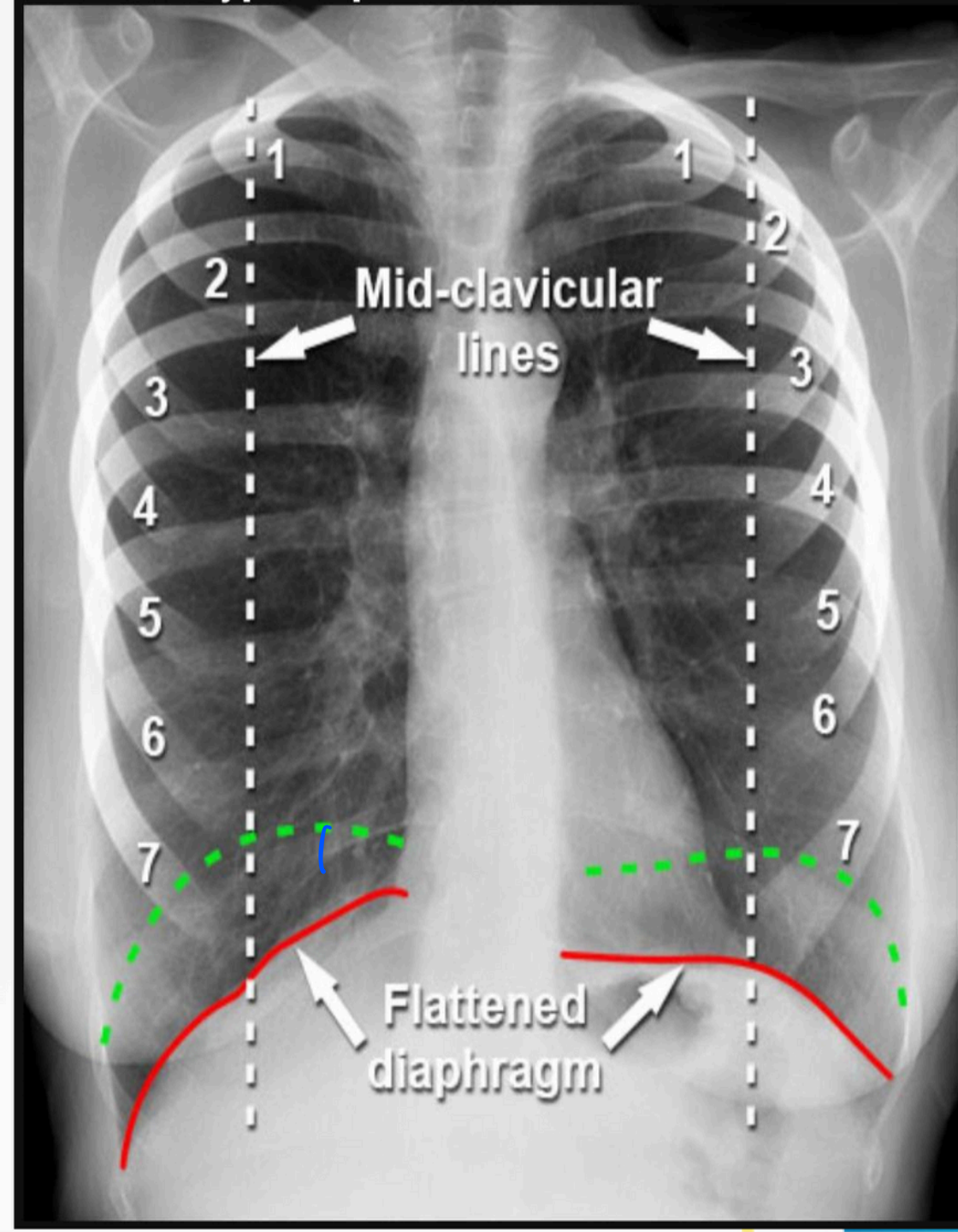
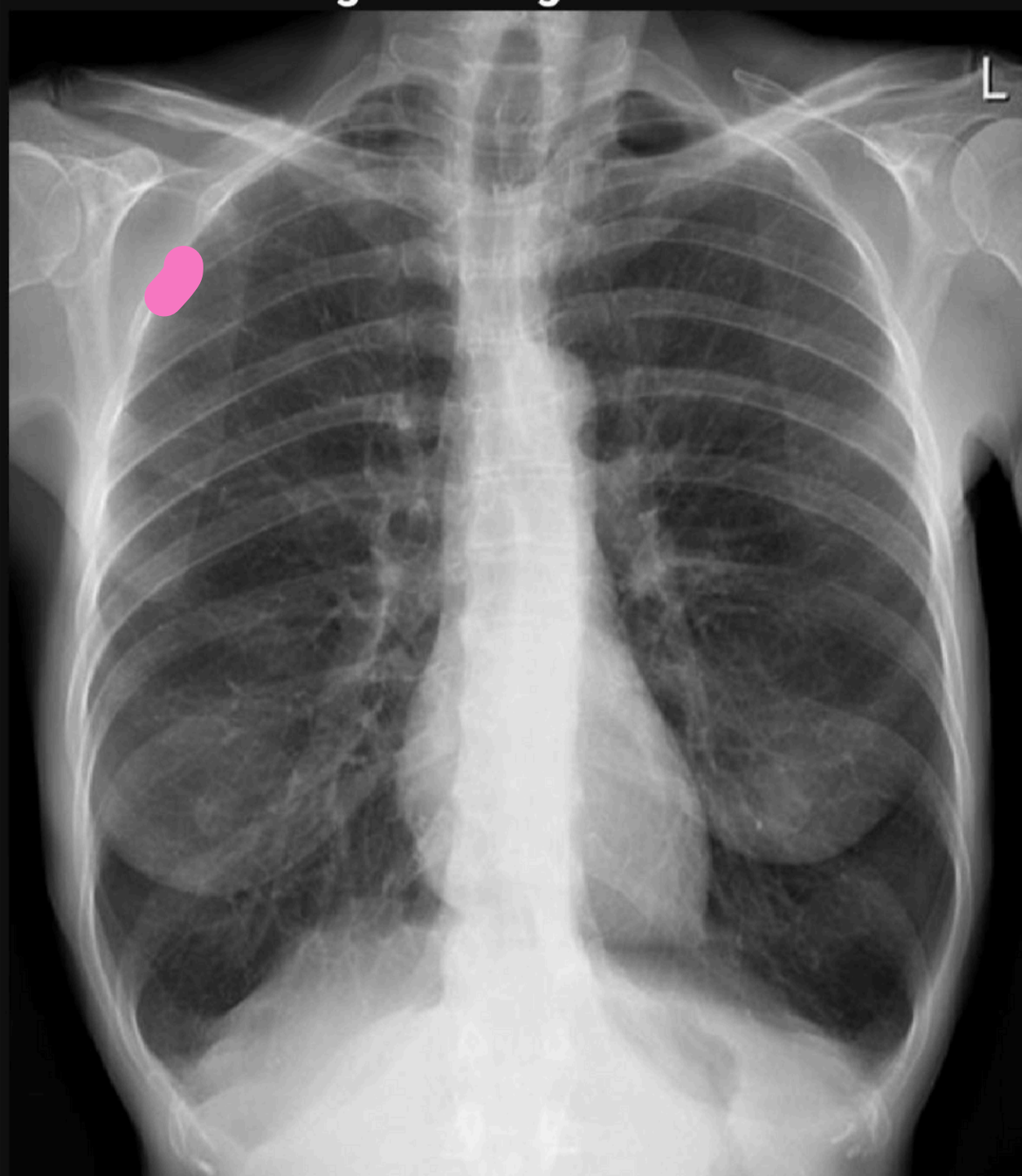
* Normal :-

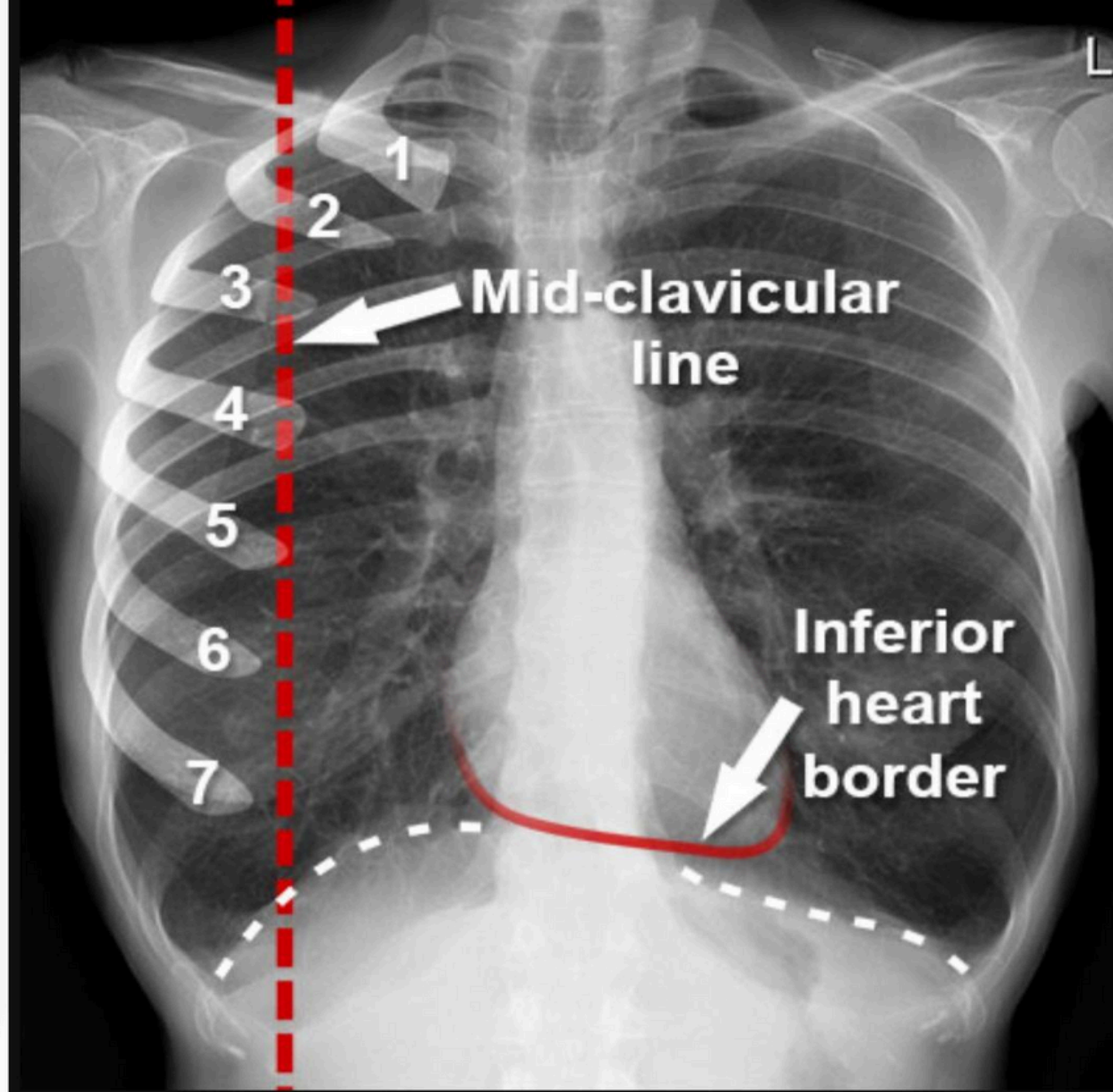
- Normal Trachea
- // Hilum
- // Costophrenic Angle
- // Bony Cage.
- No Cardiomegaly

* Lateral View

- ϕ AP diameter Barrel shape.







Hilar structures

The hilar consist of the main pulmonary vasculature and the major bronchi.

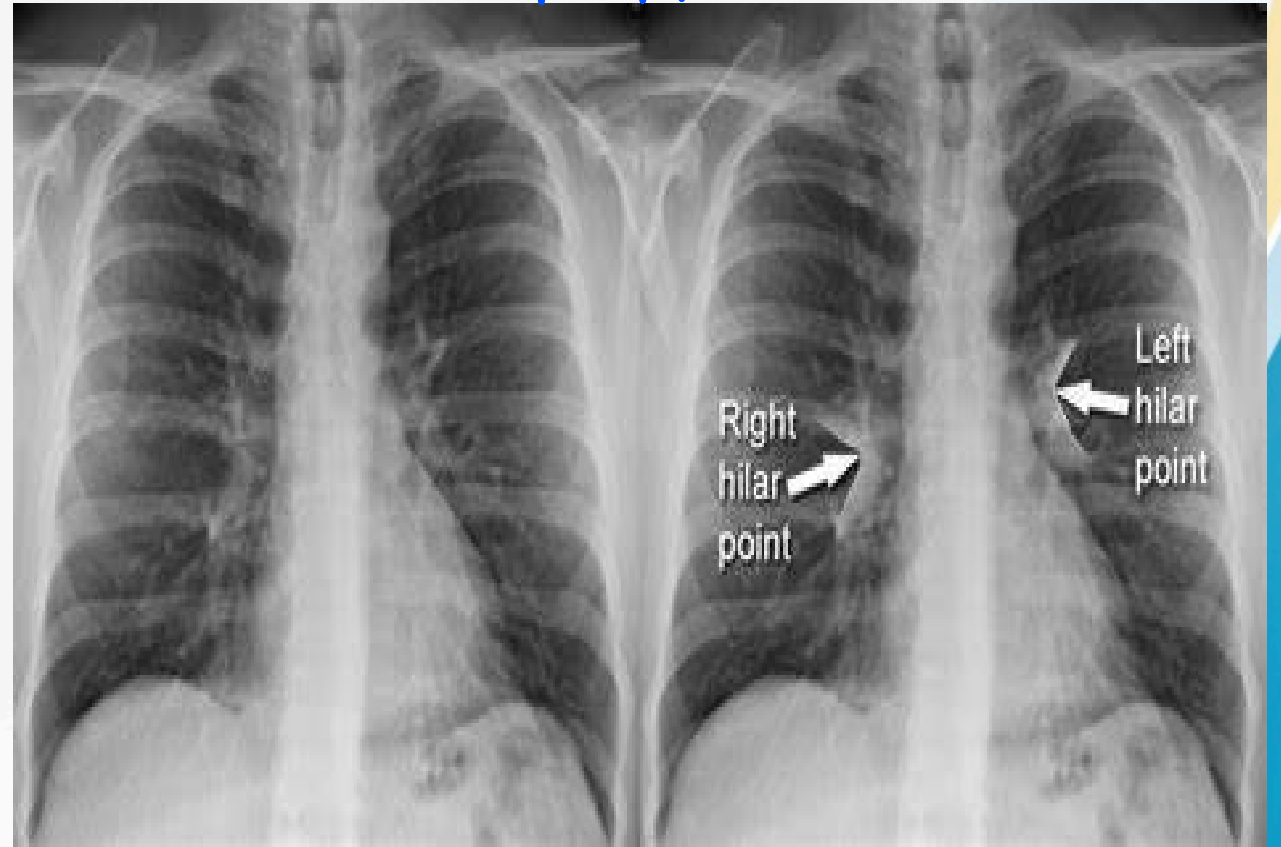
- Each hilar also has a collection of lymph nodes which aren't usually visible in healthy individuals *Sarcoidosis ← Bilateral → Unilateral → Malignancy + TB*

- The left hilum is often positioned slightly higher than the right, but there is a wide degree of variability between individuals.

The hilar are usually the same size, so asymmetry should raise suspicion of pathology.

- The hilar point is also a very important landmark; anatomically it is where the descending pulmonary artery intersects the **superior pulmonary vein**. When this is lost, consider the possibility of a lesion here (e.g. Lung tumour or enlarged lymph nodes).

Normal



Causes of hilar enlargement or abnormal position

- Hilar enlargement can be caused by a number of different pathologies:

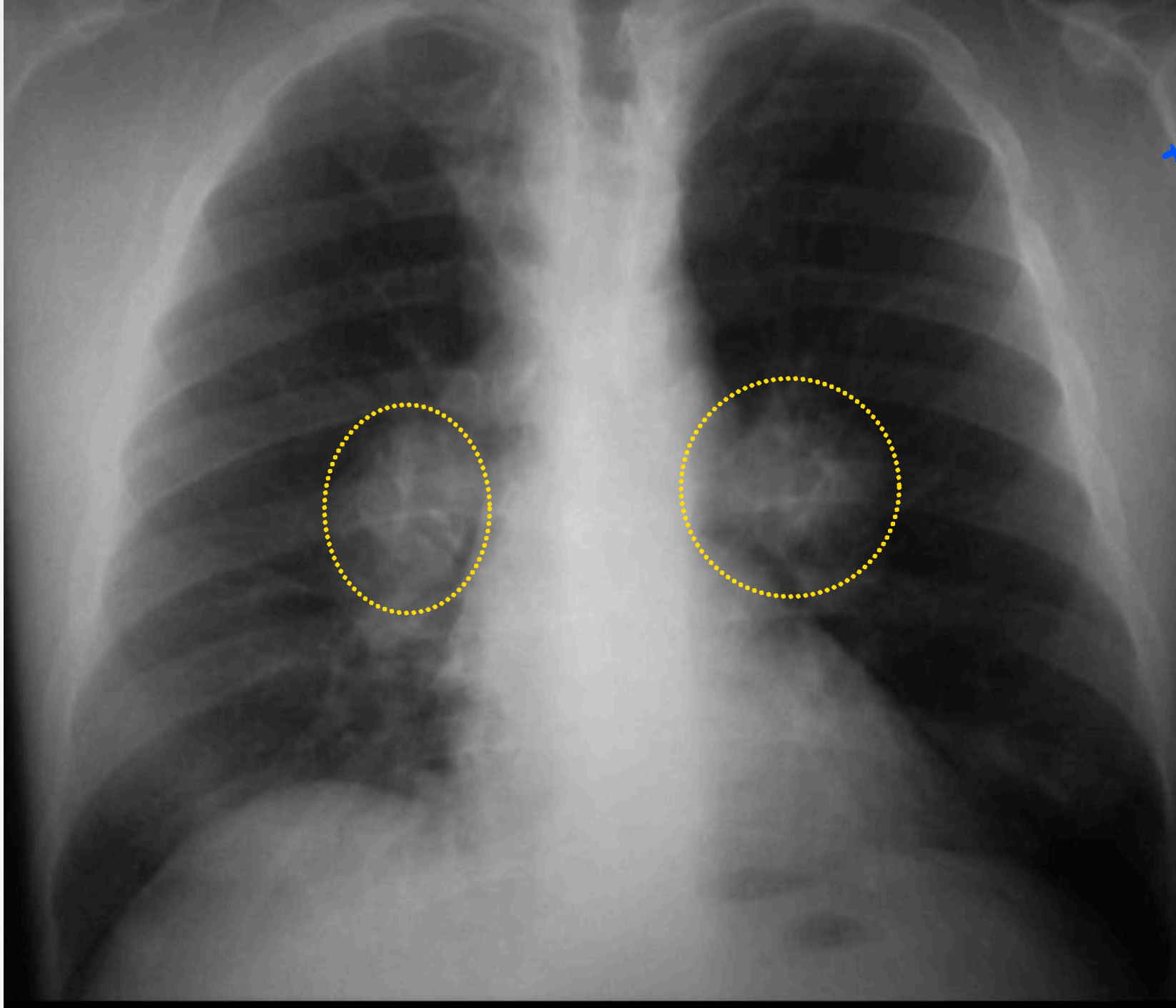
1. Bilateral symmetrical enlargement is typically associated with sarcoidosis.

الحالات التي
تسبب تضخم العقد الليمفاوية
في كلا الجانبين
مثل sarcoidosis
Nodes enlargement.

2. Unilateral/asymmetrical enlargement may be due to underlying malignancy + TB

- Abnormal hilar position can also be due to a range of different pathologies.

You should inspect for evidence of the hilar being pushed (e.g. By an enlarging soft tissue mass) or pulled (e.g. Lobar collapse)



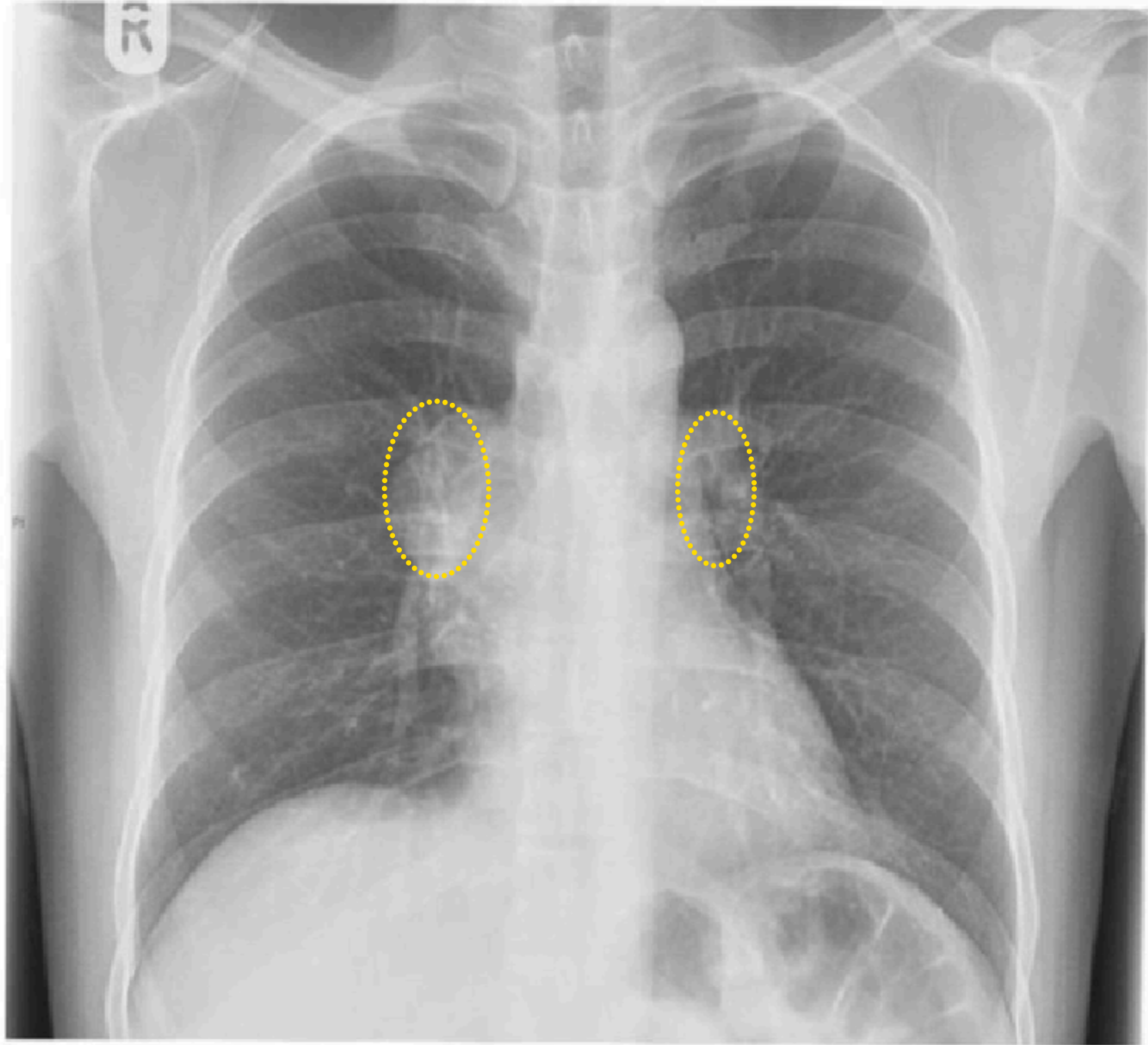
* Bilateral hilar
enlargement.

