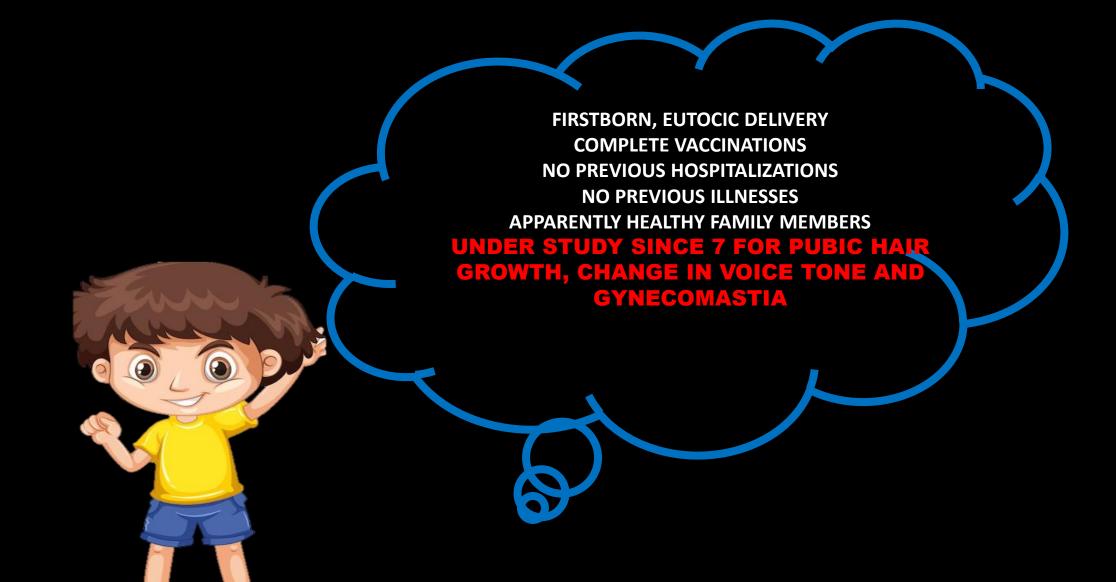
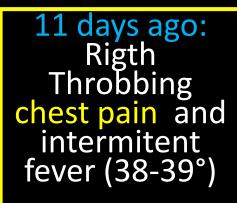
# PRIMARY MEDIASTINAL CORIOCARCINOMA IN A MALE CHILD: CASE REPORT



- ✓ To review the clinical history, radiological images and evolution of a case of primary mediastinal choriocarcinoma.
- ✓ Discussion of mediastinal germ cell tumors: classification and main features; with emphasis on primary mediastinal choriocarcinoma.

✓ Conclusions.







7 days ago:

Decreased appetite

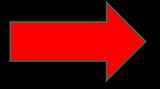


# 5 days ago:

Dry cough, fatigue and continous fever (3/d)

Treated at home with NSAIDs and Acetaminophen.

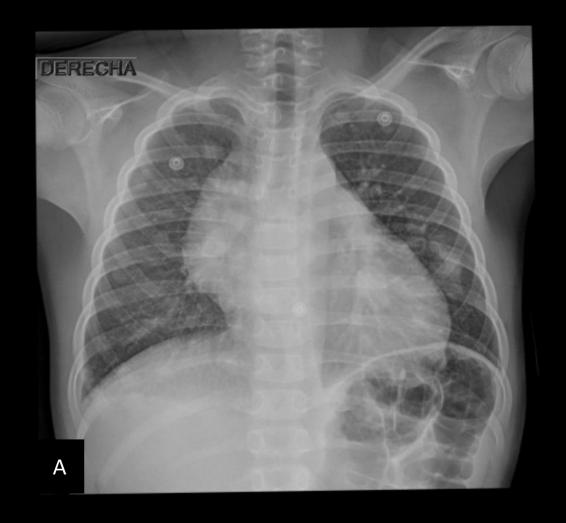
Starts requering pillows to sleep and have difficulty speaking

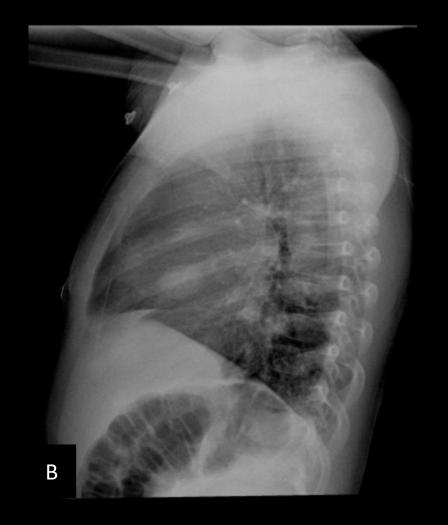


#### Today:

Referred to emergency from outpatient clinic due to respiratory difficulty, is intubated.



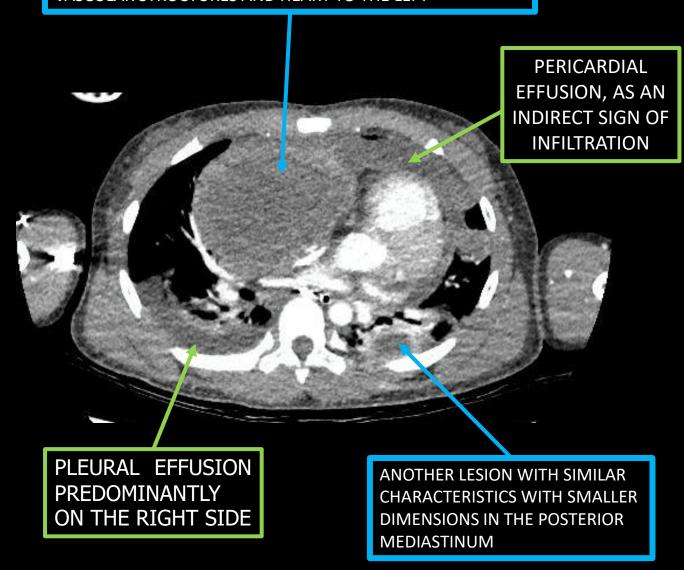


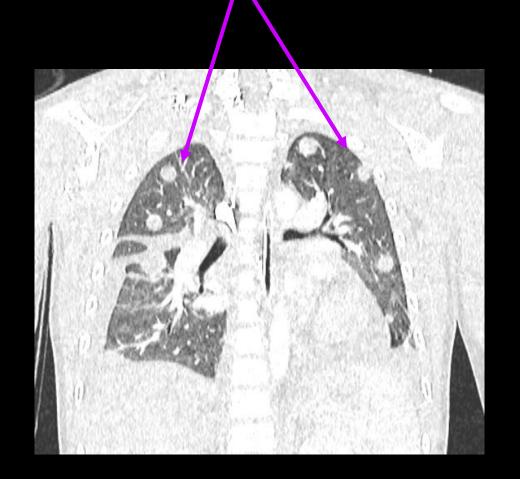


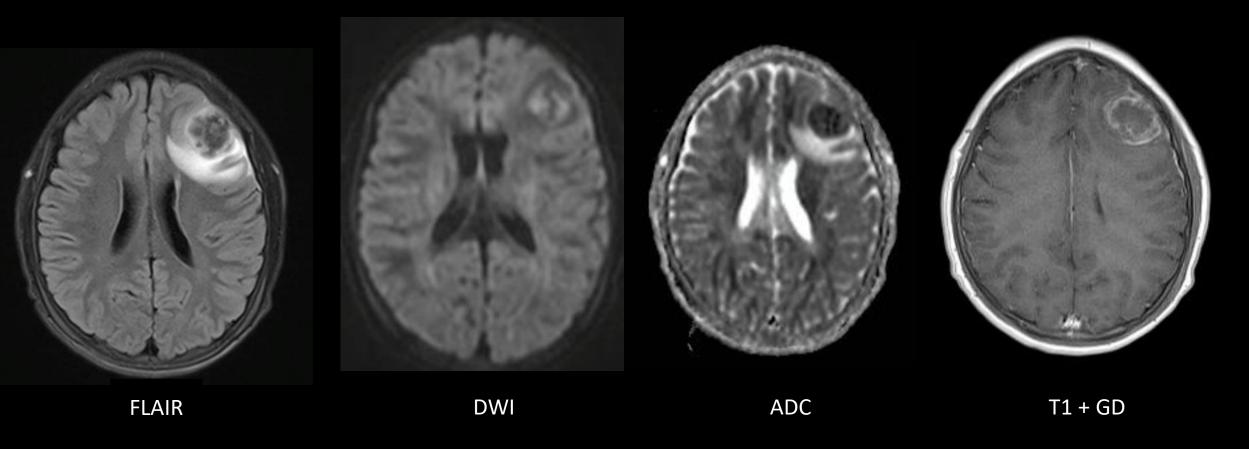
THE ANTEROPOSTERIOR (A) AND LEFT LATERAL (B) CHEST X-RAYS SHOW MARKED WIDENING OF THE MEDIASTINUM WITH INCREASED CARDIAC SILHOUETTE AND MULTIPLE NODULAR LESIONS IN BOTH LUNG FIELDS.

HETEROGENEOUS HYPODENSE LESION IN PREVASCULAR MEDIASTINUM, POORLY CONTRAST-ENHANCING, MEASURING 117 X 73 X 110 MM (T X AP X L), DISPLACING VASCULAR STRUCTURES AND HEART TO THE LEFT