- PA CXR

- Bilateral hilar lymphadenopathy

+

Normal ⑤



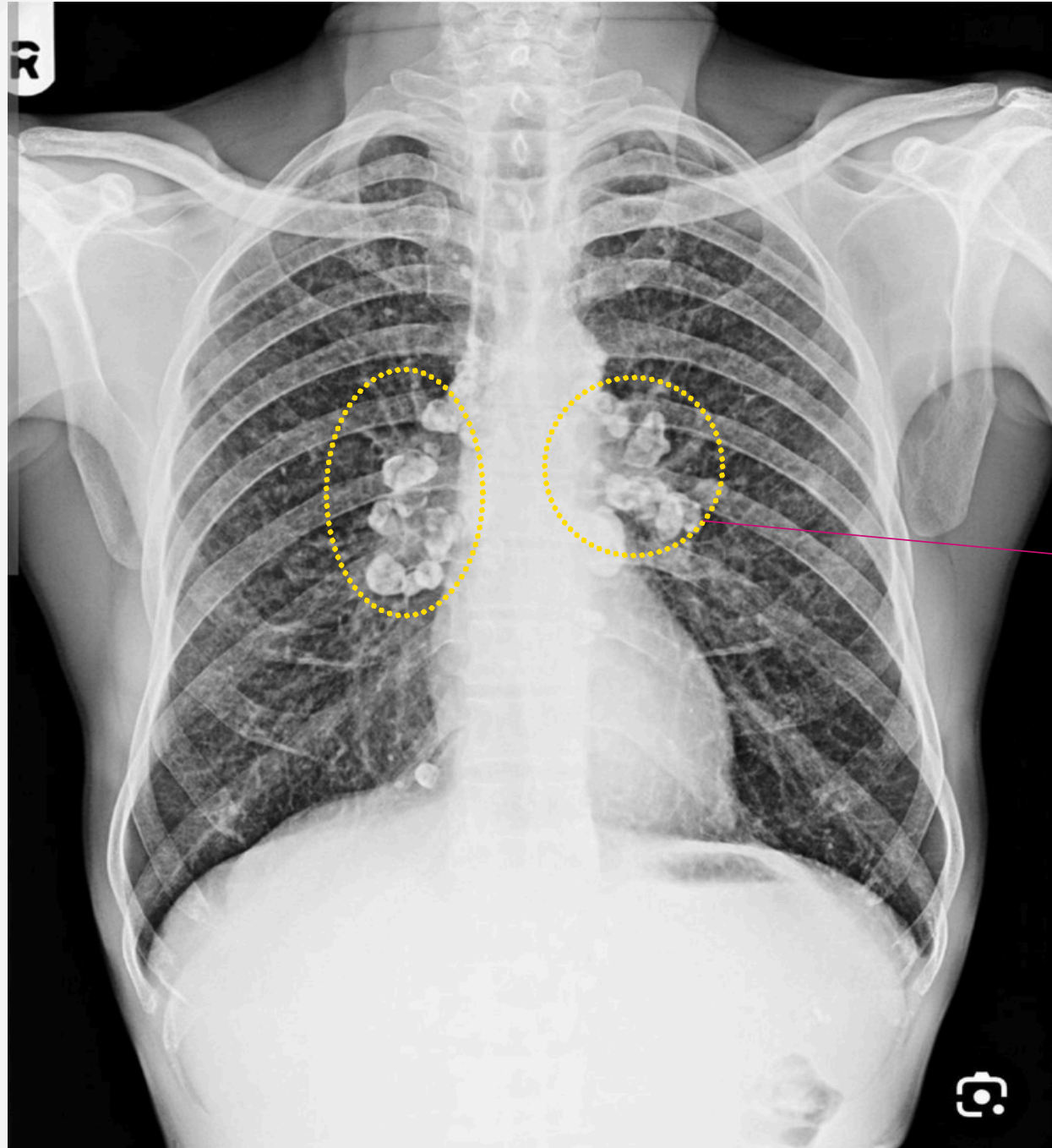
Sarcoidosis

→ Autoimmune disease
↳ middle Age Female.
↳ black more than white

- a non-caseating granulomatous multisystem disease with a wide range of clinical and radiographic manifestations.
- Pulmonary manifestations are present in ~ 90% of patients.
- Pulmonary sarcoidosis may progress to pulmonary fibrosis with honeycombing. → last stage. ④

منعاجاد بصفوف الـ CT

- The most common radiographic finding in sarcoidosis is **symmetric adenopathy**.
- Lymph nodes may contain stippled or eggshell calcification in up to 50%.
- The most common CT finding in sarcoidosis, in addition to ^②**adenopathy**, is upper lobe predominant ^②**perilymphatic nodules** of variable sizes, representing **sarcoid granulomas**.



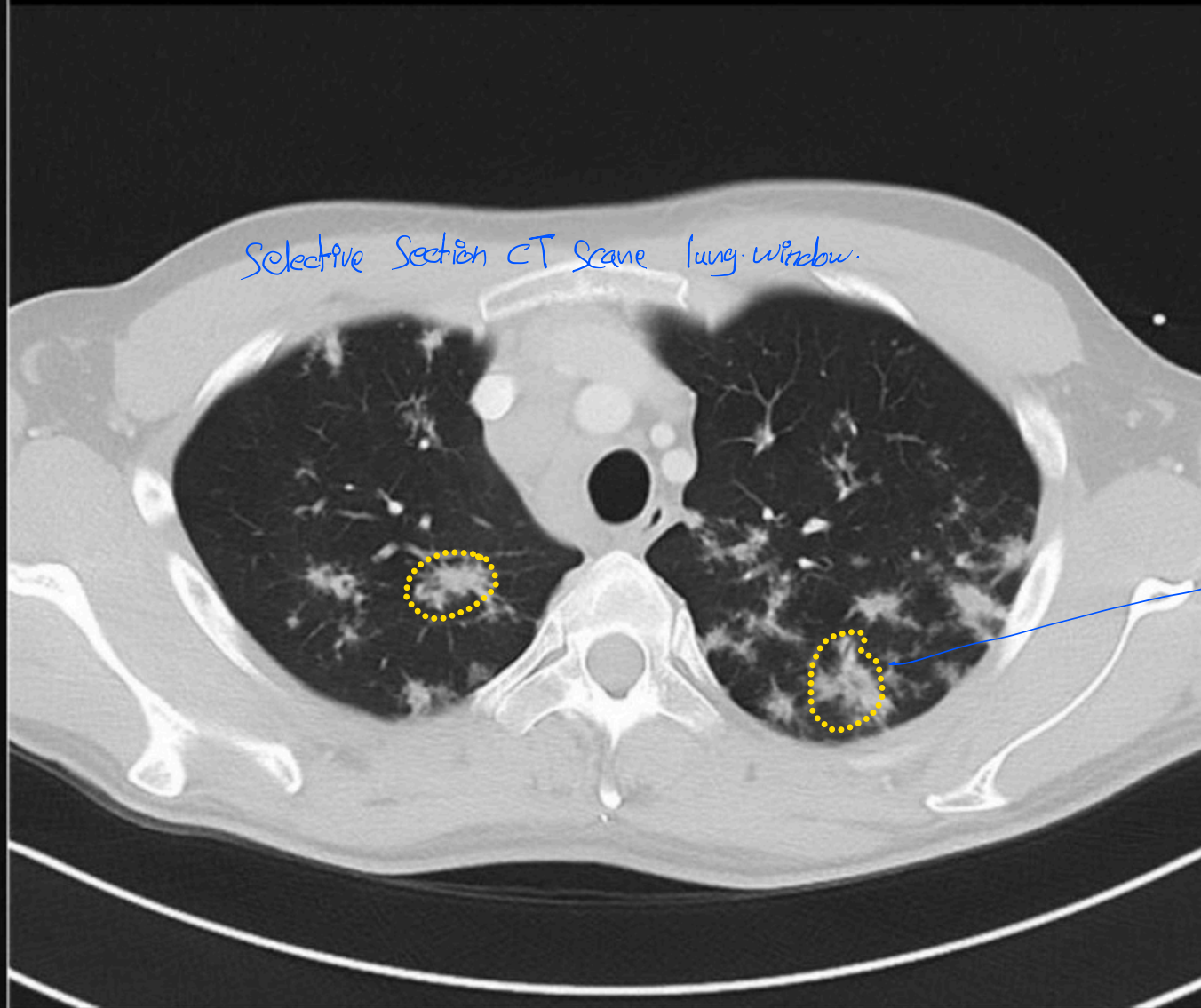
*DDX of Sarcoidosis:-

- Infection
- TB
- Malignancy

▷ Bilateral symmetrical lymphadenopathy with eggshell calcification.

- random opacity along the lung fields

- the picture suggestive of Sarcoidosis

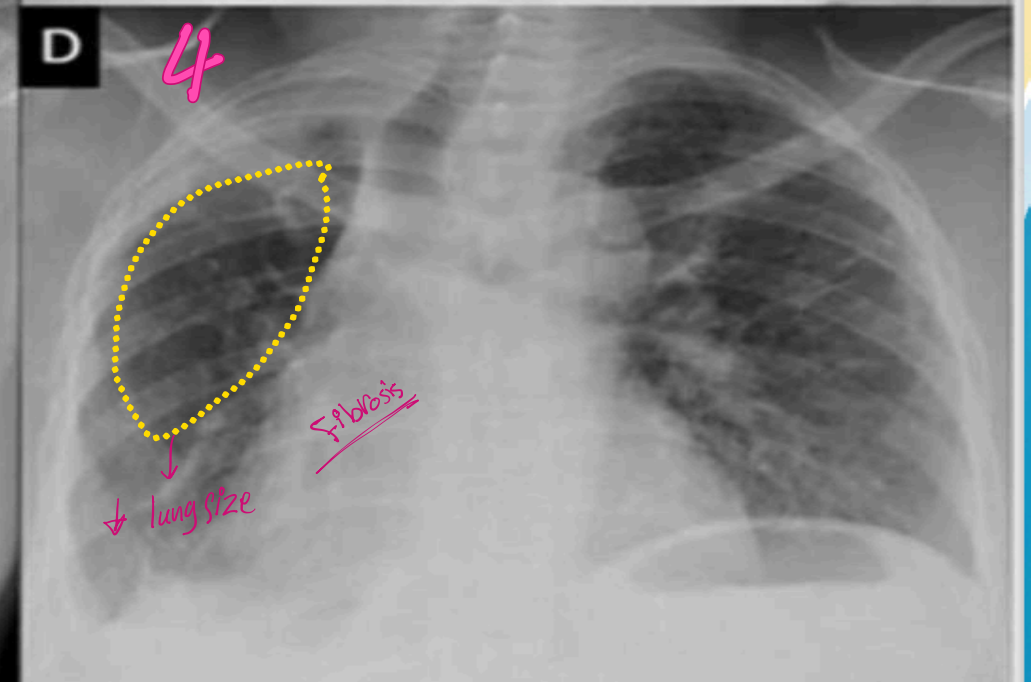
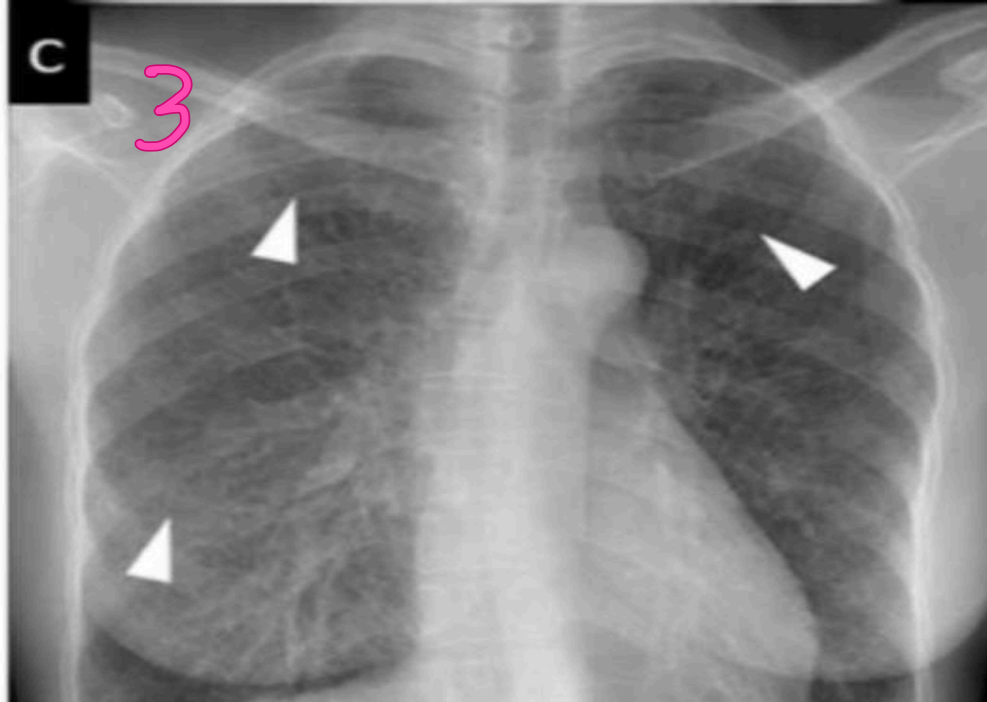
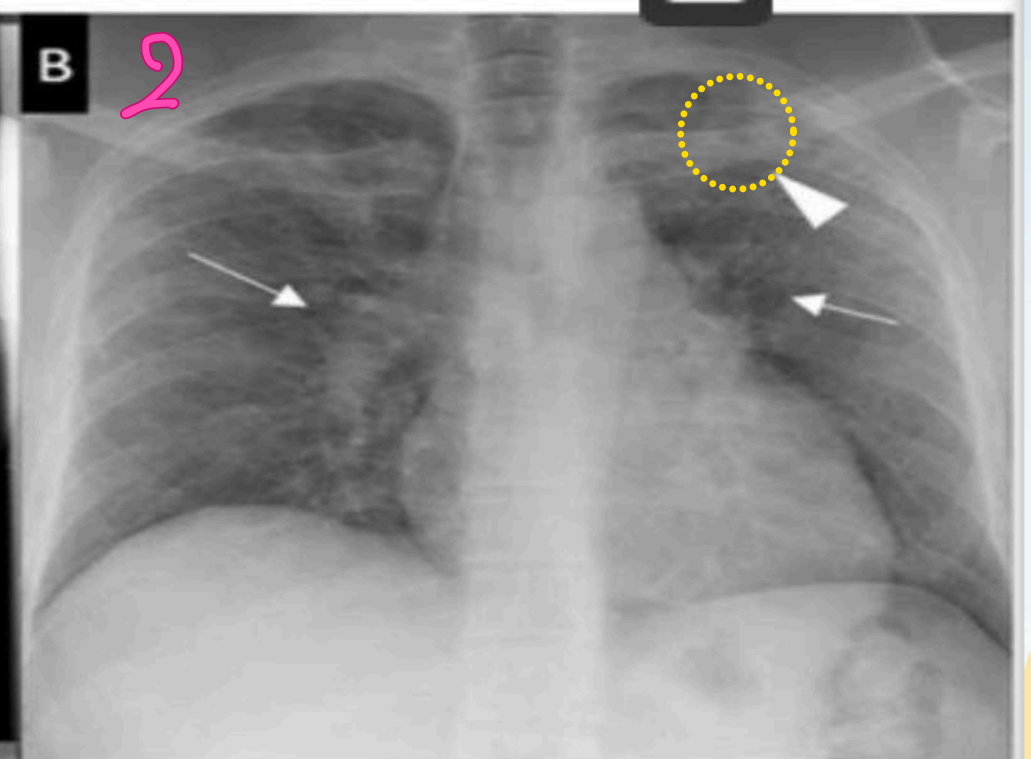
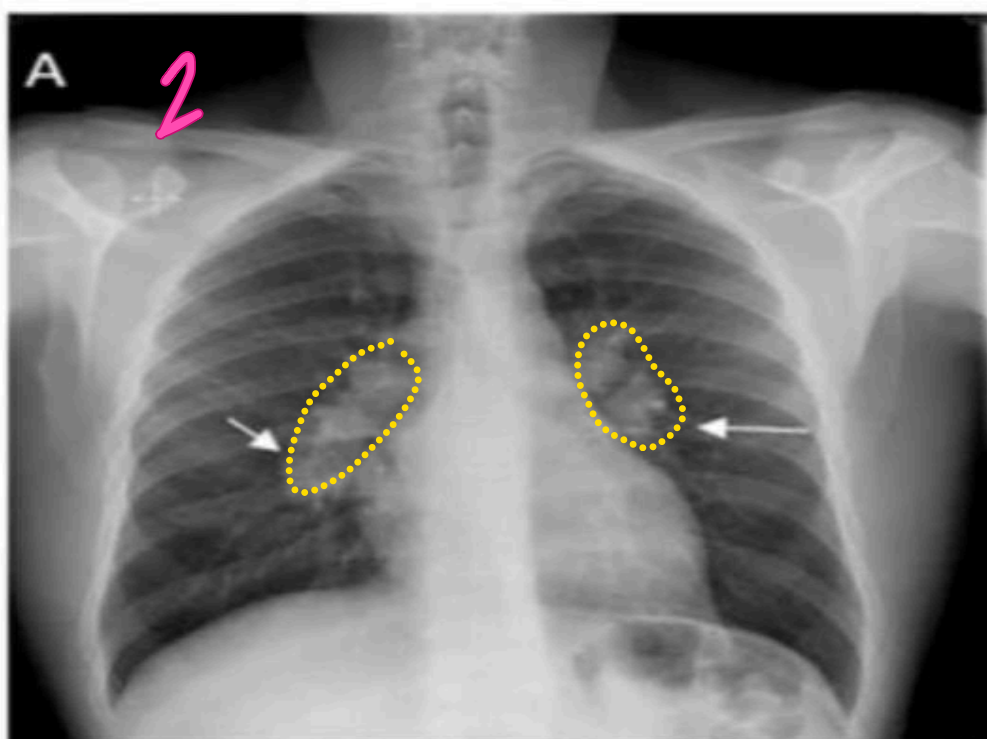


- Historical staging system has been used for radiographic findings (not CT)

Imp

Table 1 - Radiologic stages of thoracic changes of sarcoidosis.

Stage 0	Normal chest radiograph.
Stage 1	[REDACTED]
Stage 2	Lymphadenopathy with [REDACTED]
Stage 3	[REDACTED]
Stage 4	[REDACTED]



Garland triad

- also known as the 1-2-3 sign or pawnbroker's sign, is a lymph node enlargement pattern on chest radiographs which has been described in sarcoidosis:

1. Right paratracheal nodes

2. right hilar nodes

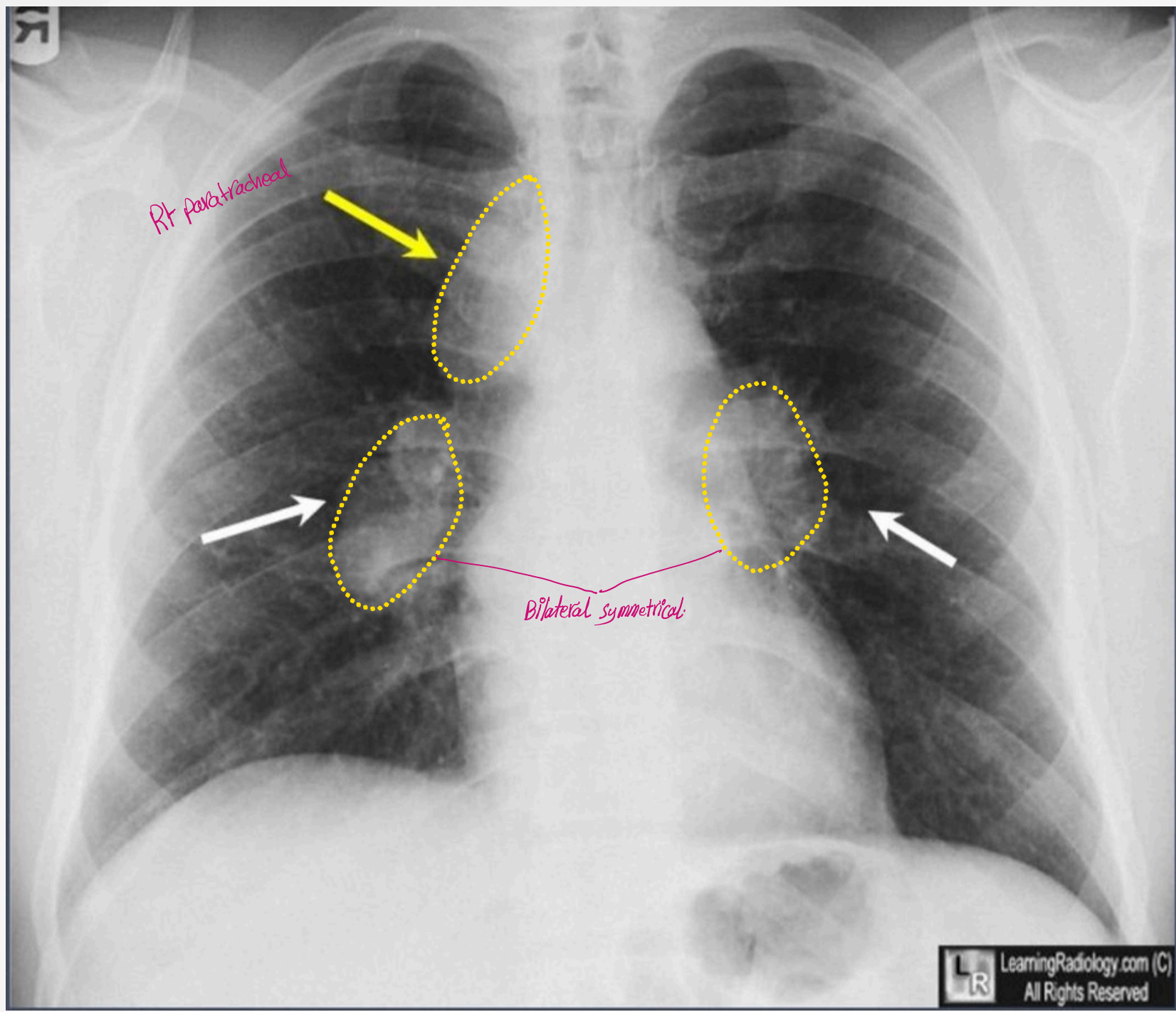
3. left hilar nodes

Bilateral symmetrical

- Hilar lymphadenopathy is **symmetrical** and usually massive.

These so-called potato nodes typically do not abut the cardiac border which distinguishes the nodal enlargement from lymphoma

* Sarcoidosis Stage 2



MEDIASTINUM

The mediastinum is a space in the thorax that contains a group of organs, vessels, nerves, lymphatics and their surrounding connective tissue.

It lies in the midline of the chest between the pleura of each lung and extends from the sternum to the vertebral column.

The mediastinum contains all the thoracic viscera except the lungs

Anterosuperior
mediastinum

4T

Middle
mediastinum

* Cardiac + Bronchogenic cyst

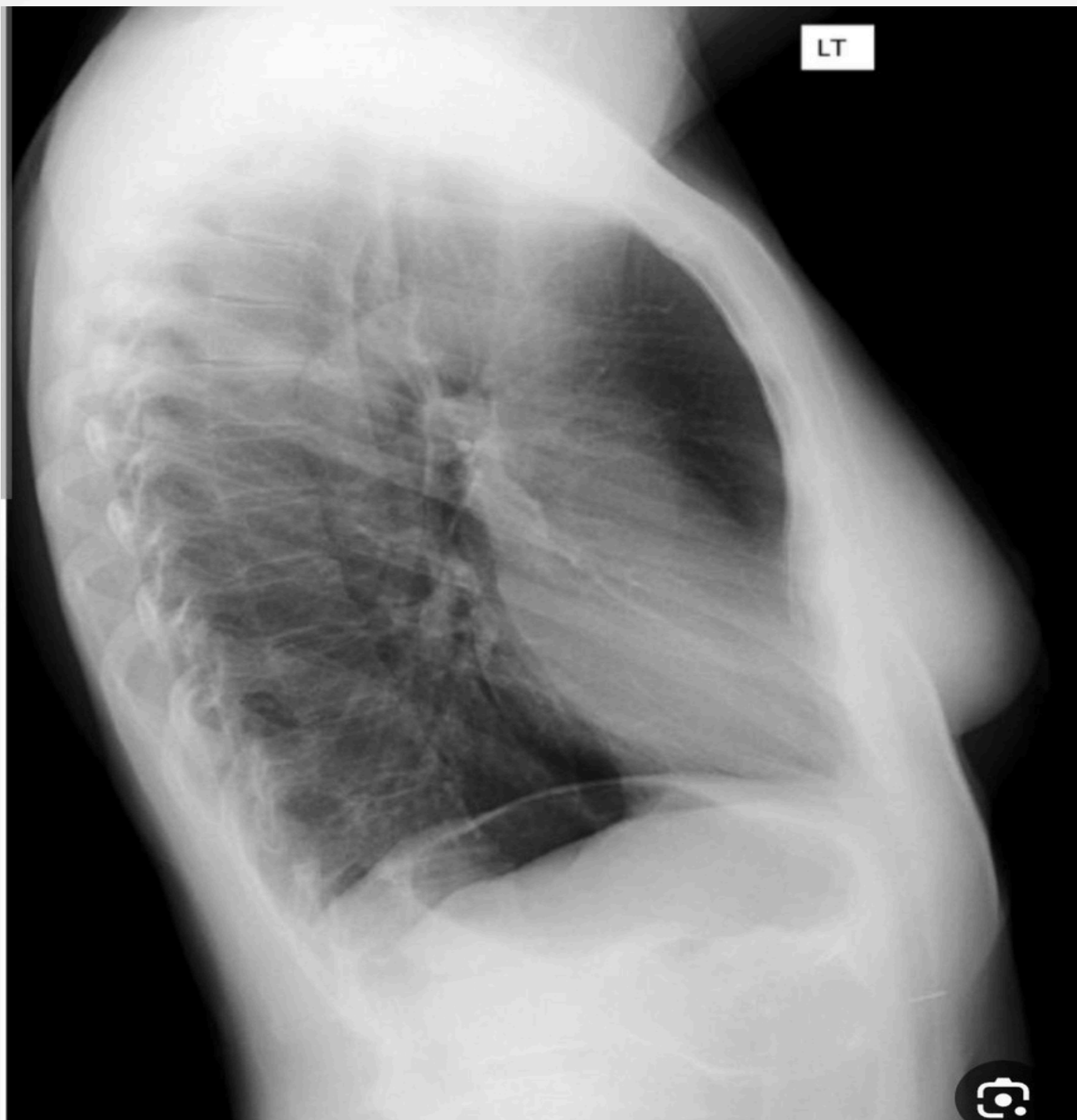
Descending Aorta

Posterior
mediastinum



Anatomic division

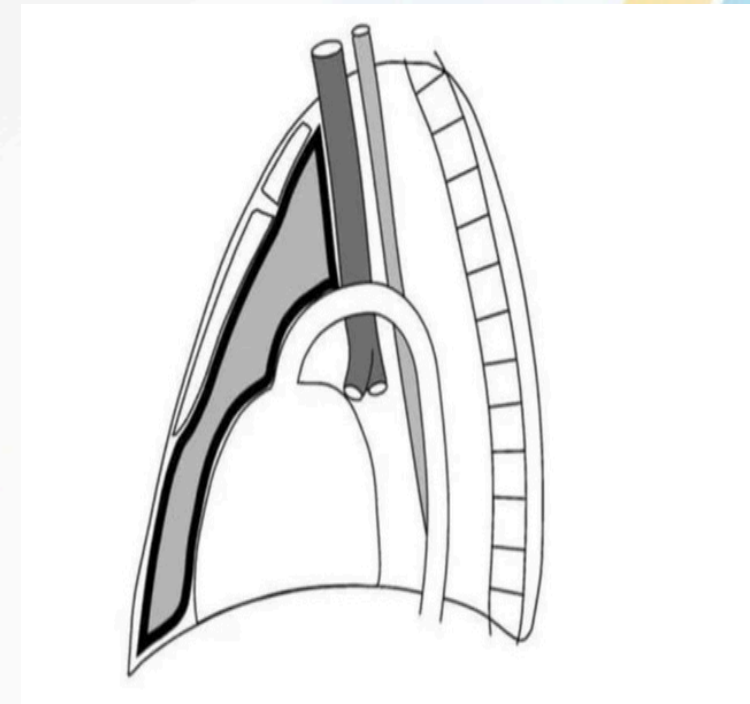
- The mediastinum is divided into three arbitrary compartments to aid in the differential diagnosis of a mediastinal mass. However, there are no anatomic planes separating these divisions and disease can spread from one “compartment” to another
1. anterior mediastinum: anterior to the pericardium
 2. middle mediastinum: within the pericardium
 3. posterior mediastinum : posterior to pericardium



Anterior mediastinum 4T

Hyroid
Thymus
T-Cell lymphoma
Teratoma

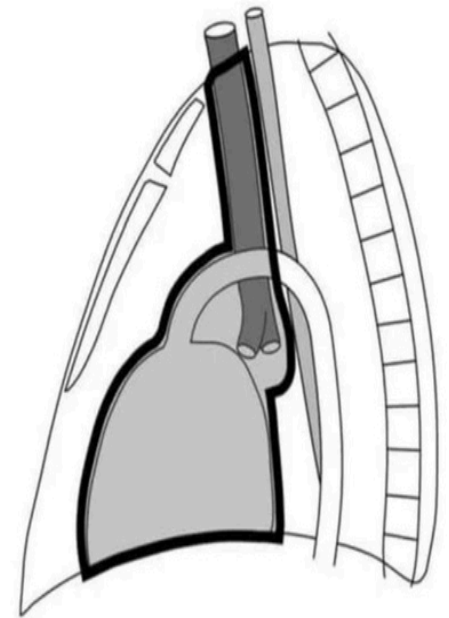
- The anterior mediastinum is the space between the sternum and the pericardium inferiorly and ascending aorta and brachiocephalic vessels superiorly.
- The anterior mediastinum can be thought of as two compartments – the **prevascular** compartment superiorly and the **precardiac** compartment inferiorly.
- The contents of the **prevascular** anterior mediastinum include:
 - Thymus. → *Thymoma; Teratoma*
 - Lymph nodes.
 - Enlarged thyroid gland, if it extends inferiorly into the mediastinum.
- The **precardiac anterior mediastinum** is a potential space.



Middle mediastinum

- The anterior border of the middle mediastinum is the anterior pericardium and the posterior borders are the posterior pericardium and posterior tracheal wall.
- The contents of the middle mediastinum include:

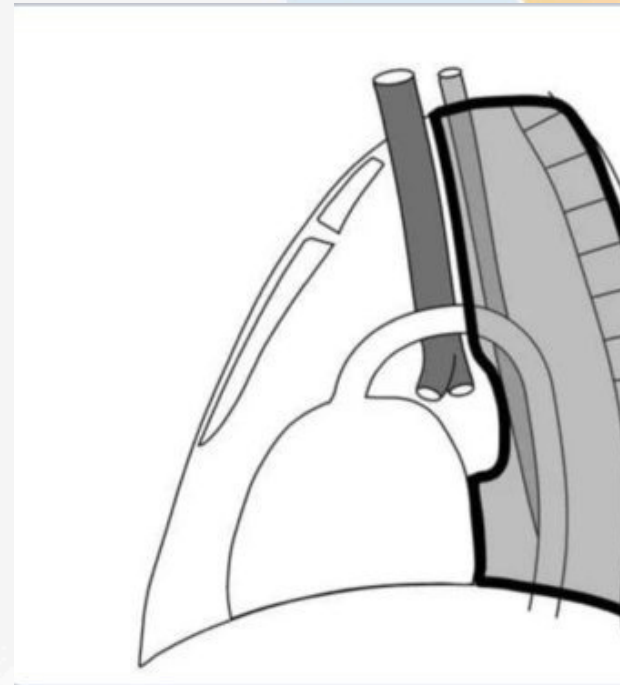
1. Ascending aorta and aortic arch.
2. Great vessels including SVC, IVC, pulmonary arteries and veins, and brachiocephalic vessels
3. Trachea and bronchi
4. Lymph nodes.
5. Phrenic, vagus, and recurrent laryngeal nerves (all of which pass through the AP window).



Posterior mediastinum

The anterior border of the posterior mediastinum is the posterior trachea and posterior pericardium.

- The posterior border is somewhat loosely defined as the anterior aspect of the vertebral bodies; however, paraspinal masses are generally included in the differential of a posterior mediastinal mass
- The contents of the posterior mediastinum include:



Esophagus.

Descending thoracic aorta.

Azygos and hemiazygos veins.

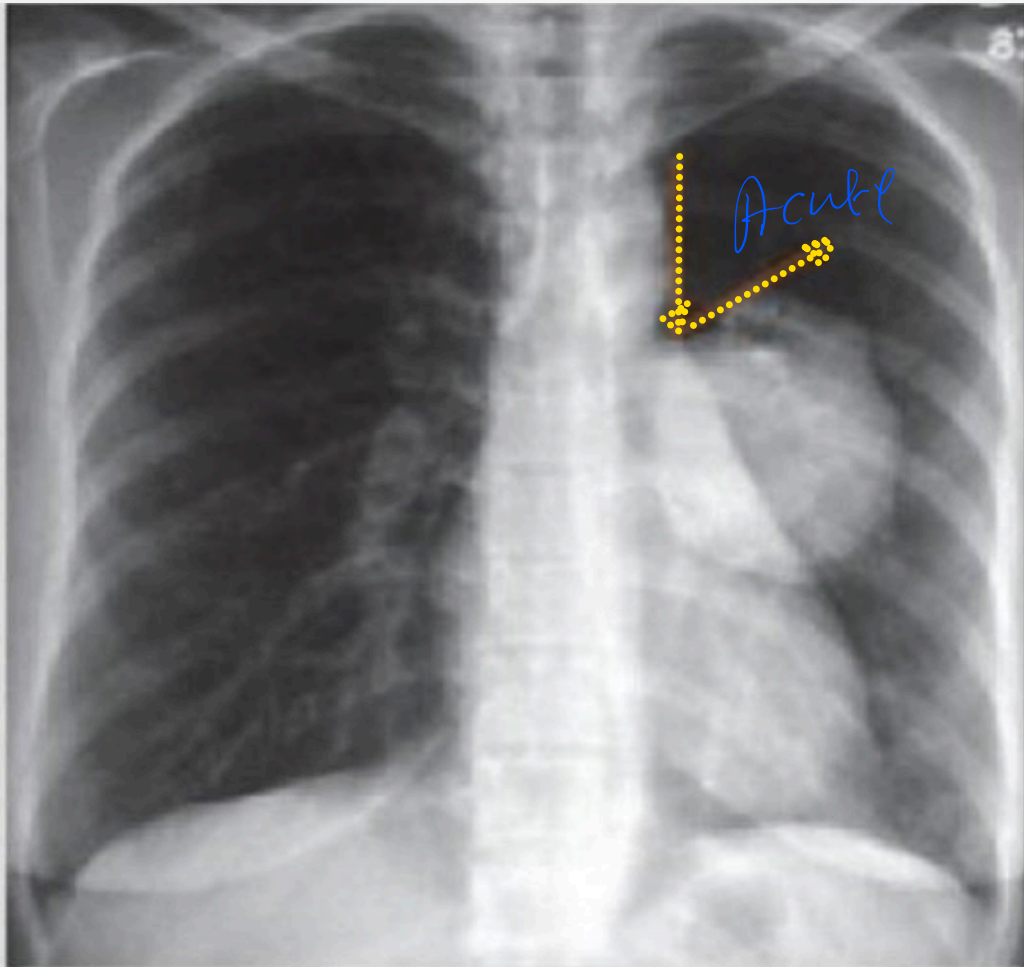
Thoracic duct.

Vagus nerves.

Lymph nodes.

The following characteristics indicate that a lesion originates within the mediastinum

1. Unlike lung lesions, a mediastinal mass will not contain air bronchograms.
2. The margins with the lung will be obtuse.
3. There can be associated spinal, costal or sternal abnormalities.



Pulmonary mass

- [redacted] with mediastinum
- [redacted] silhouetting of mediastinum

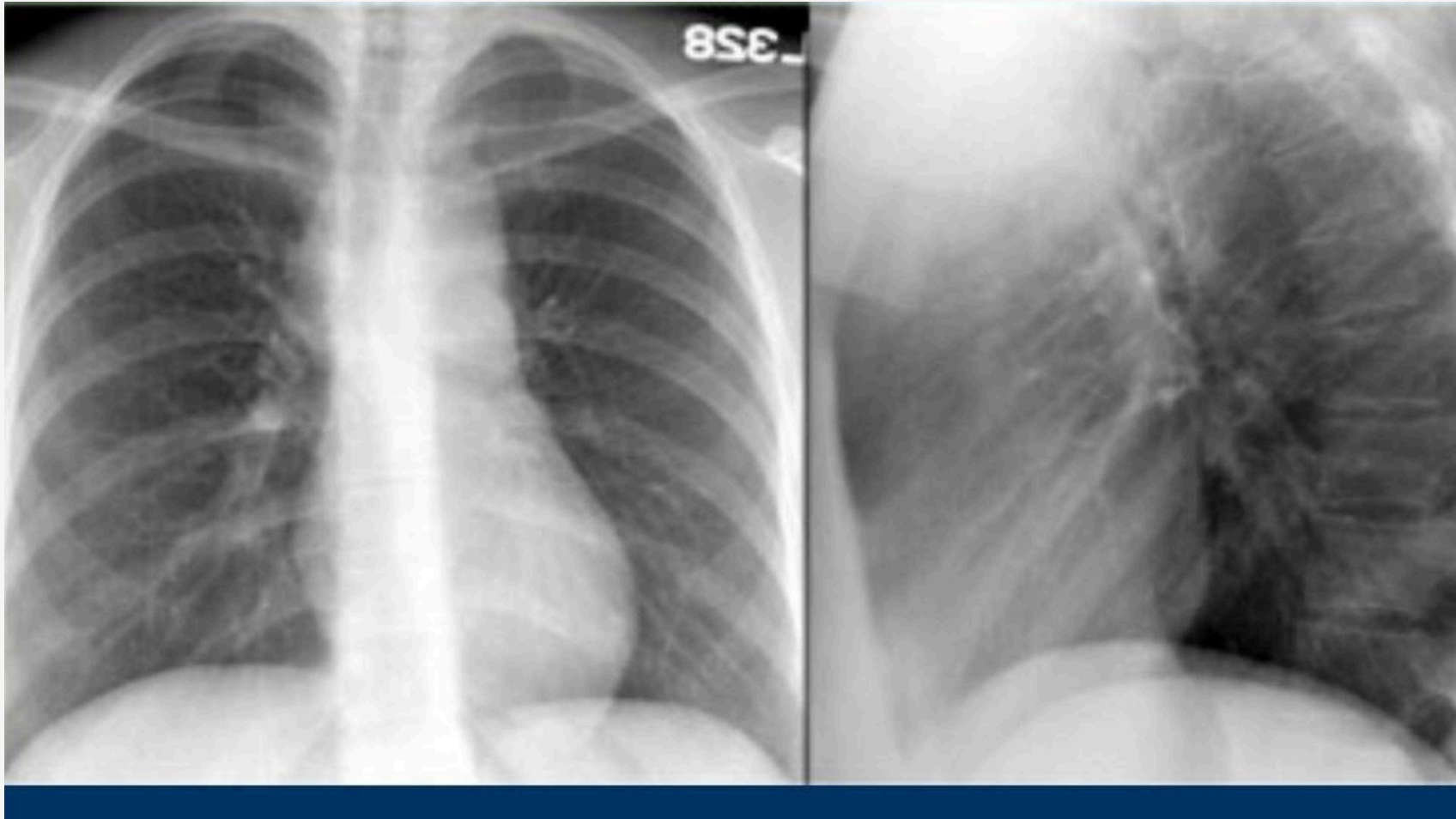


Mediastinal mass

- [redacted] with mediastinum
- Silhouetting of mediastinum

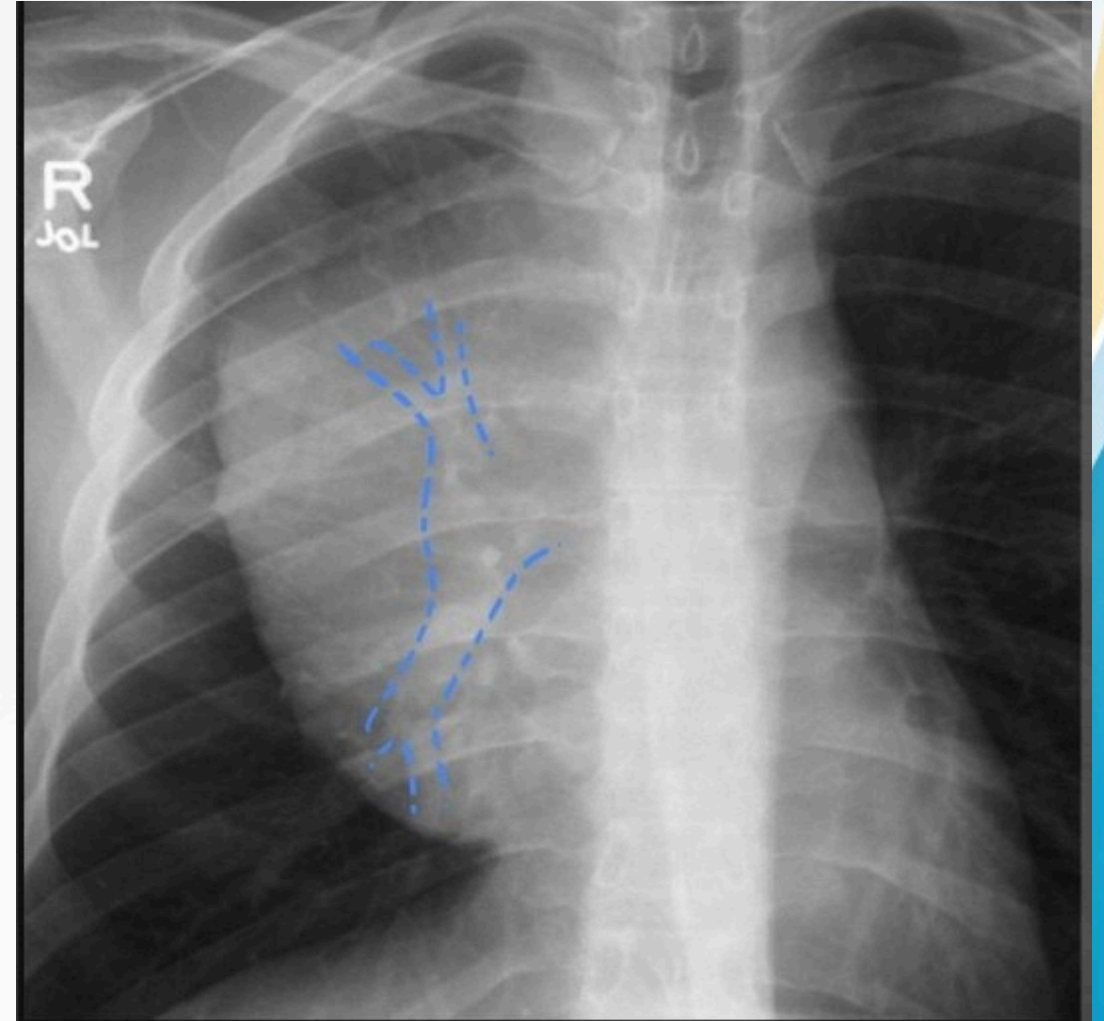
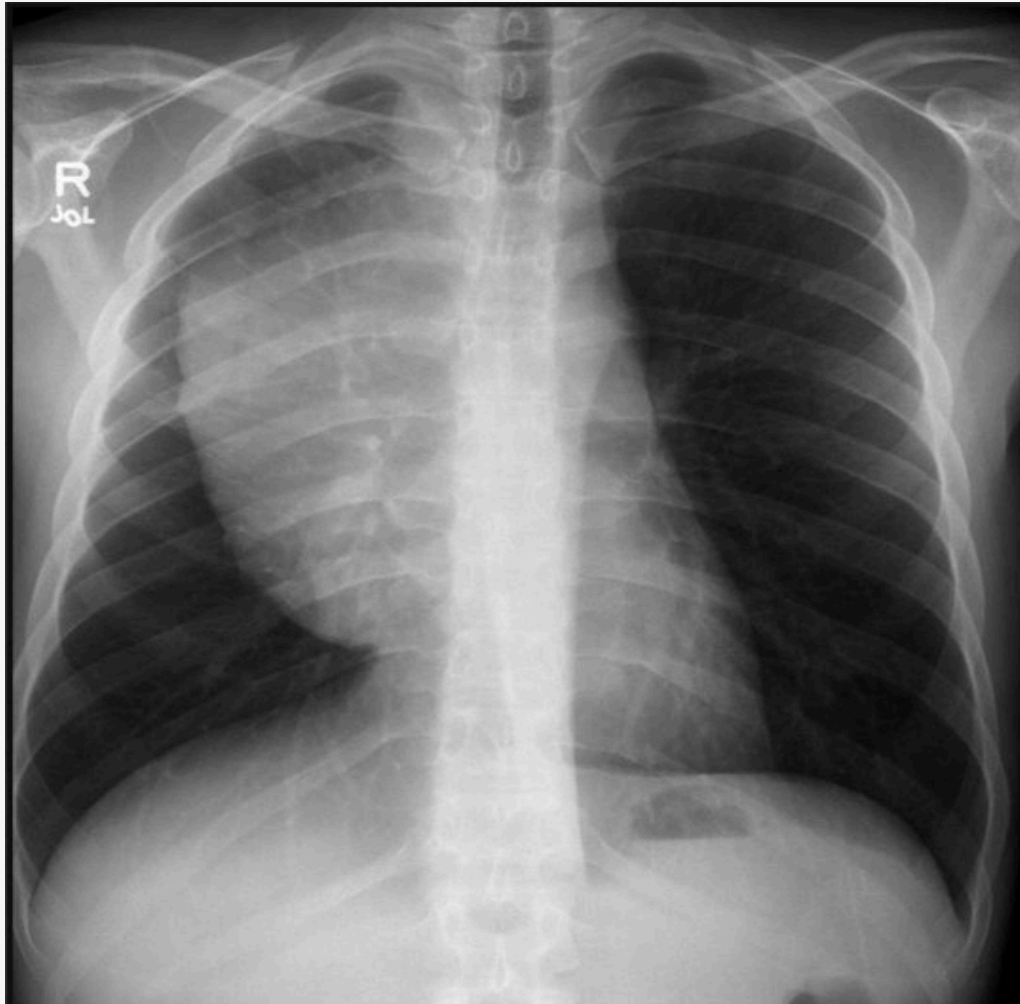
Mass in anterior mediastinum