ASSOCIATED WITH MULTIPLE METASTATIC NODULAR LESIONS IN DIFFERENT LUNG SEGMENTS SUGGESTIVE OF CANNOBALL MESTASTASES

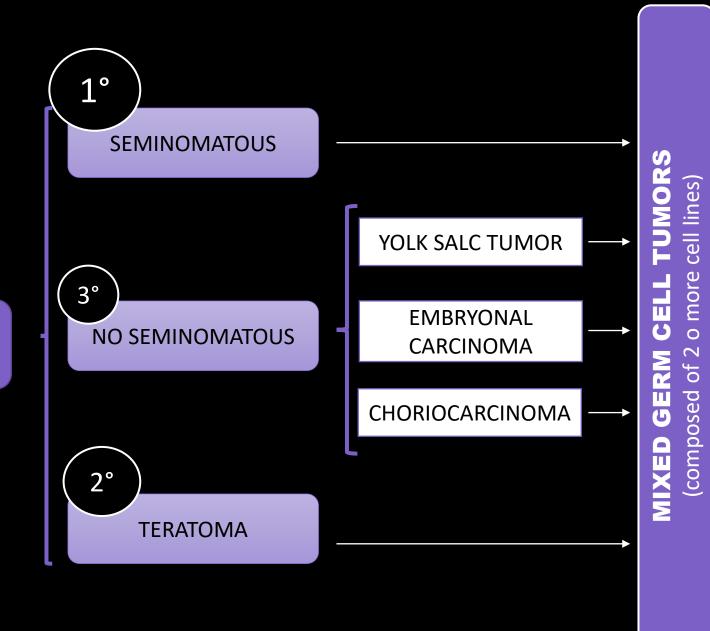






INTRAXIAL HETEROGENEOUS SOLID EXPANSIVE LESION WITH SOME CENTRAL CYSTIC AREAS LOCATED TOWARDS THE LEFT FRONTAL CONVEXITY, *ASSOCIATED VASOGENIC EDEMA*, PRESENTS *DIFFUSION RESTRICTION* AND *SCARCE ENHANCEMENT AFTER CONTRAST ADMINISTRATION*, MEASURES DT:23MM X DAP:21MM, CONTANTED WITH DURA MATER. VENTRICULAR SYSTEM PRESERVED.NO SIGNIFICANT DISPLACEMENT OF THE CEREBRAL MIDLINE. FINDINGS HIGHLY SUGGESTIVE OF BRAIN METASTASIS.