1. Obliterated retrosternal clear space

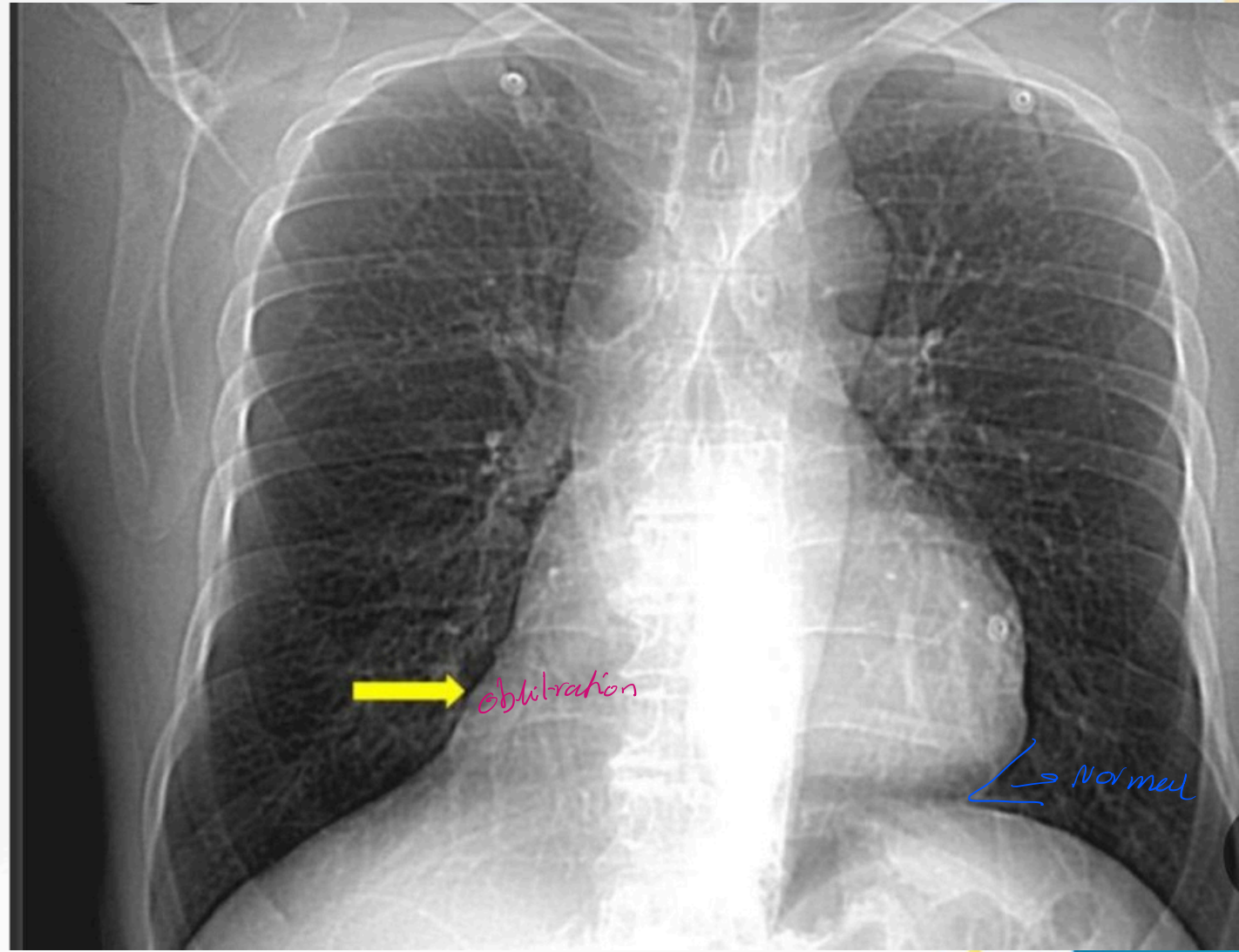


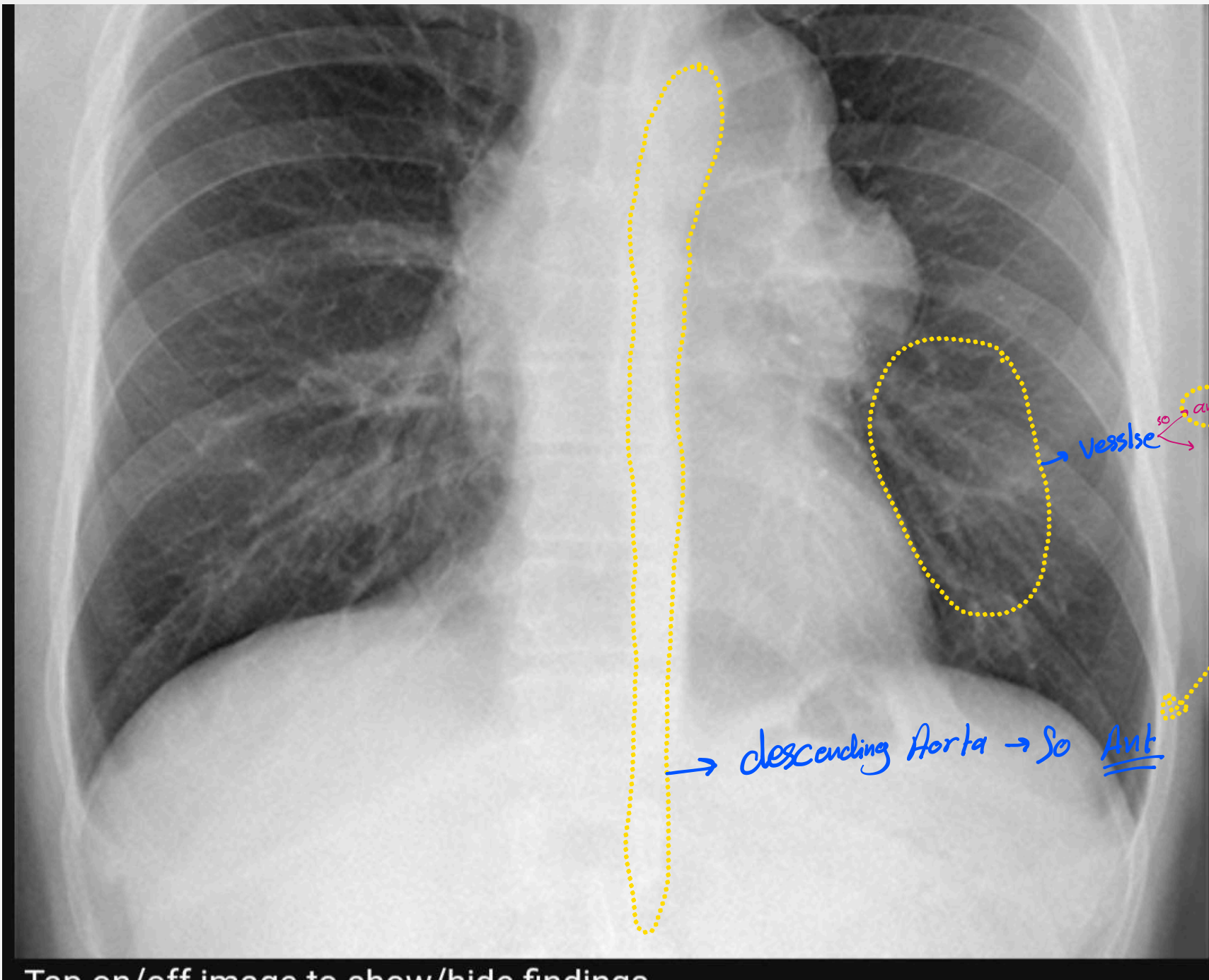
2. The hilum overlay sign :

is present on the frontal view if hilar vessels are visualized through the mass. It indicates that the mass cannot be in the middle mediastinum. The mass may be in the anterior (most likely) or posterior mediastinum.

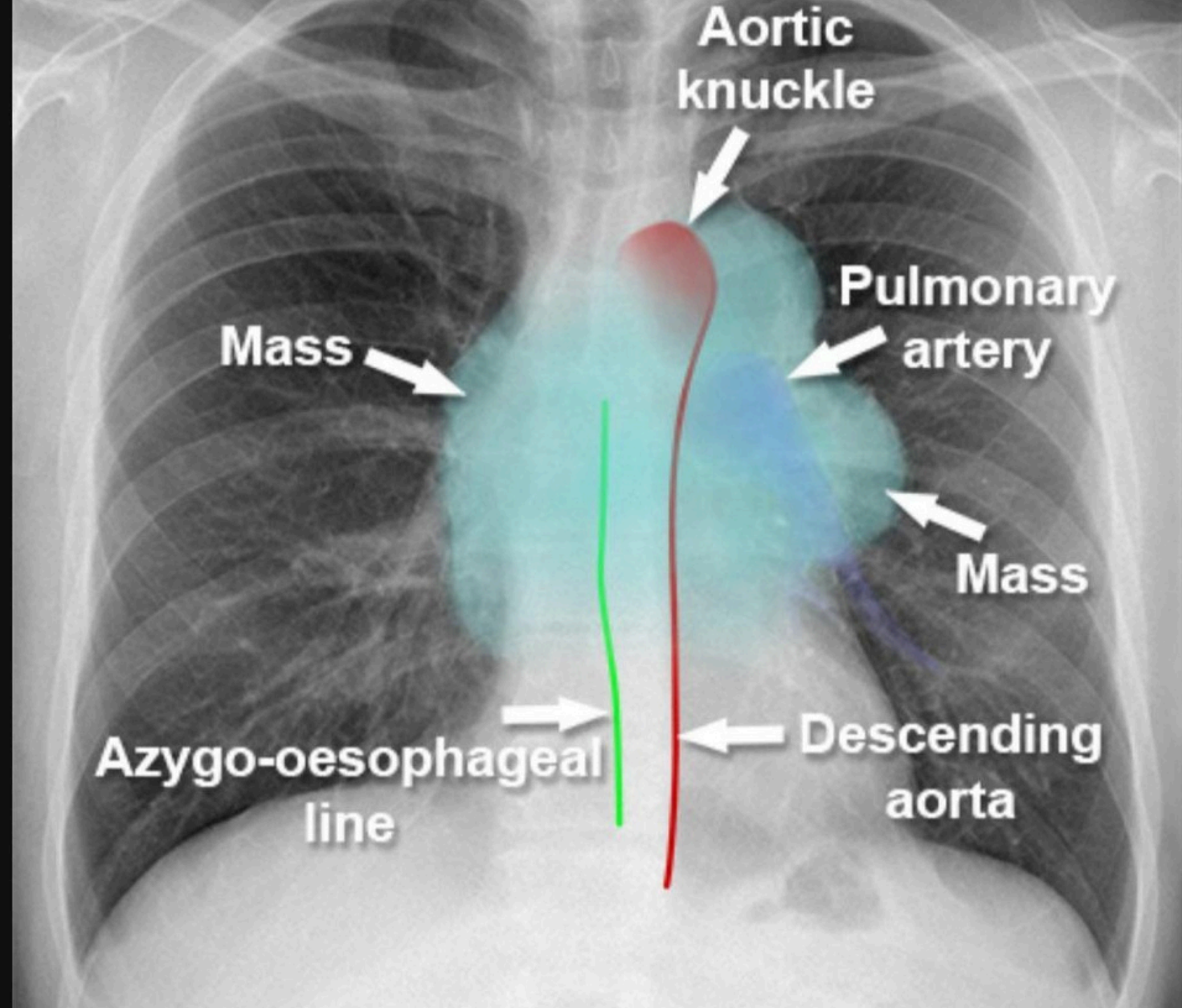


3. Obliteration of cardiophrenic angle





Tap on/off image to show/hide findings



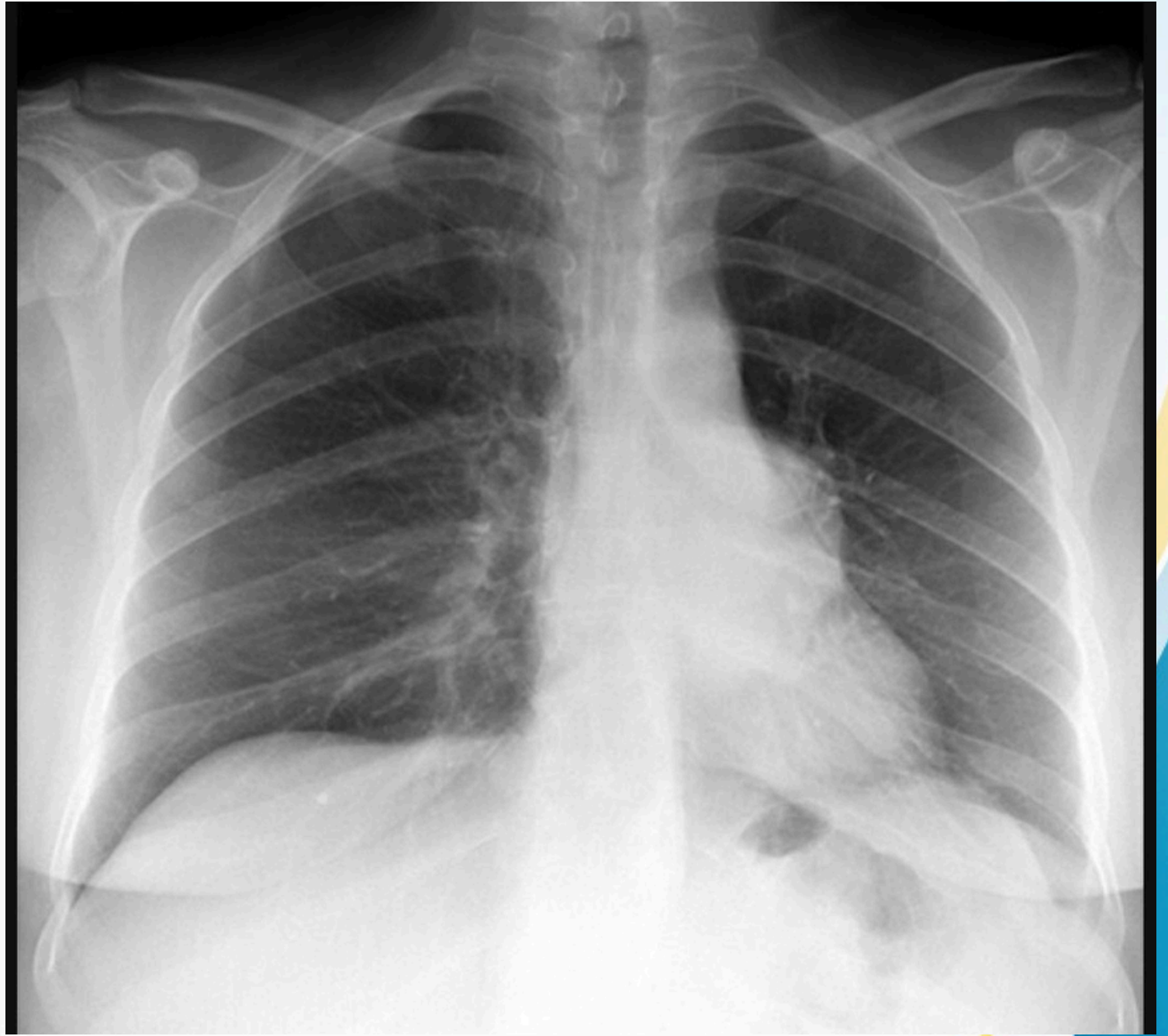
Posterior mediastinal mass

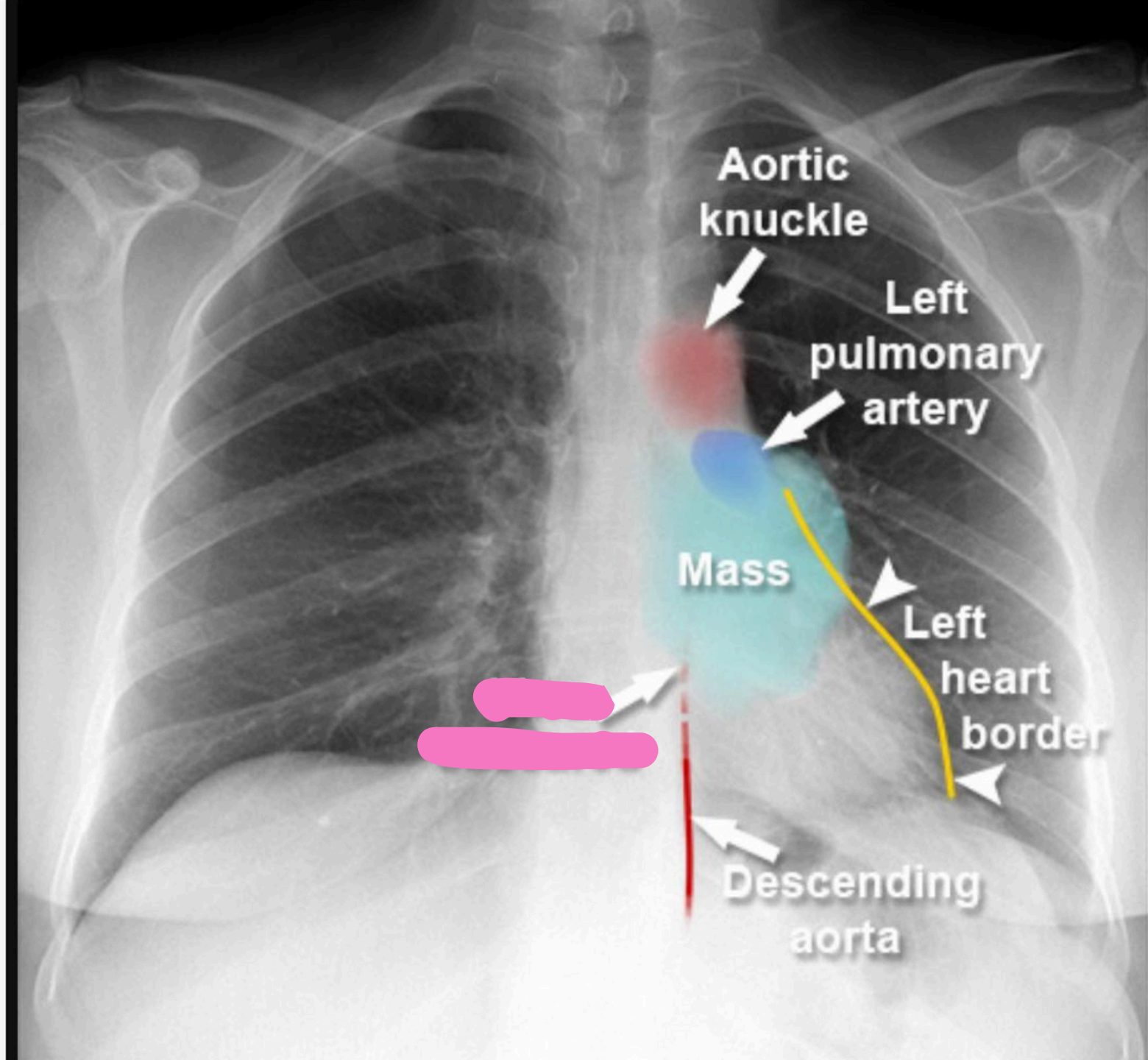
1. Hilum overlay sign
2. Distinct heart border [not in middle mediastinum]
3. Seen on lateral view posterior to heart
4. On left side obscuring the descending aorta
5. Adjacent to vertebral body on lateral view
6. Cervico thoracic sign → mass above the clavicle

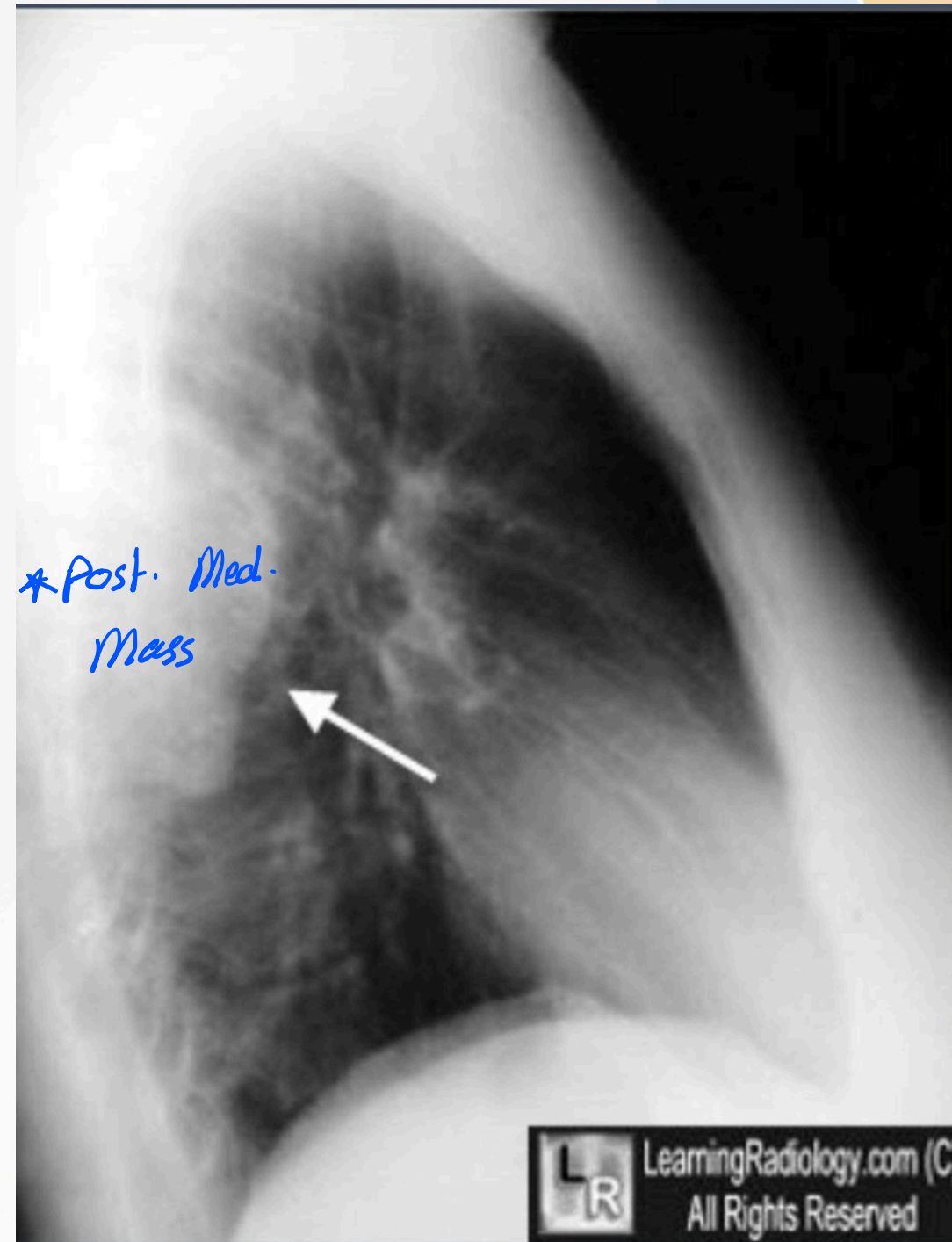
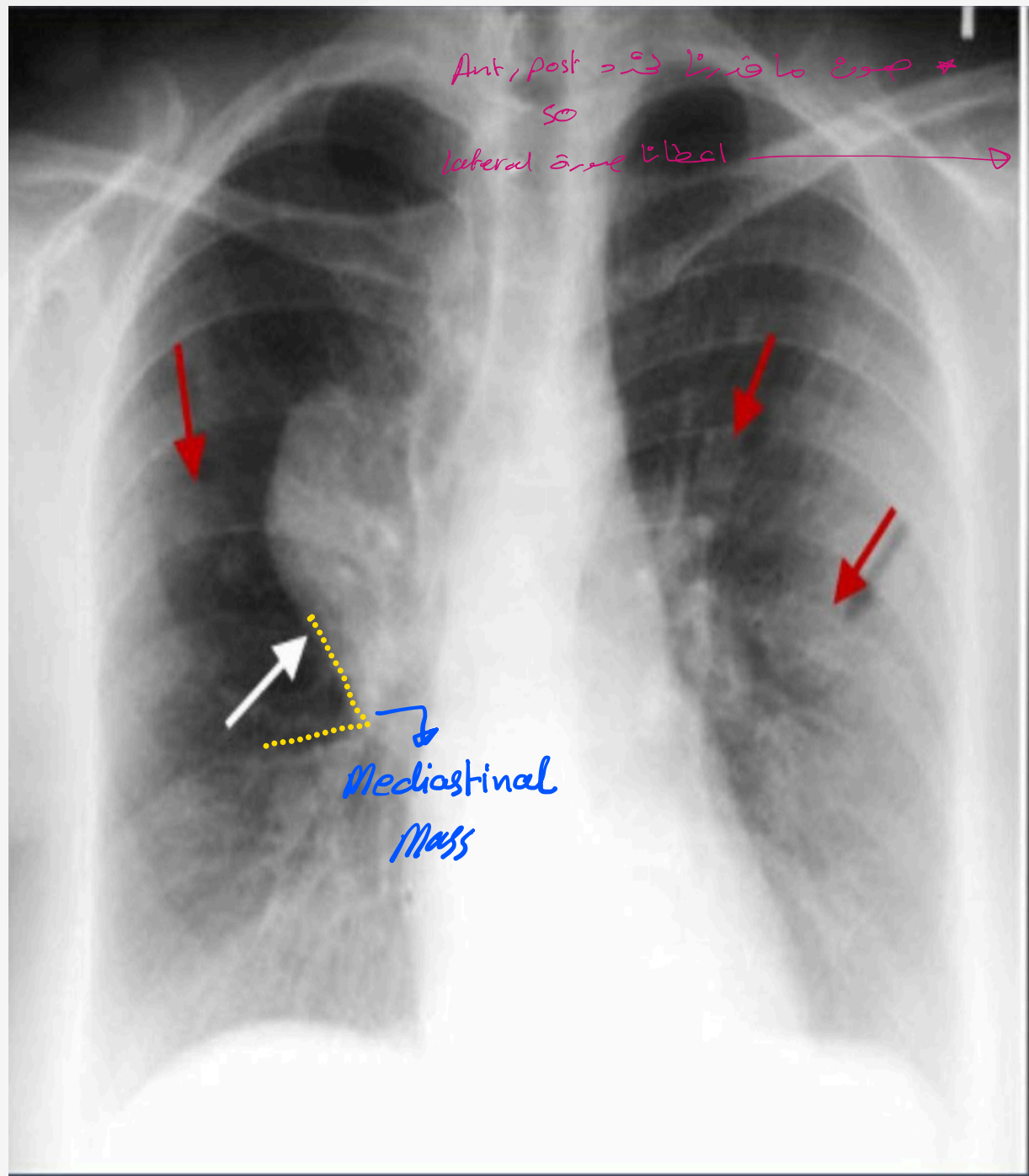
Posterior mediastinal mass

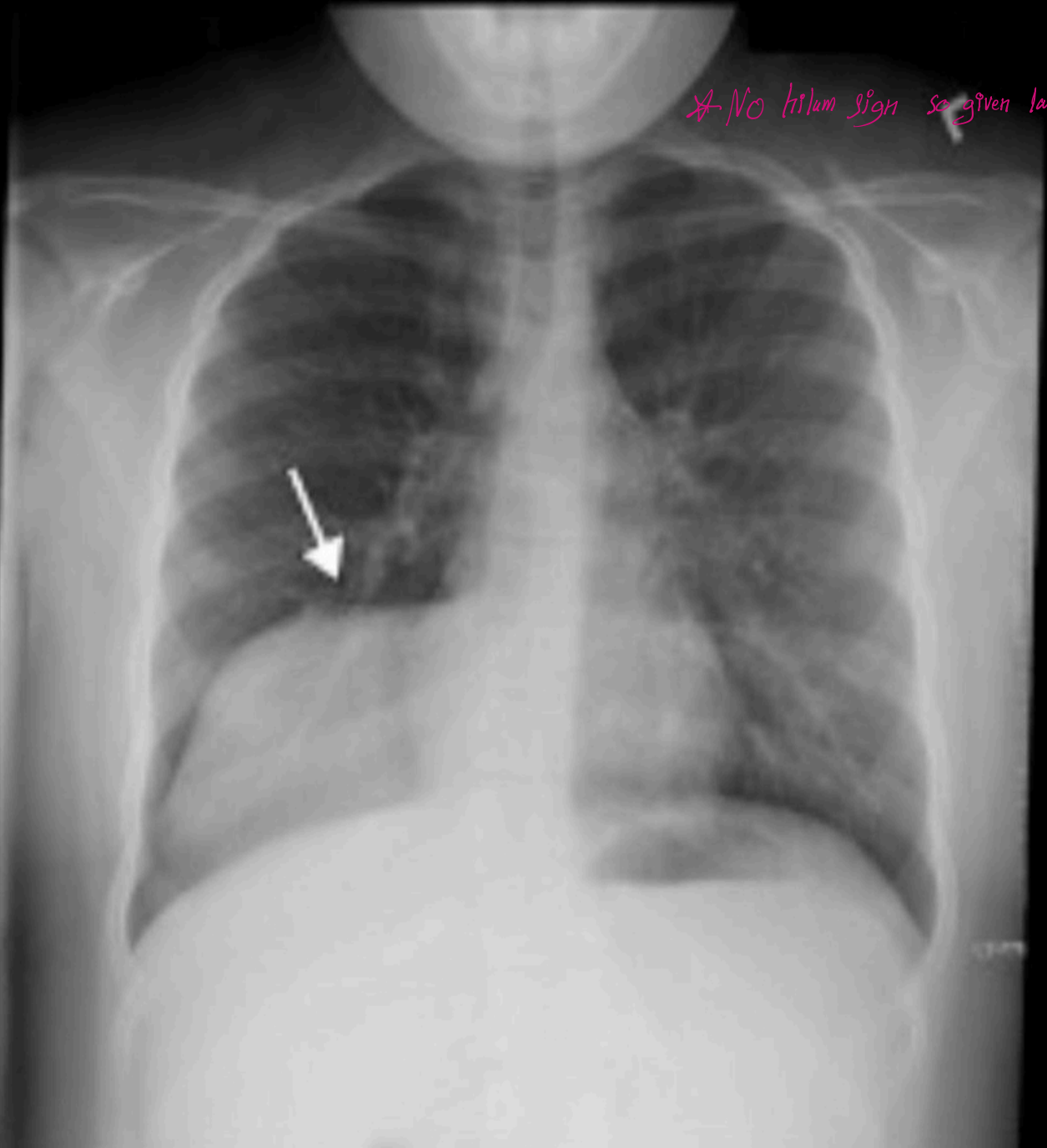
— PA
— post mediastinal mass

+
Normal

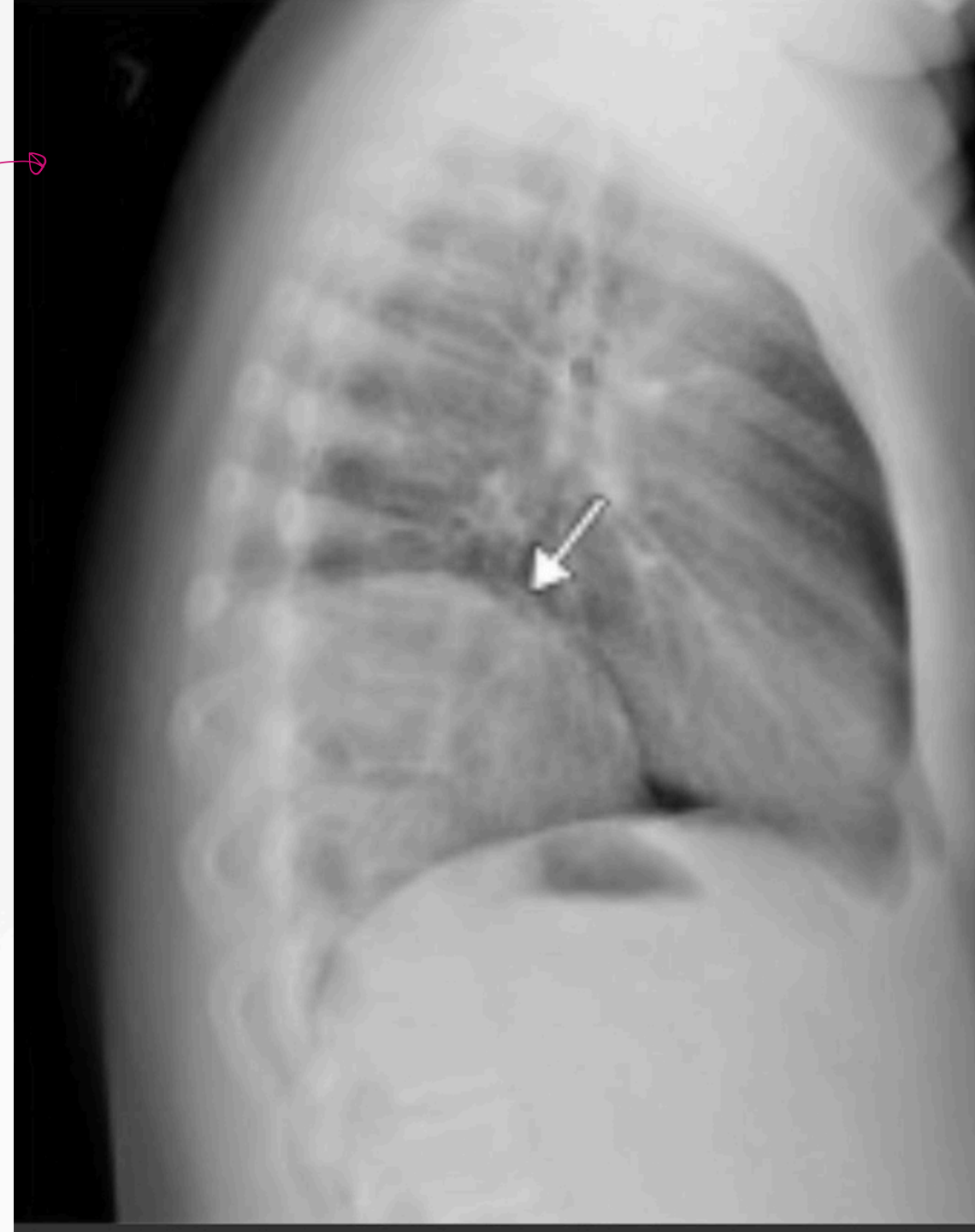








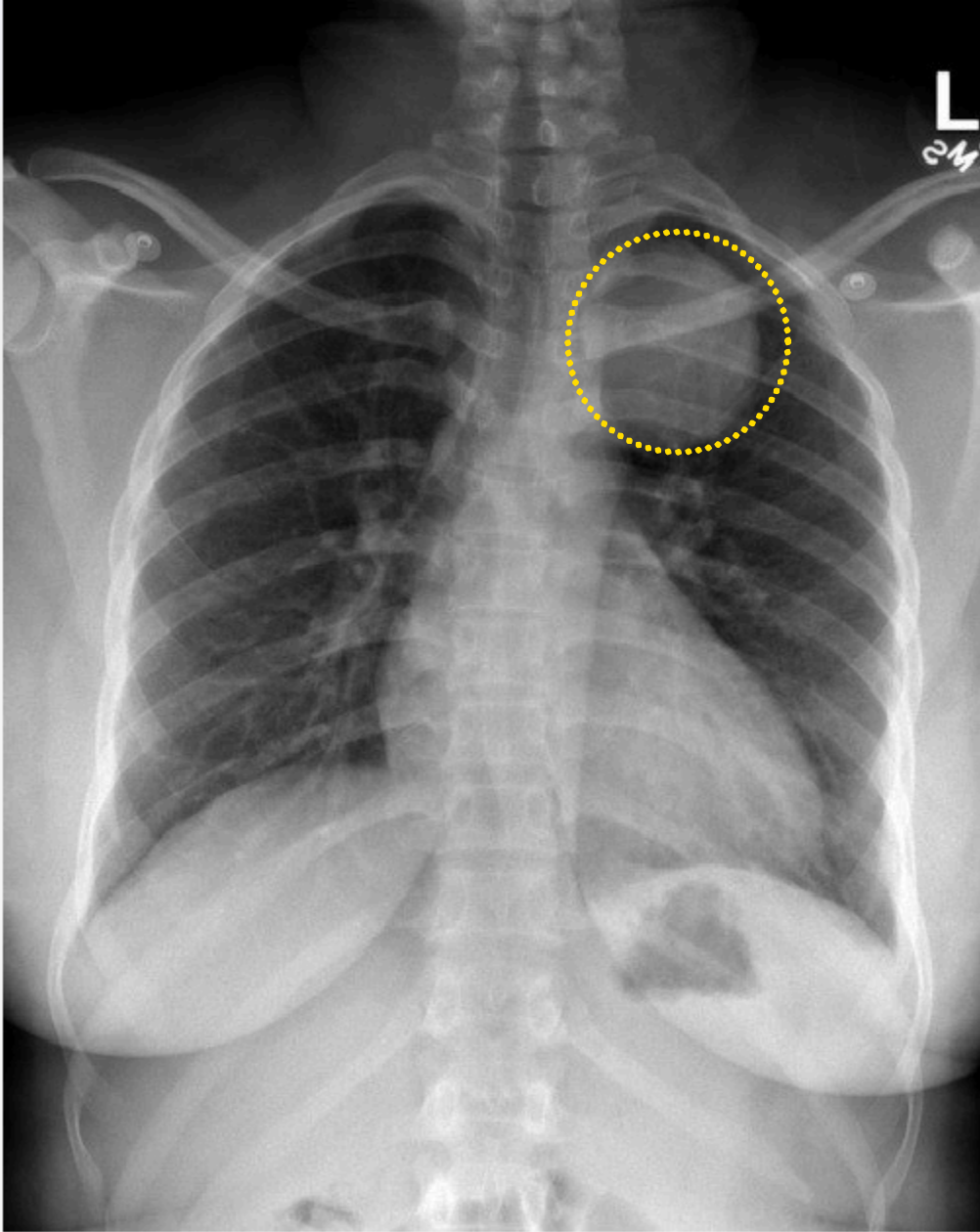
** No hilum sign so given lateral view*



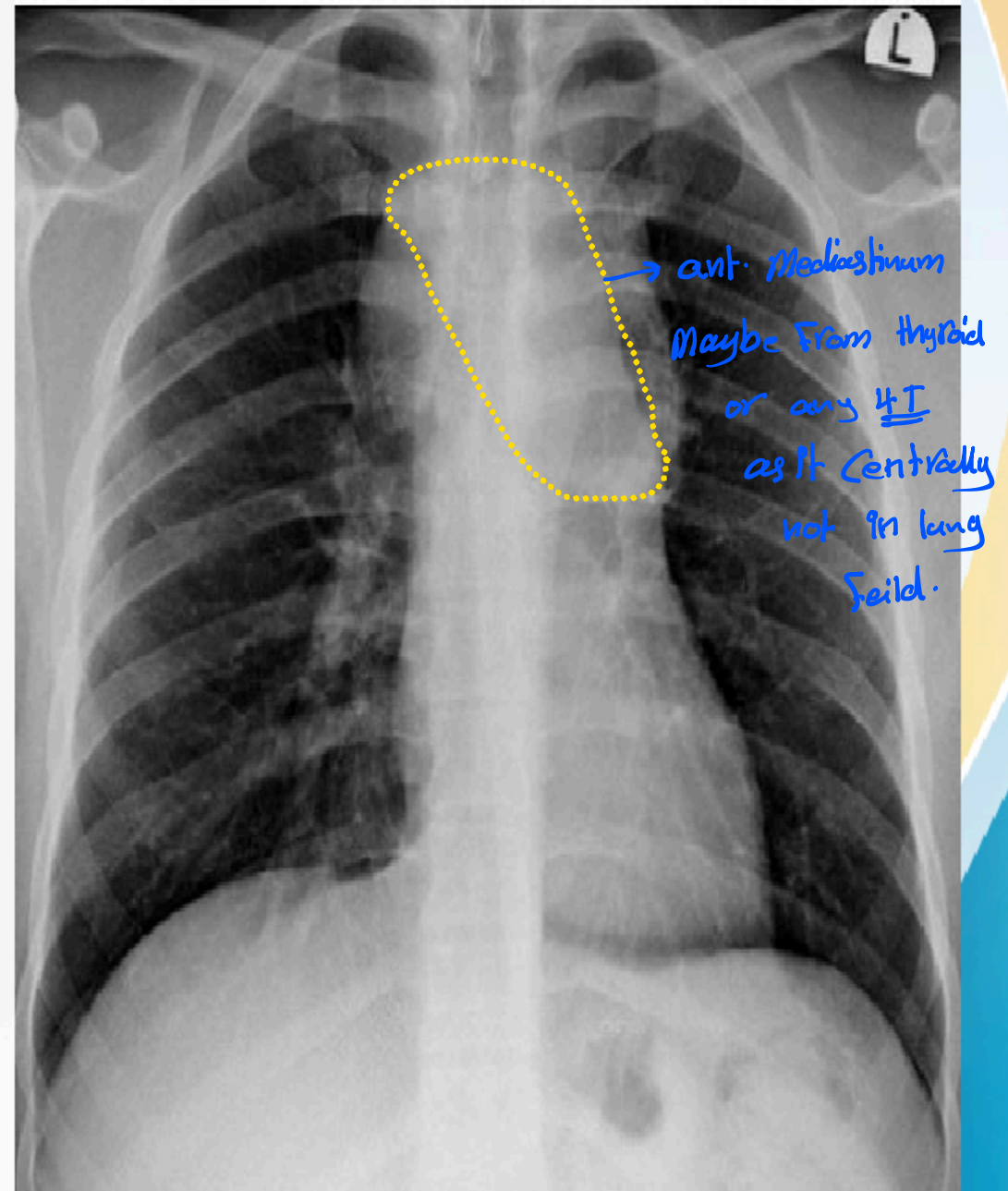
cervicothoracic sign

● **positive cervicothoracic sign** occurs when a thoracic lesion contacts the neck or extends into it, thereby obliterating the upper borders of the lesion on radiographs

- Because the anterior mediastinum ends at the level of the clavicles
- In contrast, a lesion posterior to the trachea can have its upper border outlined by lung above the level of the clavicles (**negative cervicothoracic sign**), as the posterior mediastinum ends much higher

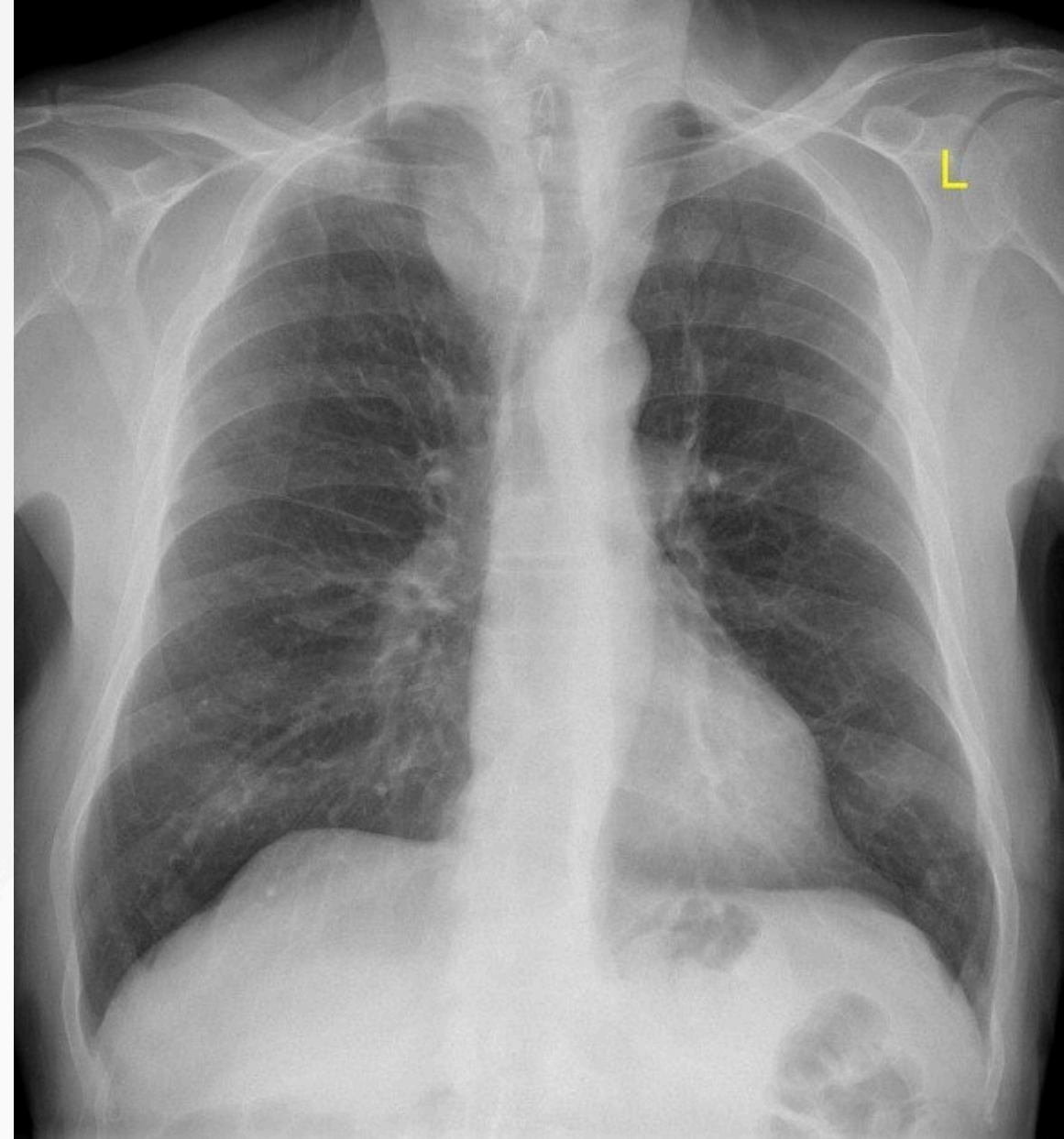


posterior mediastinal mass



anterior mediastinal mass

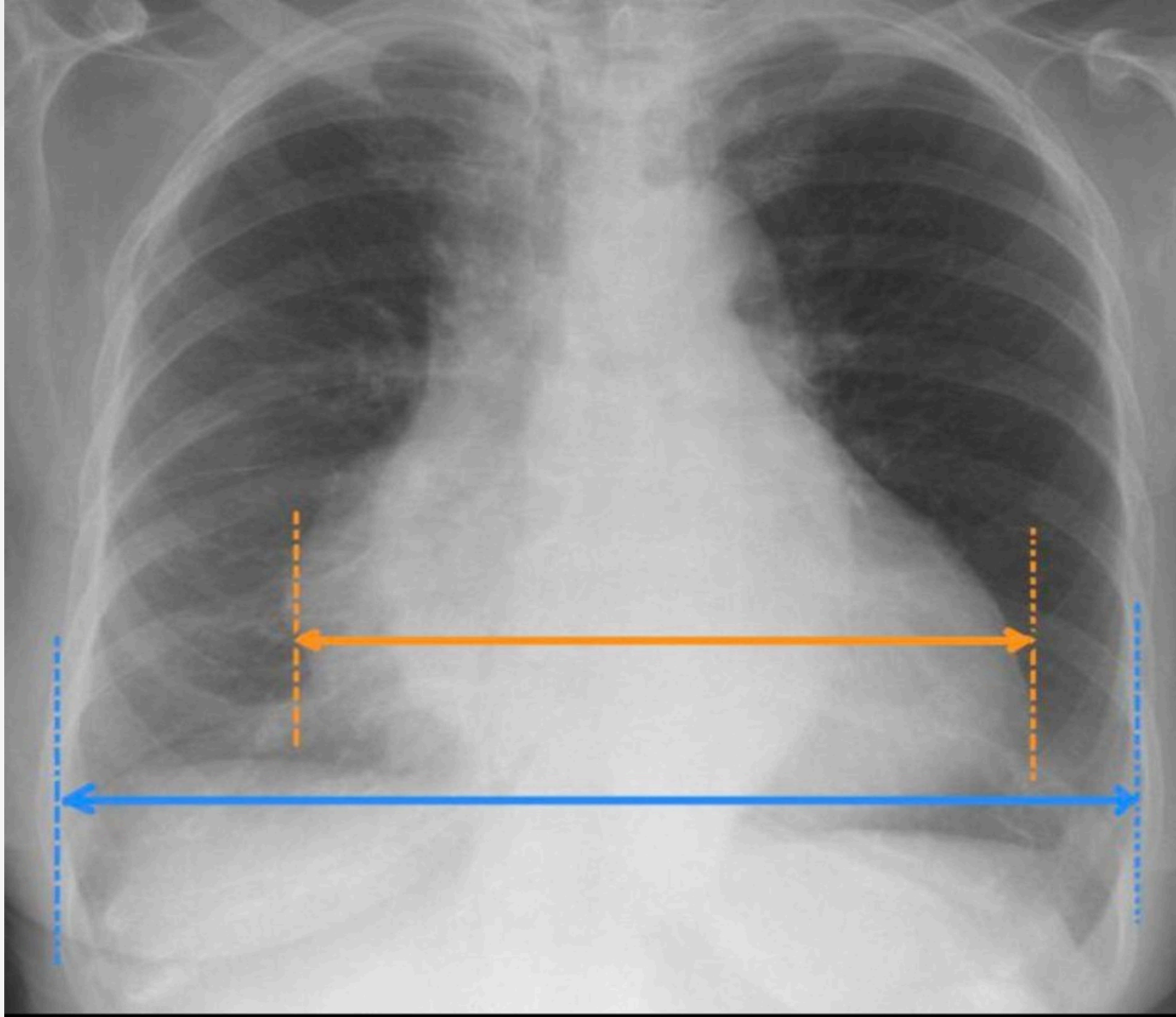
- if a thoracic mass is in direct contiguity with the soft tissues of the neck, the borders delineating their point of contact will be lost or obscured



Cardiomegaly

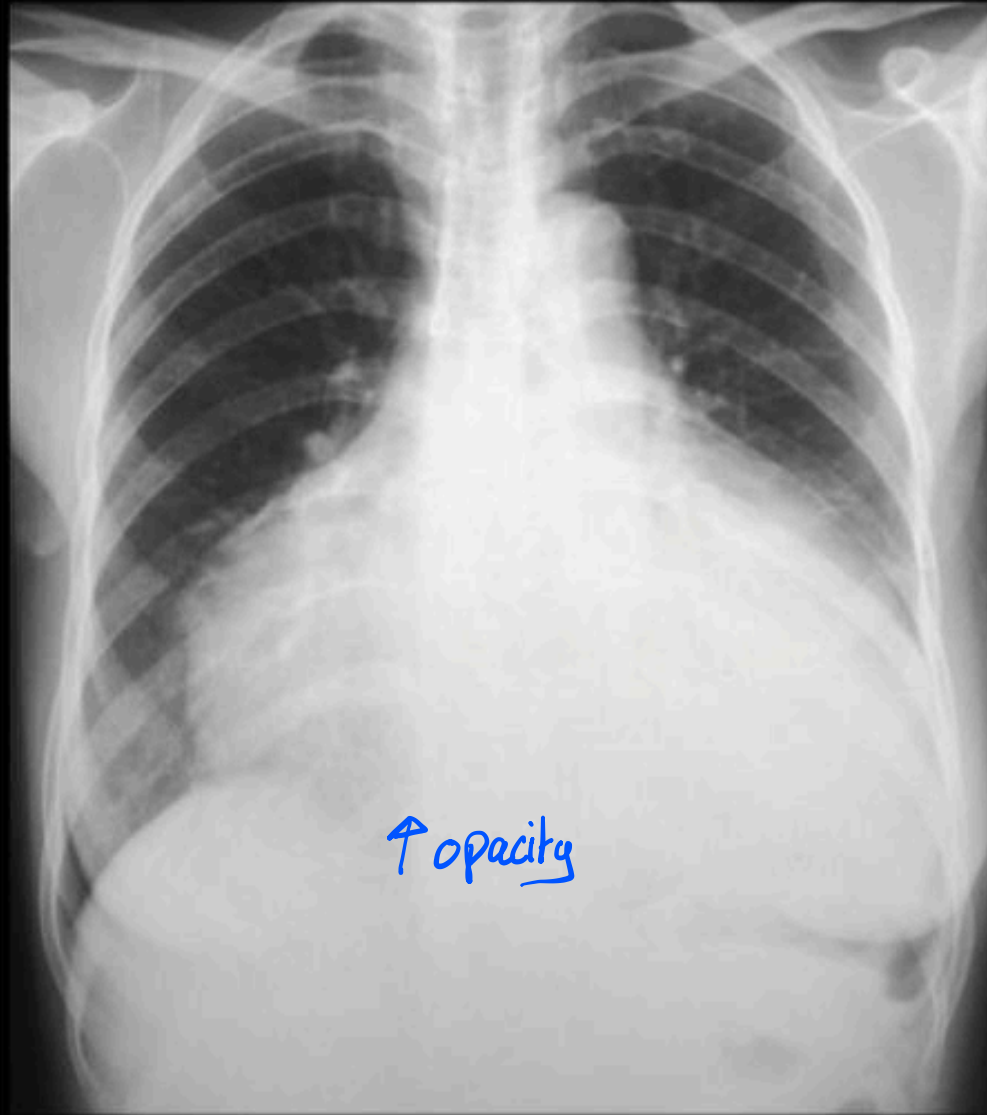
- means enlargement of the heart.
- The definition is when the transverse diameter of the cardiac silhouette is greater than or equal to 50% of the transverse diameter of the chest (increased cardiothoracic ratio) on a posterior-anterior projection of a chest radiograph or a computed tomography





Pericardial effusion

- Pericardial effusions occur when excess fluid collects in the pericardial space (a normal pericardial sac contains approximately 30-50 mL of fluid).
- There can be globular enlargement of the cardiac shadow giving a water bottle configuration



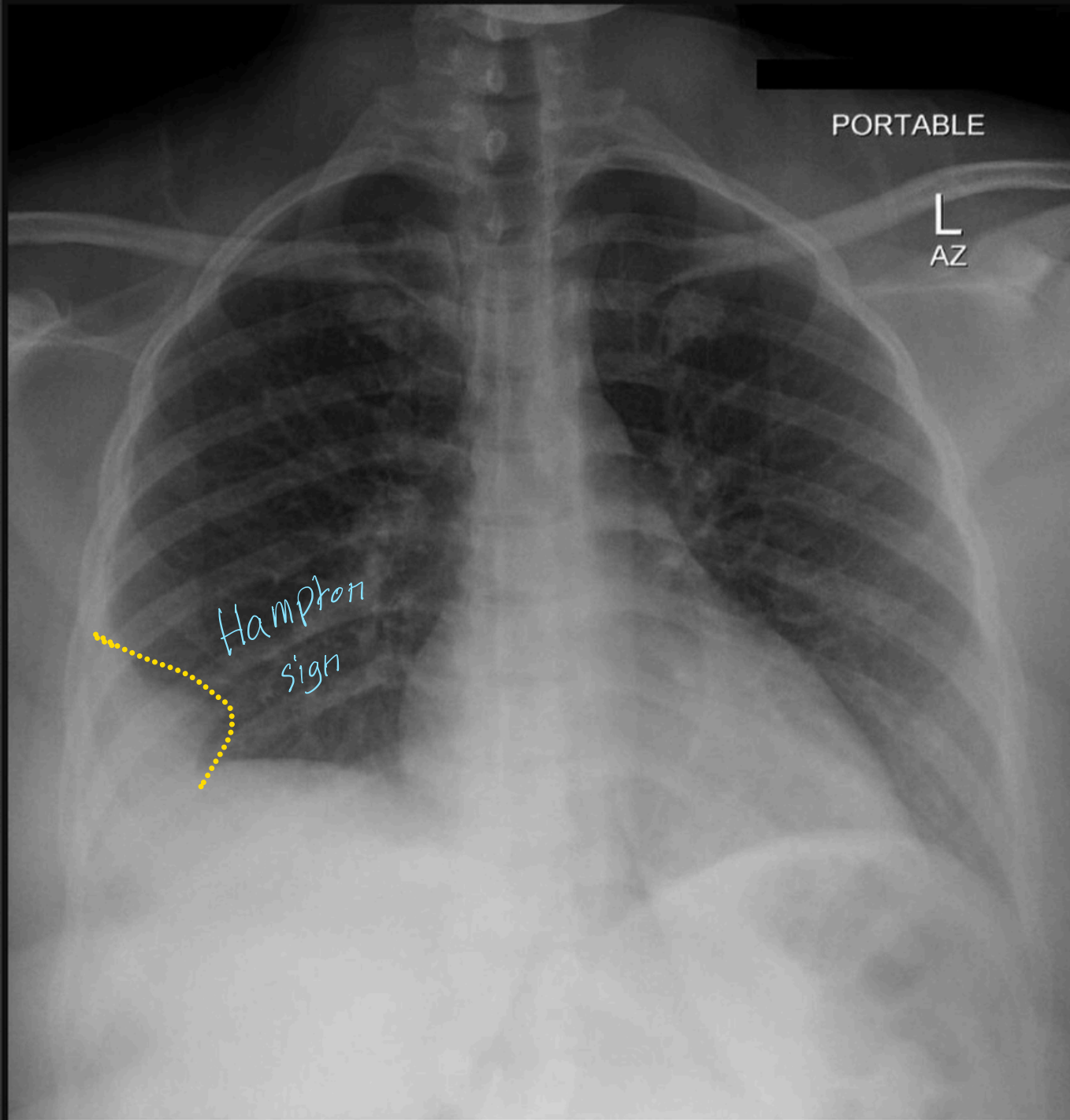
Pulmonary embolism (PE)

To diagnose depend on:

- ① Clinical Features
- ② D-dimer
- ③ X-ray

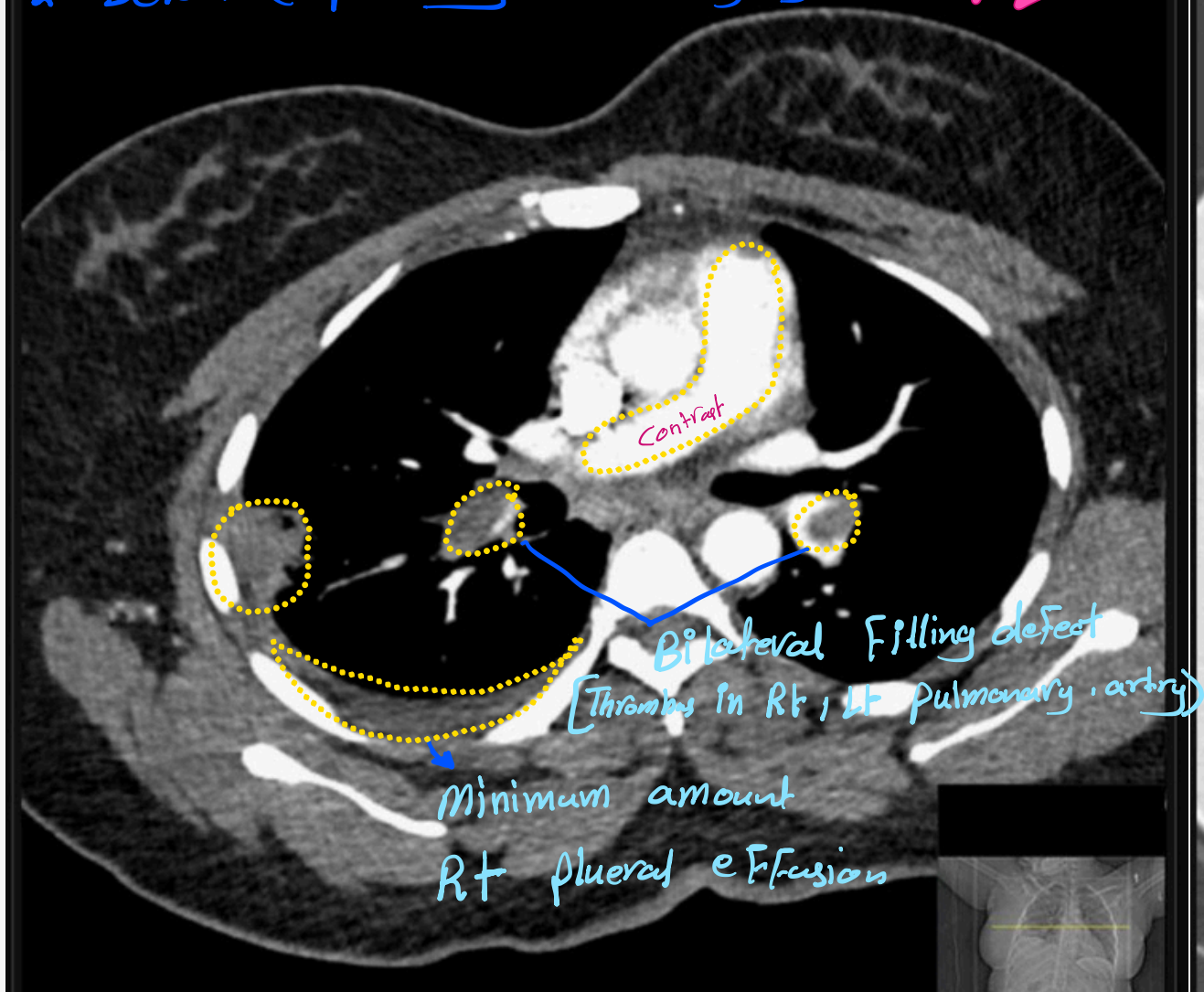
** But must make pulmonary CT Angiogram
Contrast dye used to detect pulmonary filling defect (CPE)*

- refers to embolic occlusion of the **pulmonary arterial system**. The majority of cases result from thrombotic occlusion, and therefore the condition is frequently termed pulmonary thromboembolism
- Chest radiography is **neither sensitive nor specific** for a pulmonary embolism. It is used to assess differential diagnostic possibilities such as pneumonia and pneumothorax rather than for the direct diagnosis of PE.
- **Hampton hump** refers to a **dome-shaped**, pleural-based opacification in the lung most commonly due to **pulmonary embolism** and **lung infarction**



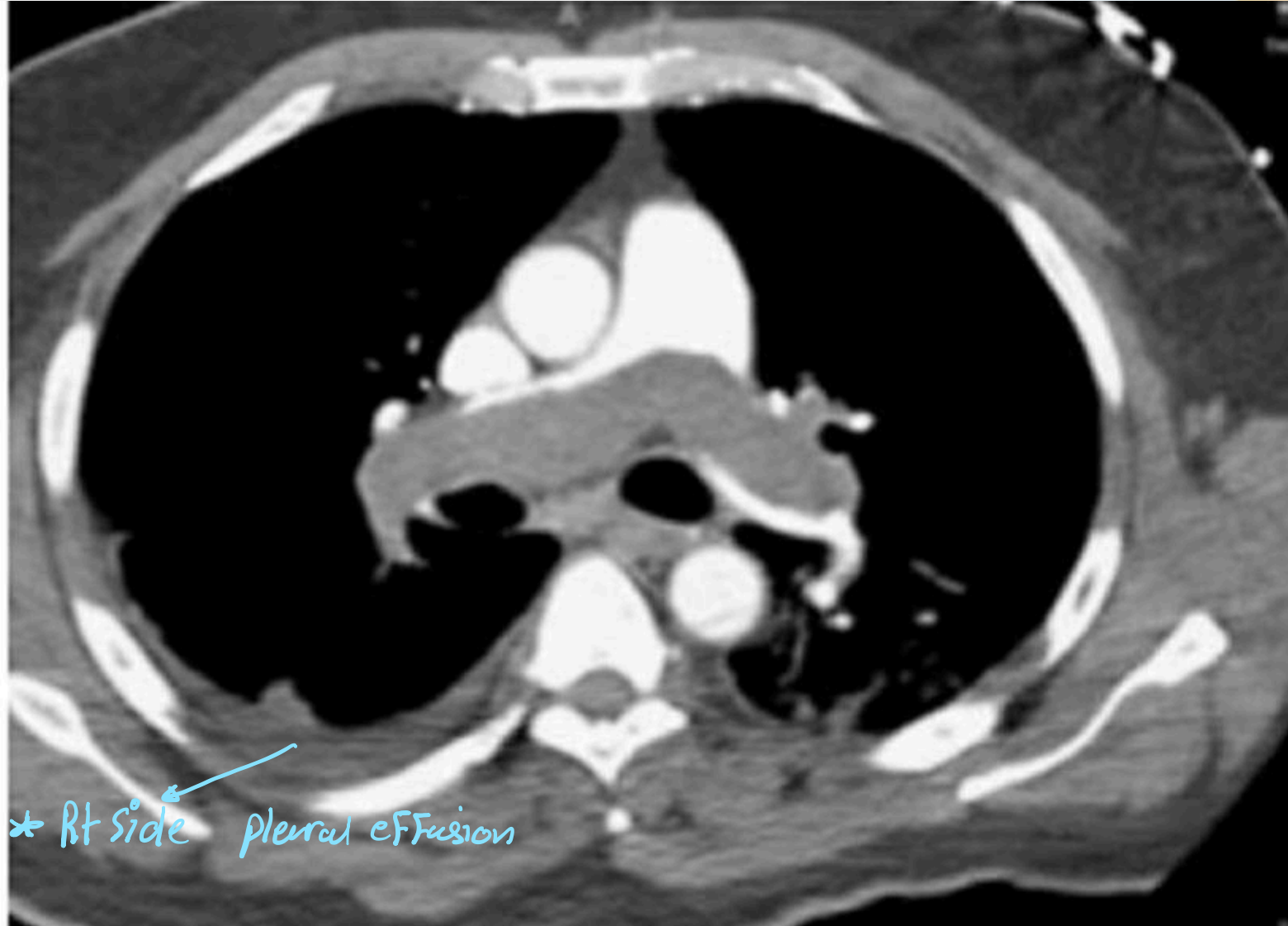
- CT pulmonary angiography (CTPA) will show filling defects within the pulmonary vasculature with acute pulmonary emboli.
- When the artery is viewed in its axial plane the central filling defect from the thrombus is surrounded by a thin rim of contrast, which has been called the **Polo Mint sign**.

* Selective pulmonary CT Angiogram PE



Saddle pulmonary embolism

- commonly refers to a large pulmonary embolism that ^{*}straddles the bifurcation of the pulmonary trunk, extending into the left and right pulmonary arteries.

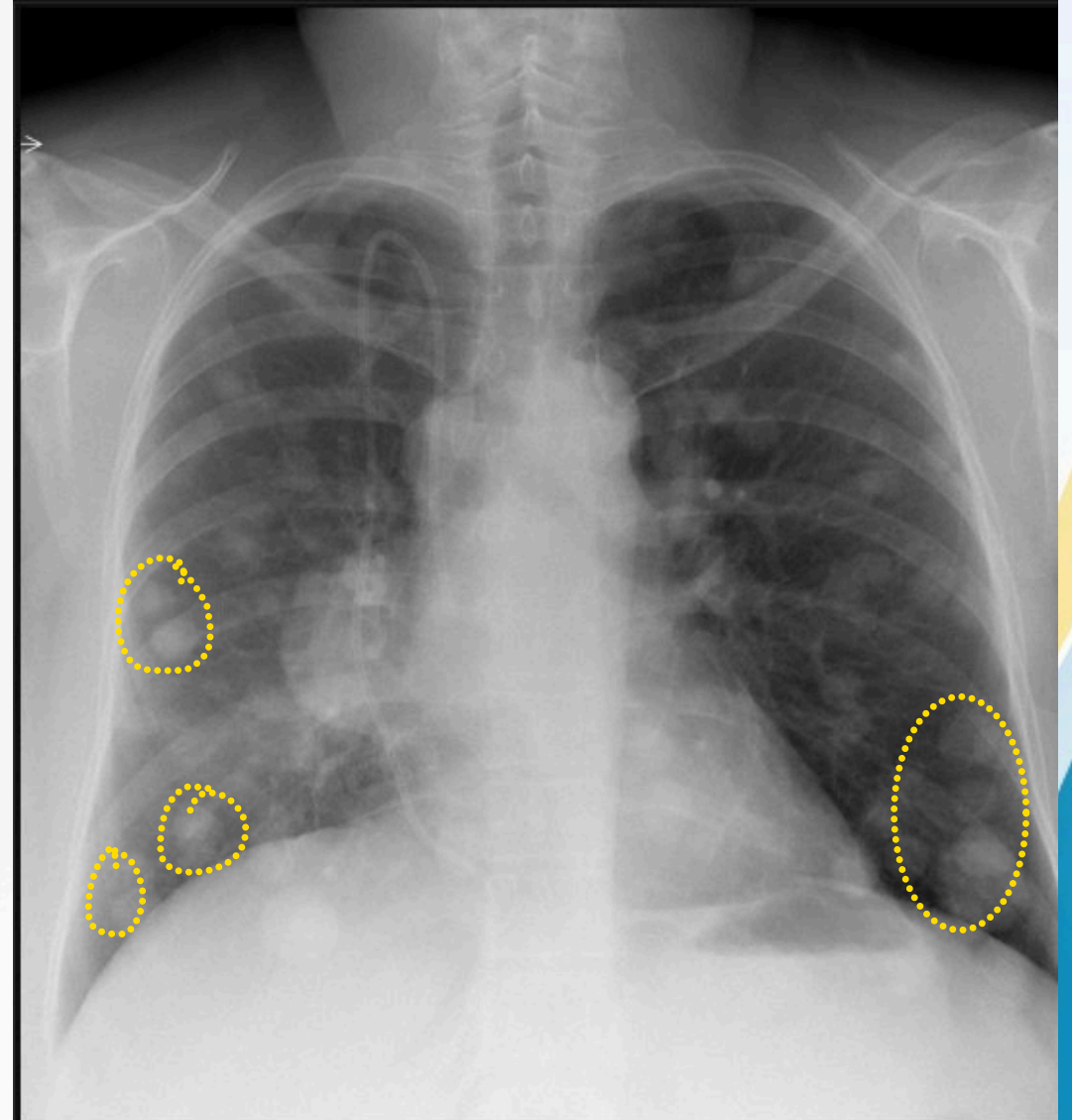


Metastases

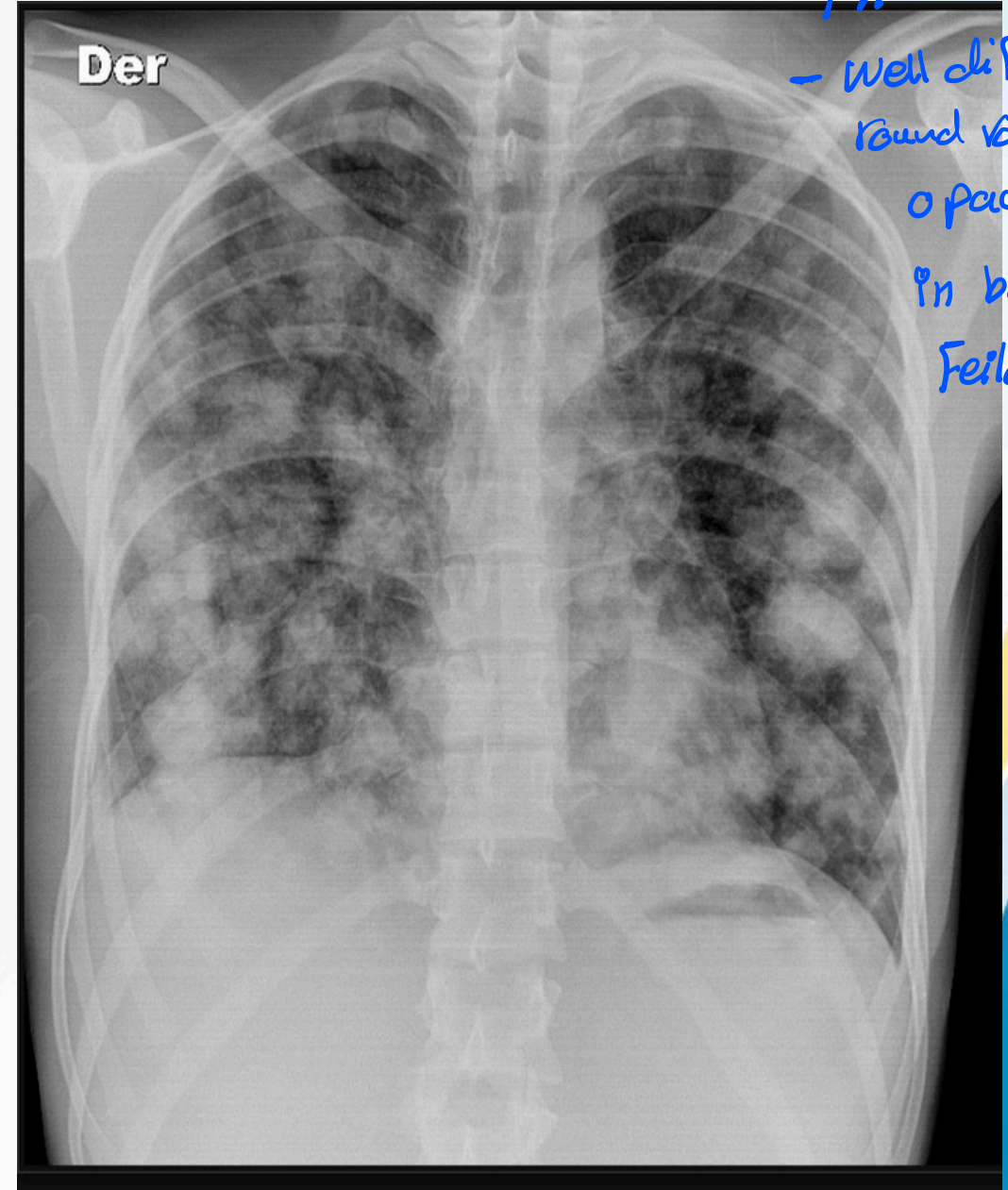
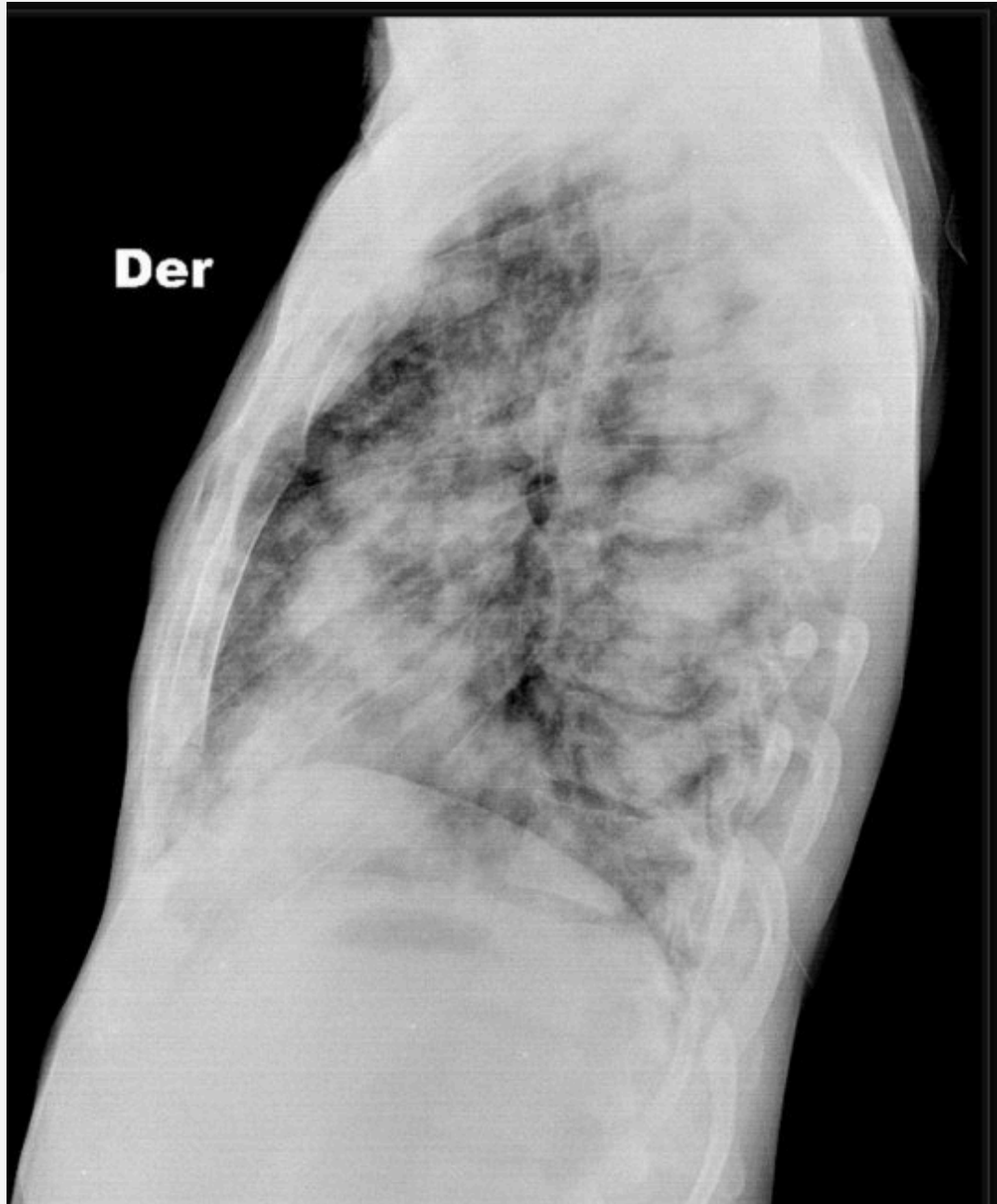
- Pulmonary metastases refer to distant tumor spread from a variety of primary tumors to the lungs via the blood or lymphatics.
- The most common malignancies to present with pulmonary metastases:
 1. Lung cancer (most common primary site)
 2. colorectal cancer
 3. renal cell carcinoma
 4. pancreatic cancer
 5. breast cancer

Plain radiograph

- Plain films are insensitive, although frequently able to make the diagnosis, as often pulmonary metastases are large and numerous.
- **Cannonball metastases** refer to multiple large, **well-circumscribed**, round pulmonary metastases that appear not unsurprisingly like cannonballs



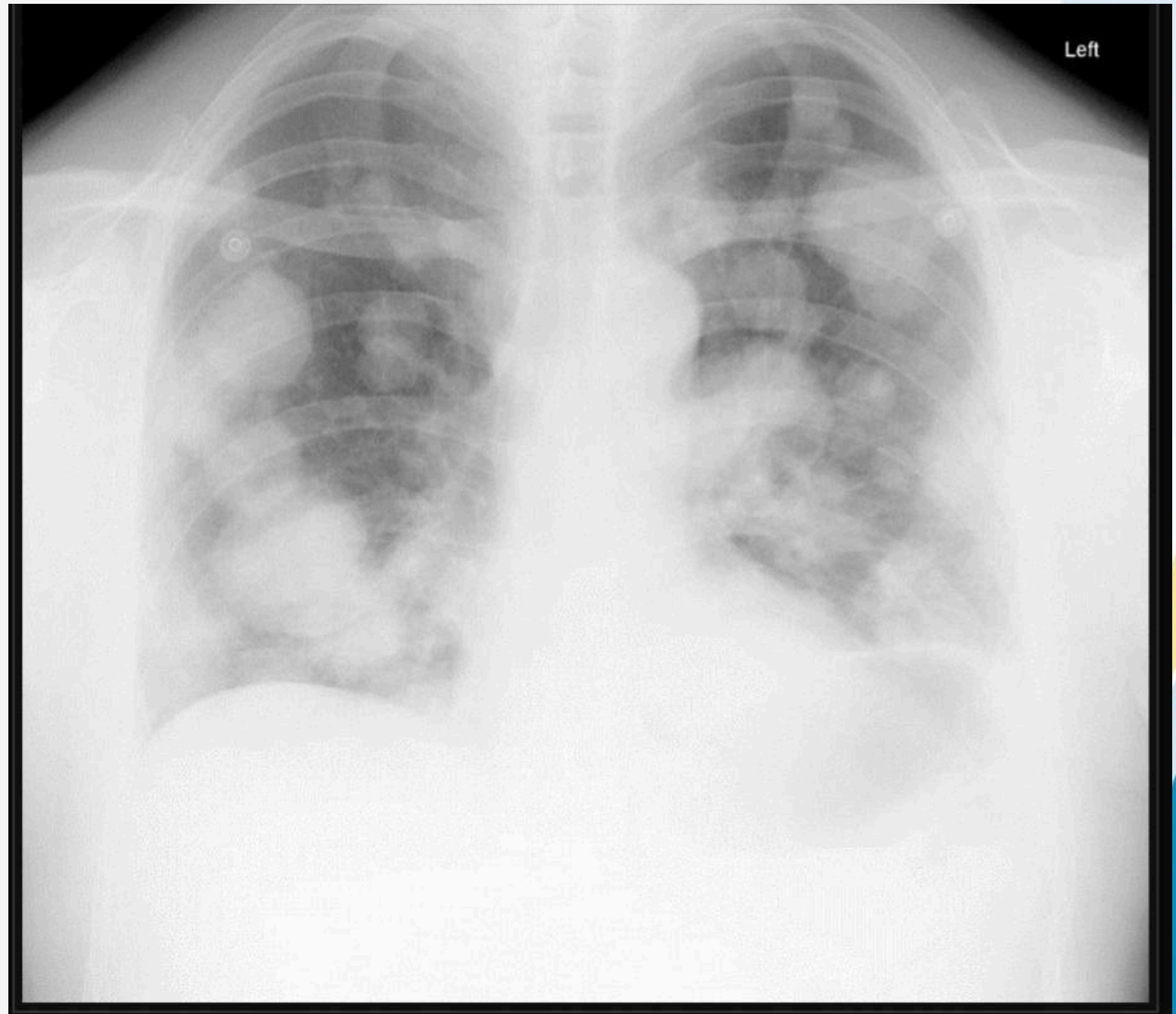
*در ۳۳ انچه ی در Normal پارامترها بتلصیب قیلم --
و کمان ادهاگان فیه Metastatic ی در Breast عیان اعرق ادهاگان در Metastatic



- PA

- well defined
round random
opacity
in both lung
field

Cannonball



* CT with Contrast

من مطلوب في كذا

فرق في
Angio
Amount of Contrast
+ earlier

Catiron ball

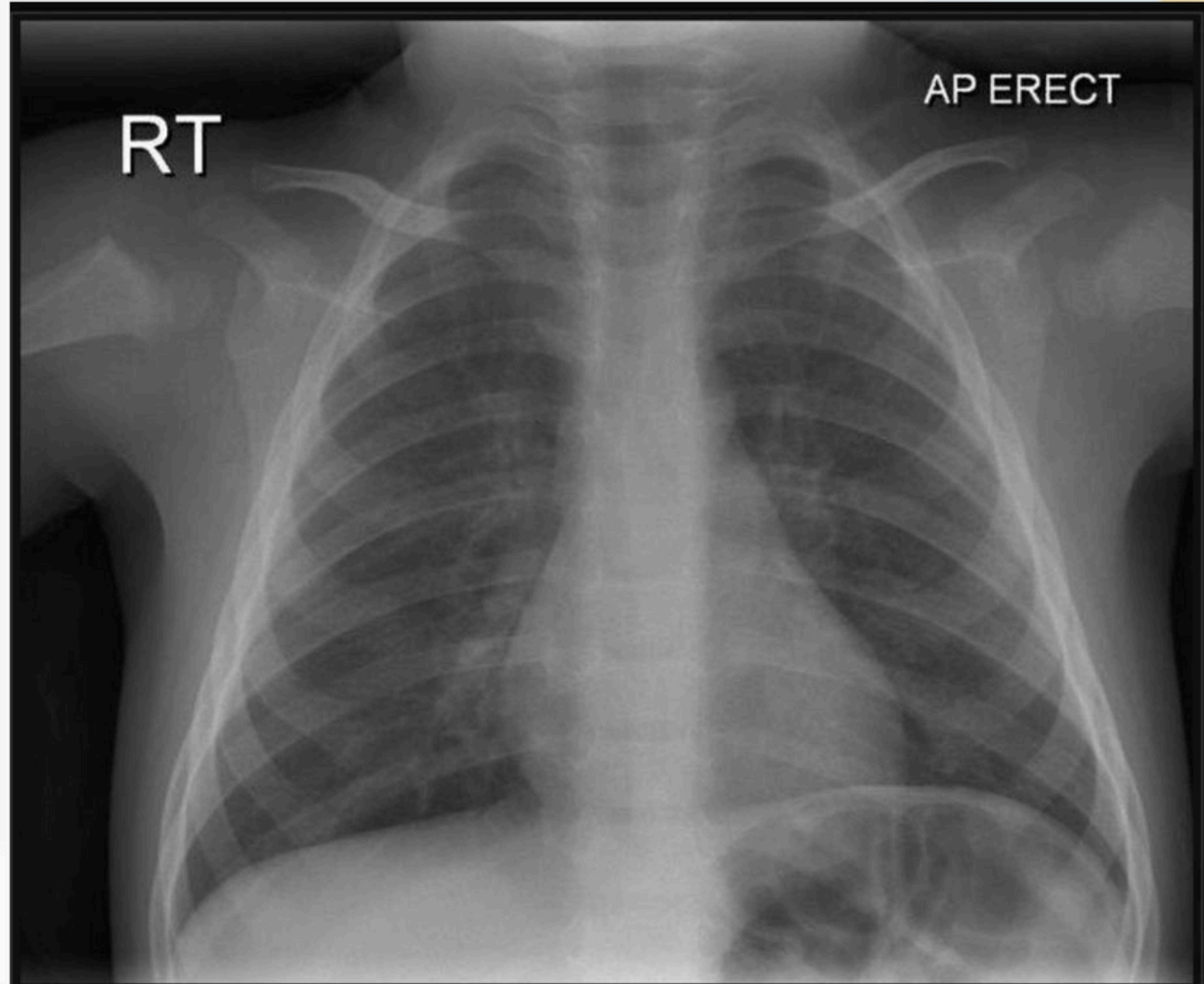
more in Rt

Bilateral pleural effusion.



Chest radiograph (pediatric)

- Depending on the patients' age, the difficulty of the examination will vary,
- often requiring a specialist trained radiographer familiar with a variety of distraction and immobilization techniques.



Standard projections

- As pediatrics vary in their level of cooperation, various projections can be utilized to suit the patient's needs and age:

- **PAerect**

performed on older patients (teenage years), not advisable for younger patients due to their attention span

- **APerect**

ideal for cooperative younger children (i.e. Between 3-7 years old) due to the ease of positioning and immobilization

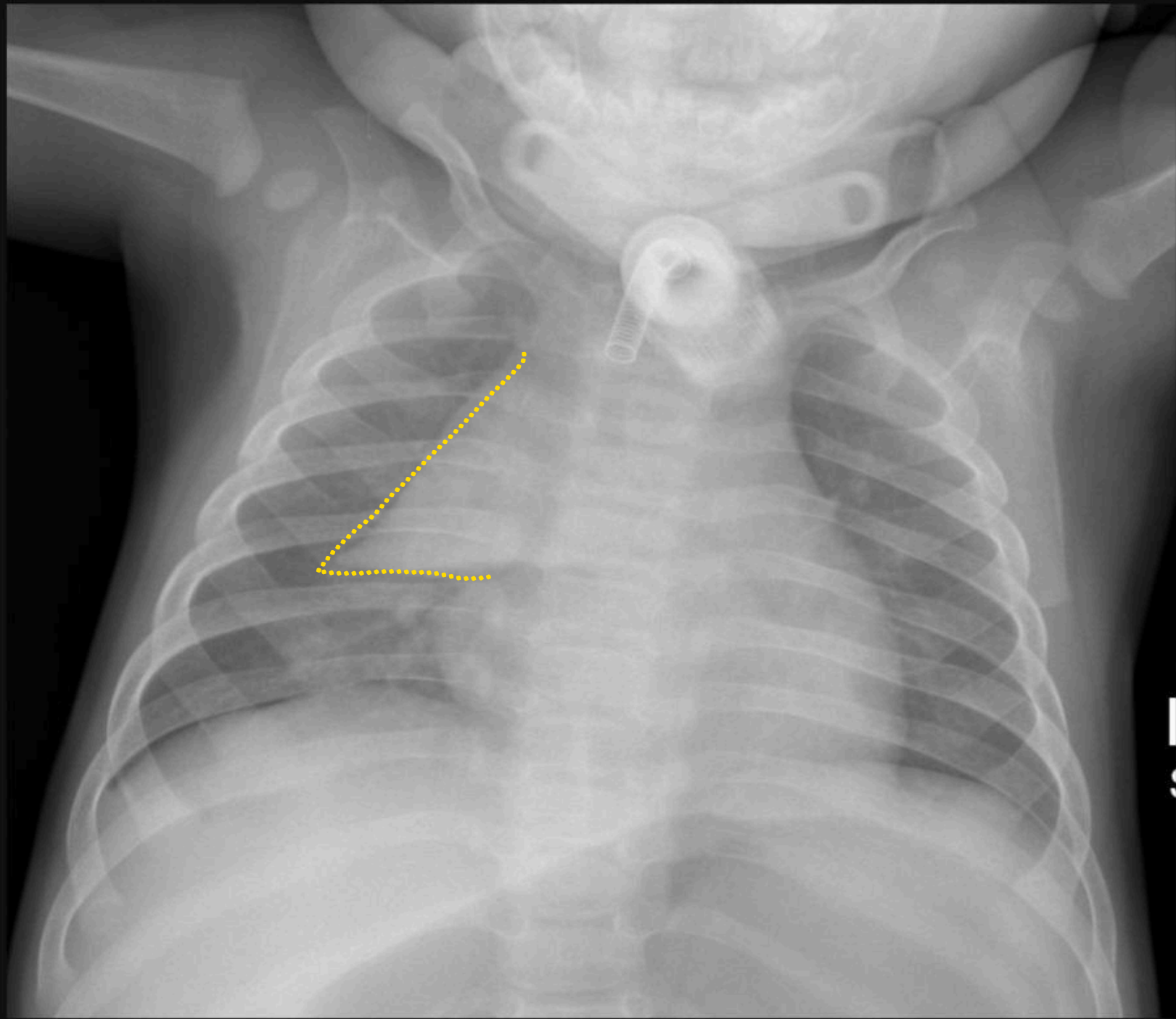
- **AP supine**

performed when imaging unconscious or uncooperative children

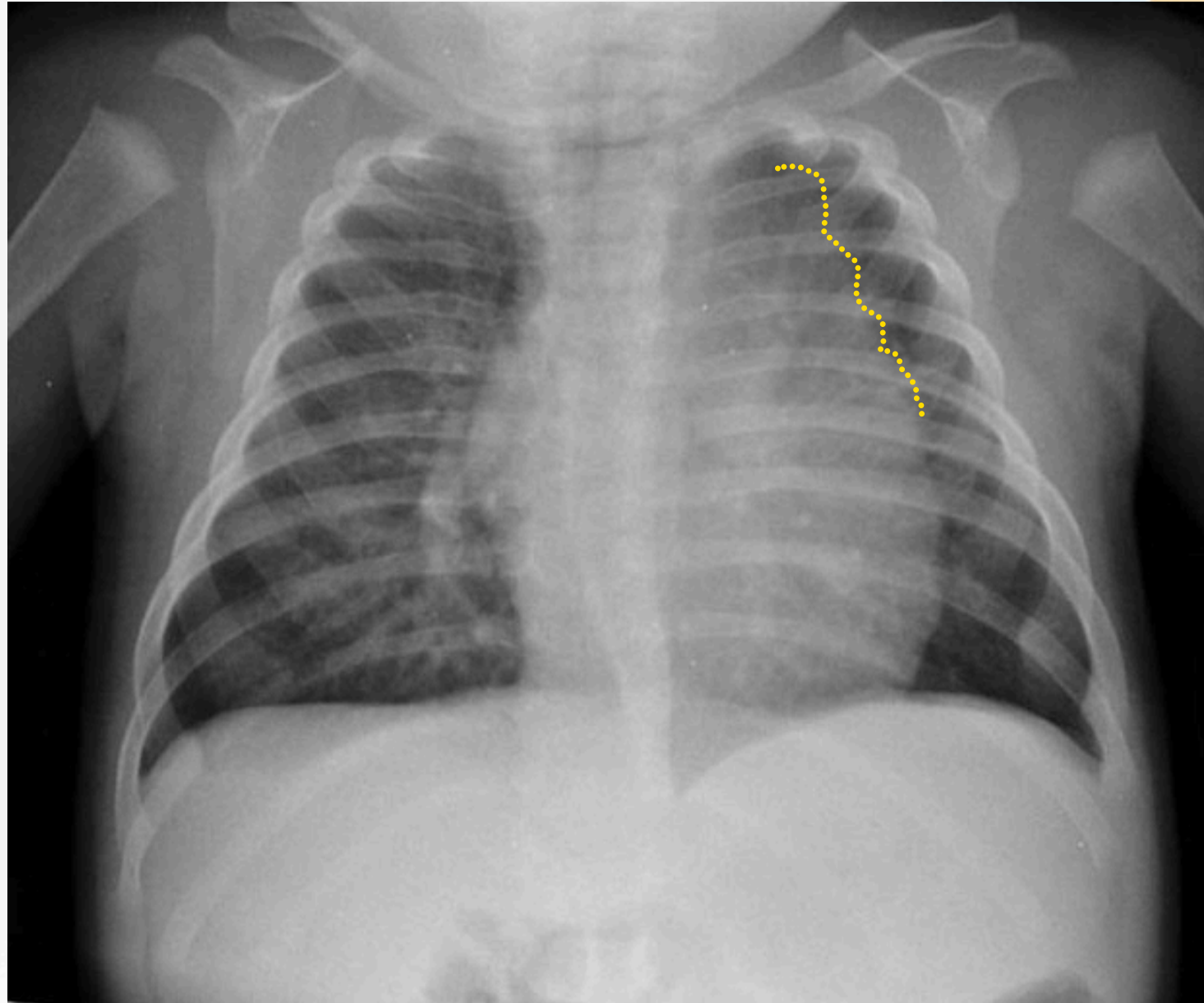
- **APsupine(neonatal)**

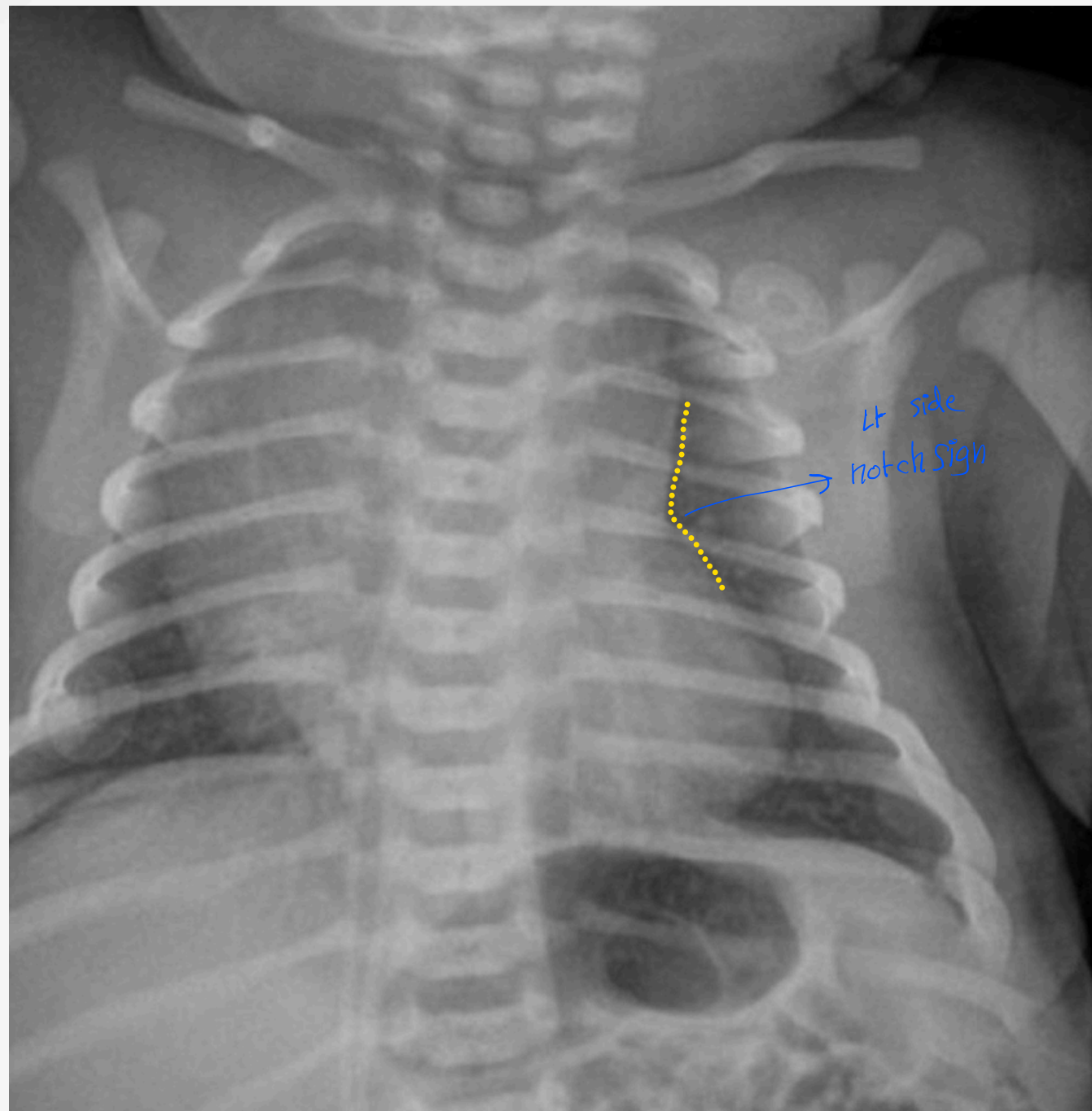
Thymic sail sign

- Represents a triangular-shaped inferior margin of the normal thymus seen on a neonatal frontal chest radiograph. It is more commonly seen on the right side, but can also be bilateral. It is seen in 3-15% of all cases.

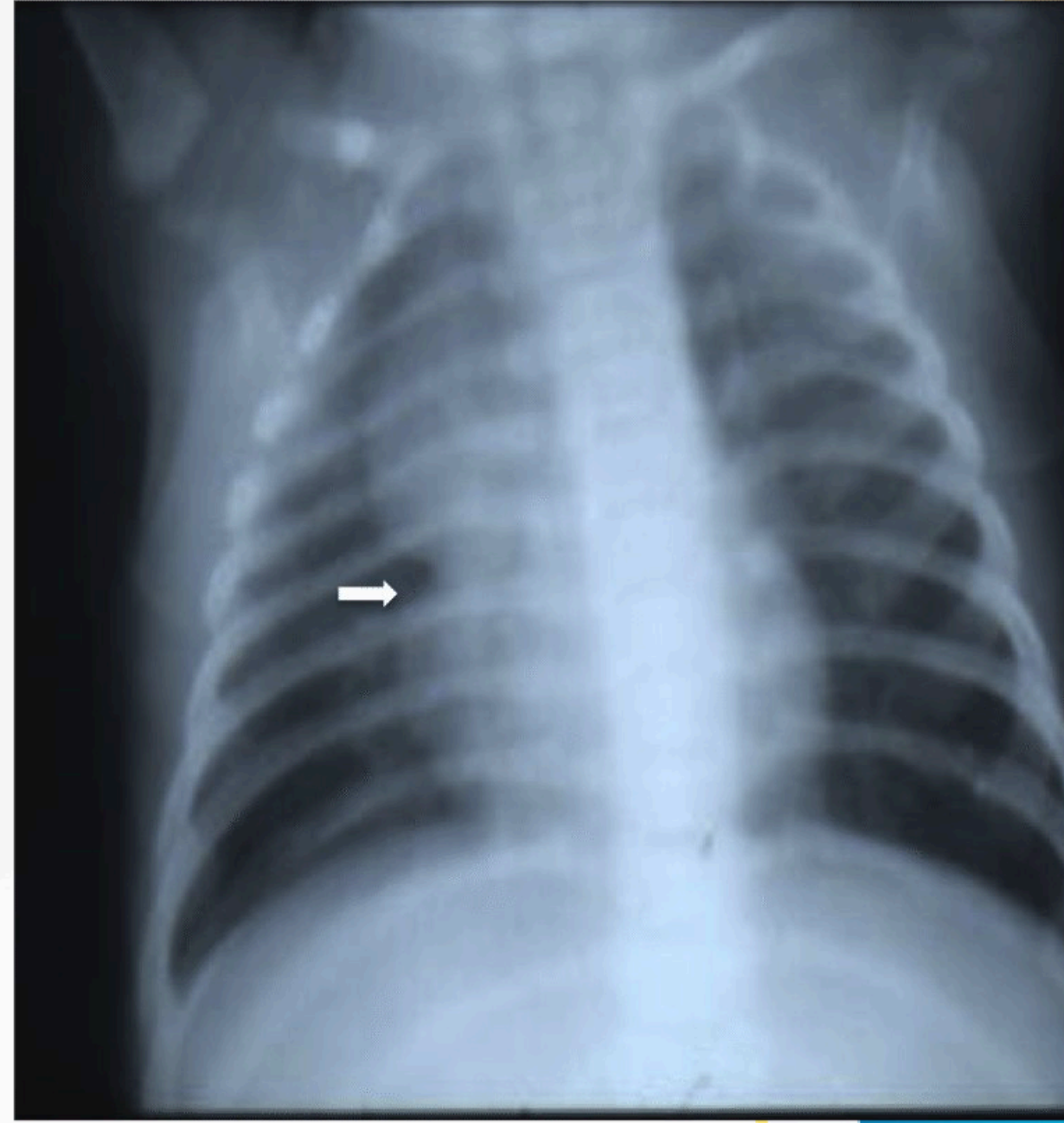


- **Thymic wave sign** refers to the indentation of the normal thymus in young children by the ribs, resulting in a wavy border on chest radiograph.





- The **thymic notch sign** represents the normal thymus in a newborn on a **frontal chest radiograph**.
- Interruption of the cardiac silhouette forms a notch, which may be seen on either side, but more frequently is seen on the left side.

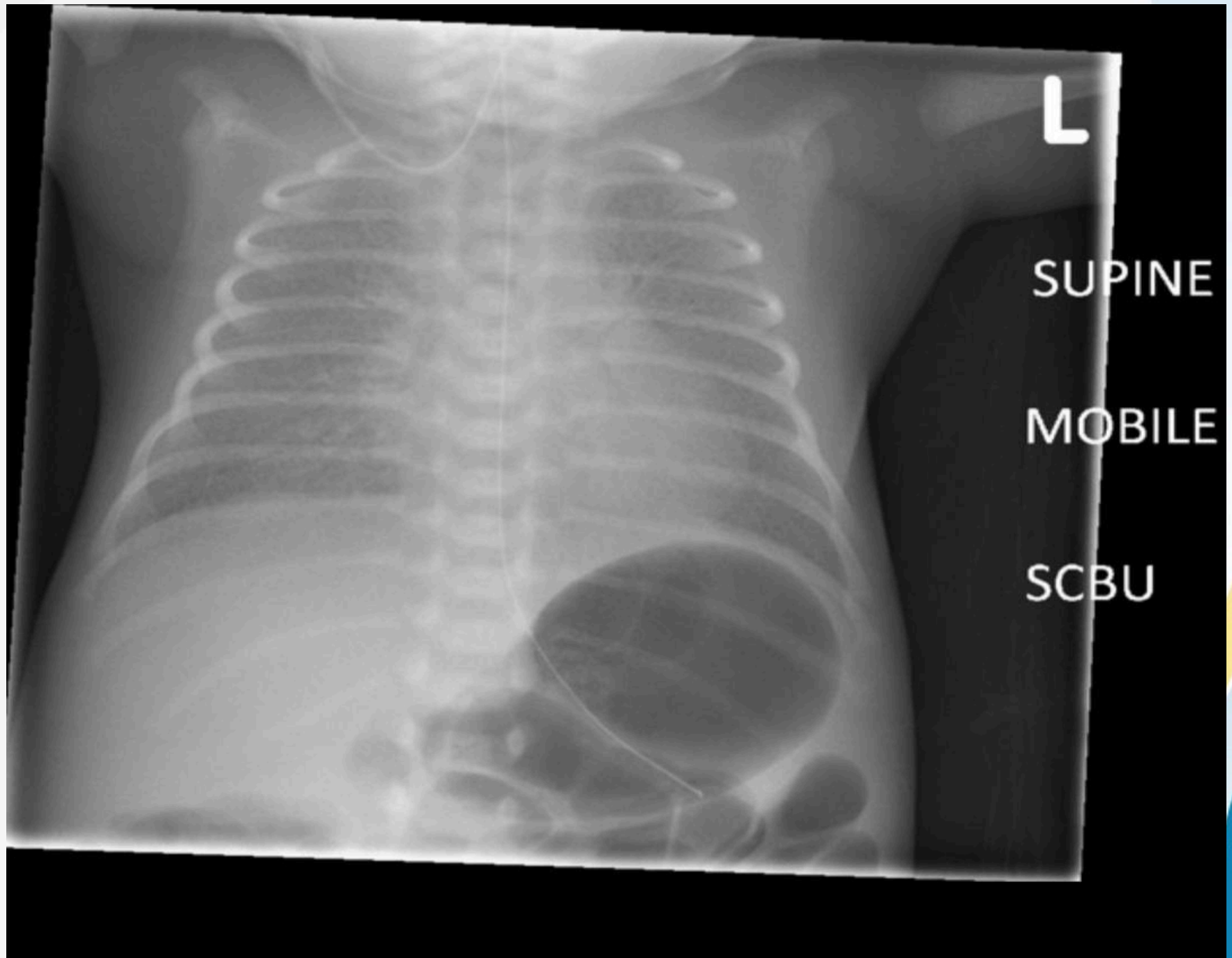


Respiratory distress syndrome (RDS)

- is a relatively common condition that occurs in preterm neonates resulting from insufficient production of surfactant.

- Risk factors

1. maternal diabetes
2. greater prematurity
3. perinatal asphyxia
4. multiple gestations



Pathology

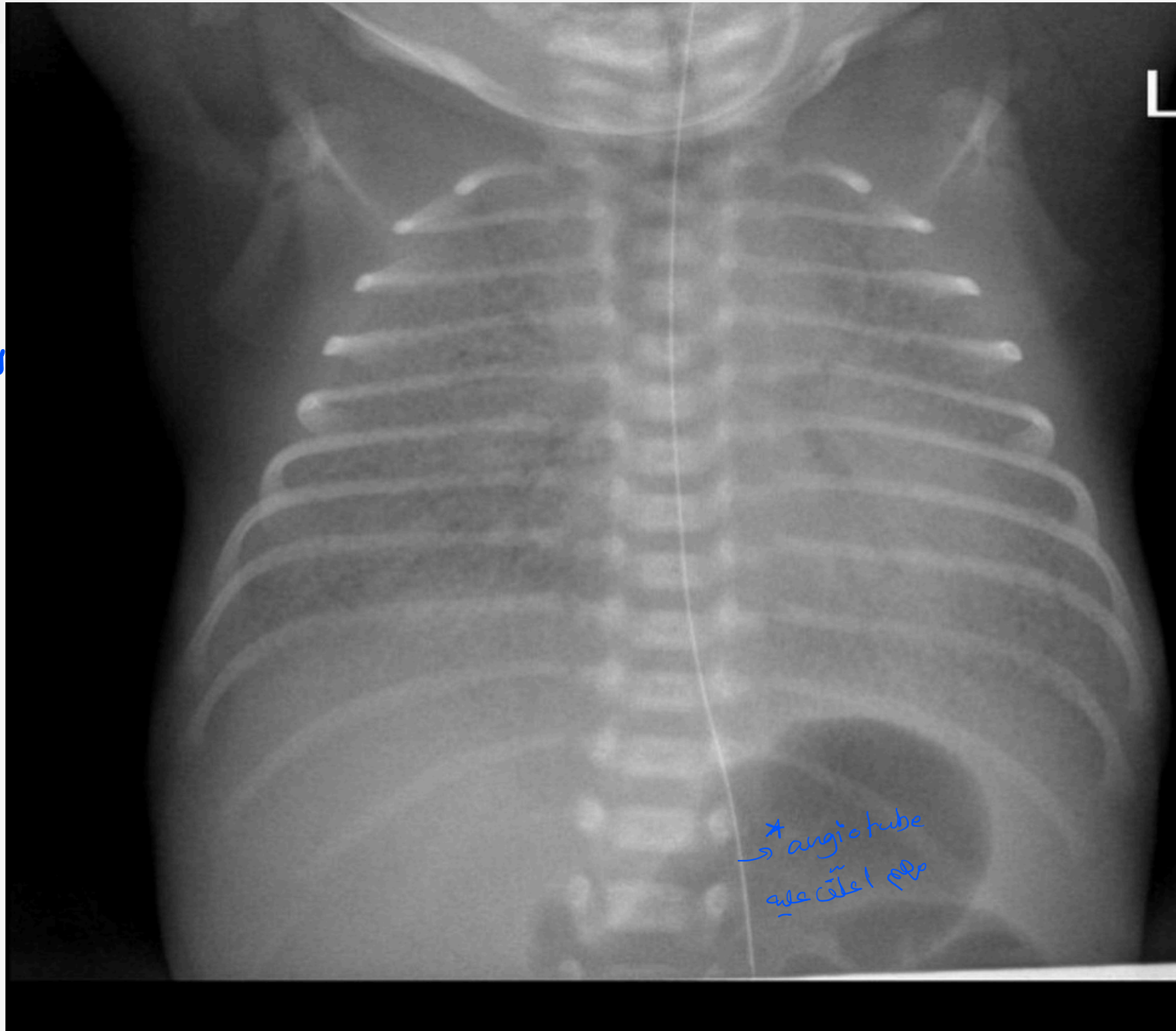
- Immature type II pneumocytes cannot produce surfactant. The lack of surfactant increases the surface tension in alveoli causing them to collapse. Patients have a decreased lecithin to sphingomyelin ratio. Damaged cells, necrotic cells, and mucus line the alveoli.
- As the alveoli are collapsed (microscopically), the lungs are collapsed macroscopically as well. It is a diffuse type of adhesive atelectasis.

Reporting :-

- Ap CXR

Bilateral symmetrical
granular lung opacity
with decreased lung
volume.

+ Normal



☆ Plain radiograph

1. low lung volumes diffuse
2. bilateral and symmetrical granular opacities
3. bell-shaped thorax
4. air bronchograms may be evident
5. Hyperinflation makes the diagnosis less likely, unless the patient is intubated.
6. If treated with surfactant therapy, there may be an asymmetric improvement as more surfactant may reach certain parts of the lungs than others.

Congenital diaphragmatic hernia

- There are two main types of congenital diaphragmatic hernia (CDH)s which are uncommon yet distinct entities that usually occur on the left side (80%) of the diaphragm

Bochdalek hernia: most common (95%), located posterolaterally and usually present in infancy *ex:- bowel continue from the abdomen to chest*

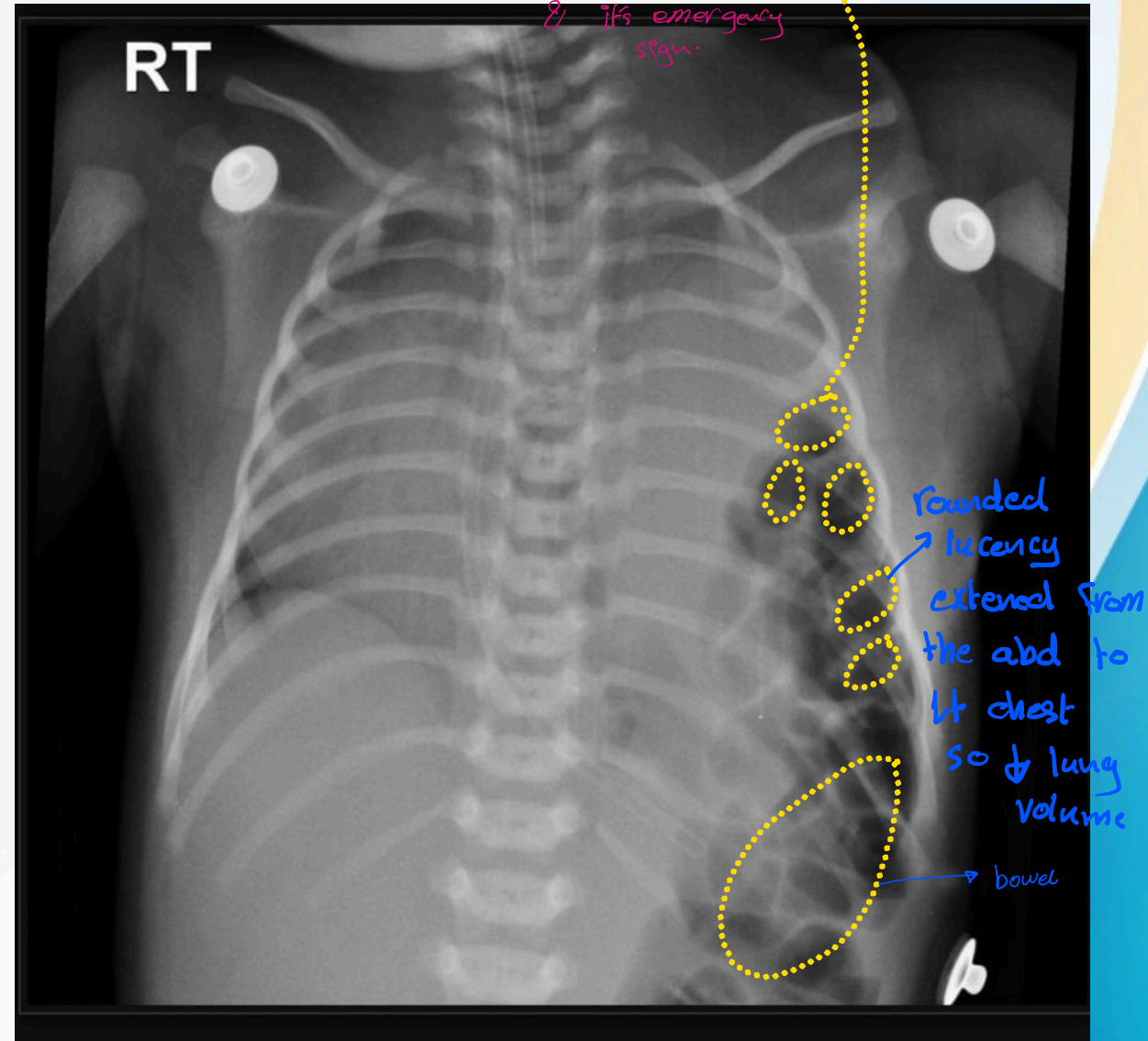
Morgagni hernia: smaller, anterior and presents later in life, through the sternocostal angles

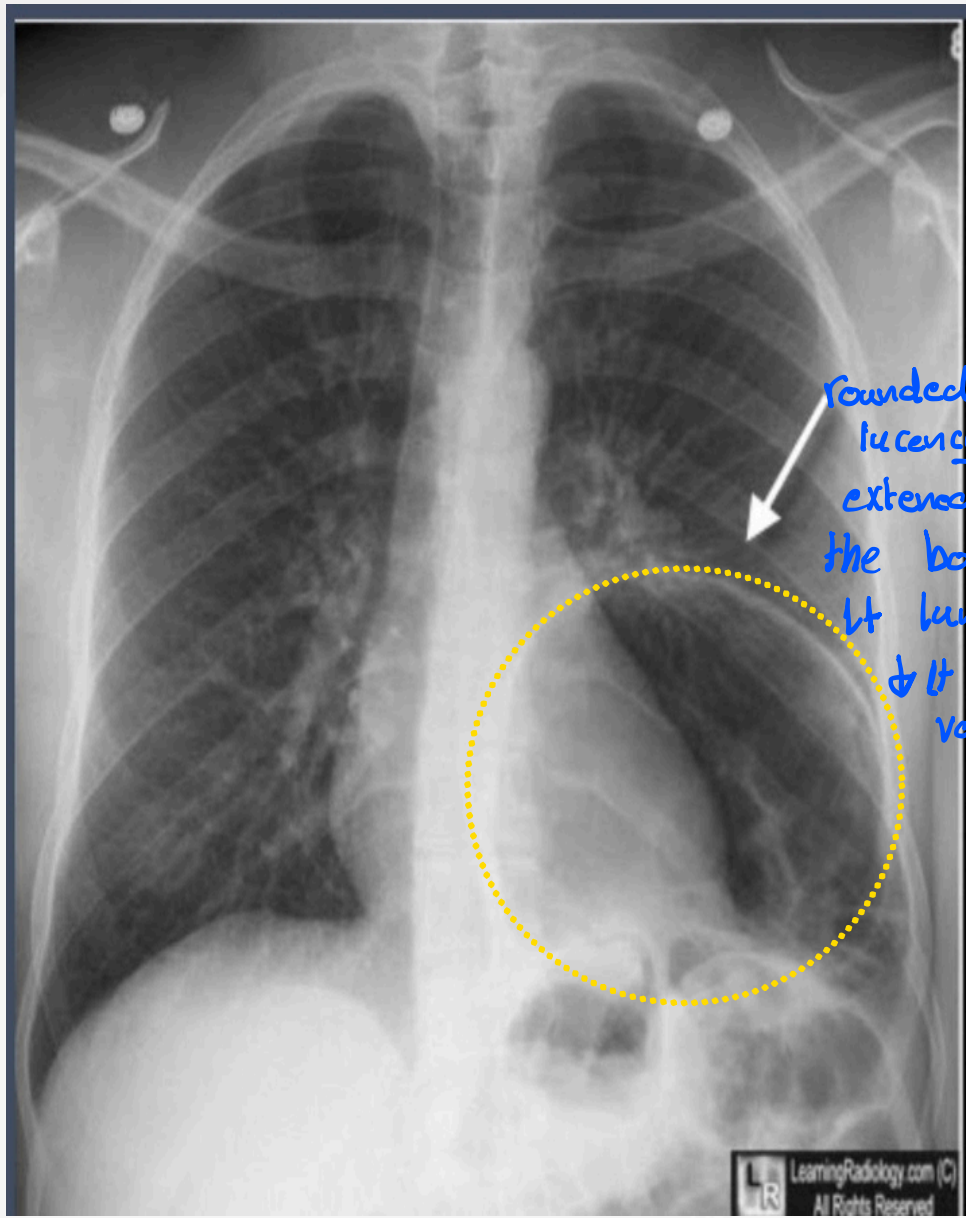
Bochdalek hernias

- also known as pleuroperitoneal hernias, (alternative plural: herniae) are the commonest type of congenital diaphragmatic hernia. They occur posteriorly and are due to a defect in the posterior attachment of the diaphragm
- **Retroperitoneal structures** may prolapse through the defect, e.g. Retroperitoneal fat or left kidney.

Plain radiograph

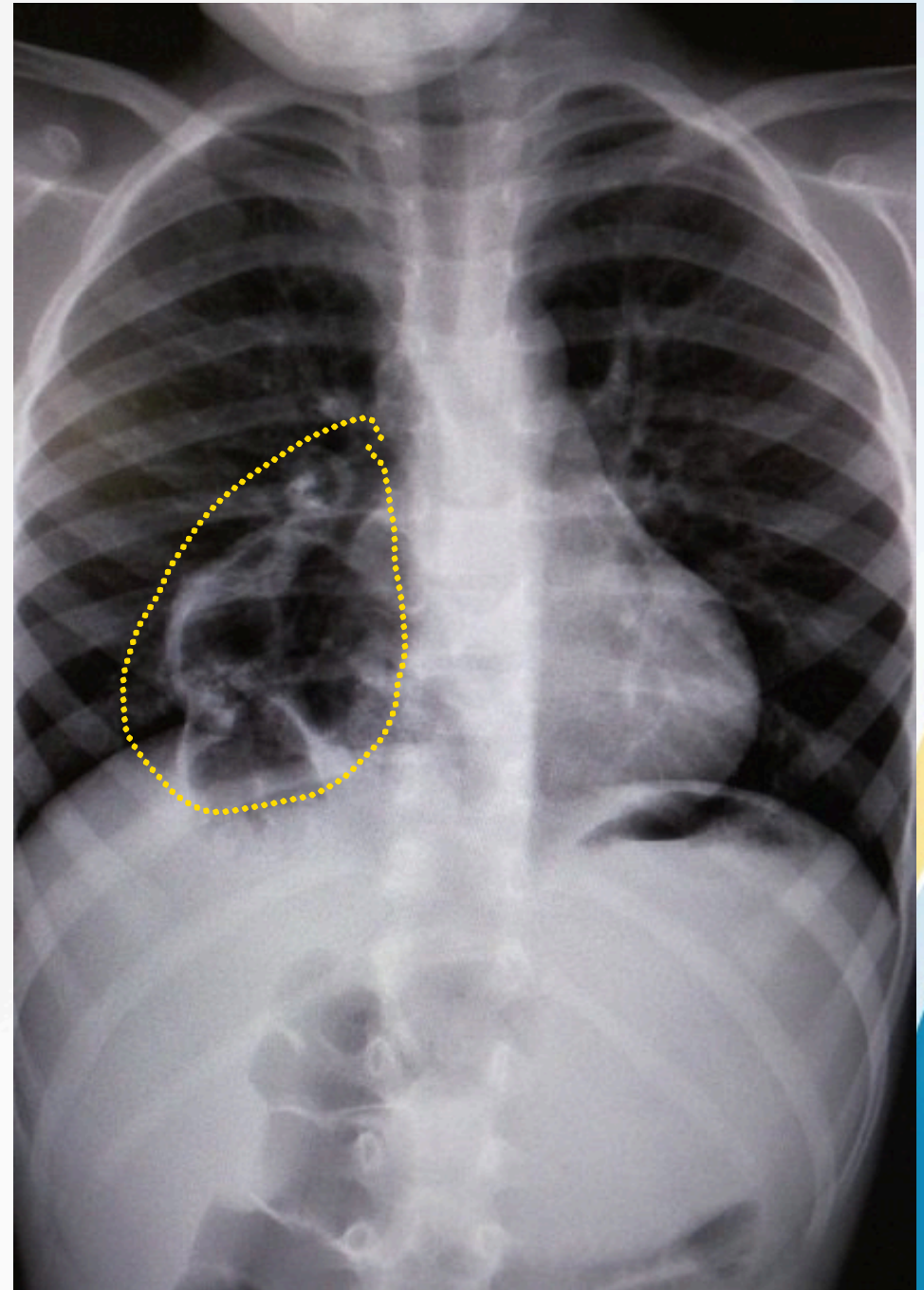
On conventional radiographs, the hernia may appear as a lung base soft-tissue opacity lesion seen posteriorly on lateral images.





Morgagni hernia

- Morgagni hernias (alternative plural: herniae) are one of the congenital diaphragmatic hernias (CDHs) and are characterized by herniation through the foramen of Morgagni.



When compared to Bochdalek hernias, Morgagni hernias tend to be:

- Anterior more often right-
- sided 90% s mal l rare 2% at
- low risk of prolapse
-
-