# DISCUSSION AND DIAGNOSIS



**Mediastinal germ** 

cell tumors

WHO 4th edition

Germ cell tumor with

**SOMATIC TYPE** 

**SOLID MALIGNANCY** 

Germ cell tumor with ASSOCIATED

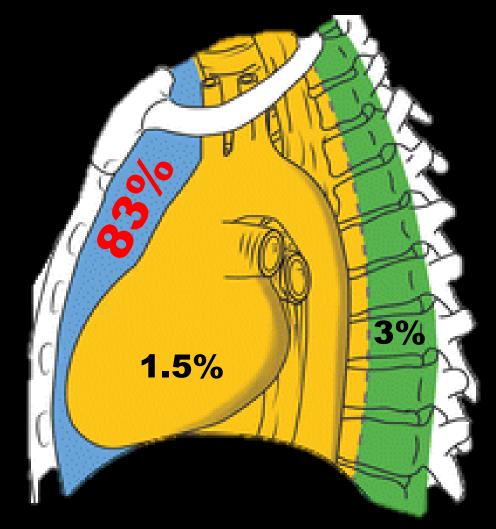
**HEMATOLOGICAL** 

**MALIGNANCY** 

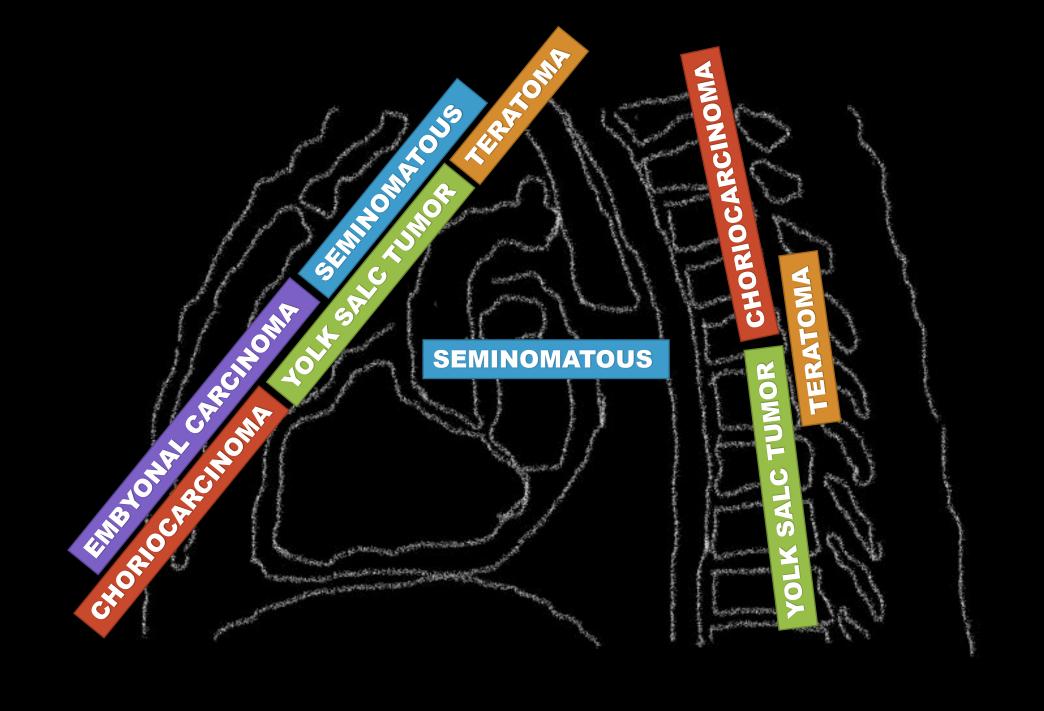


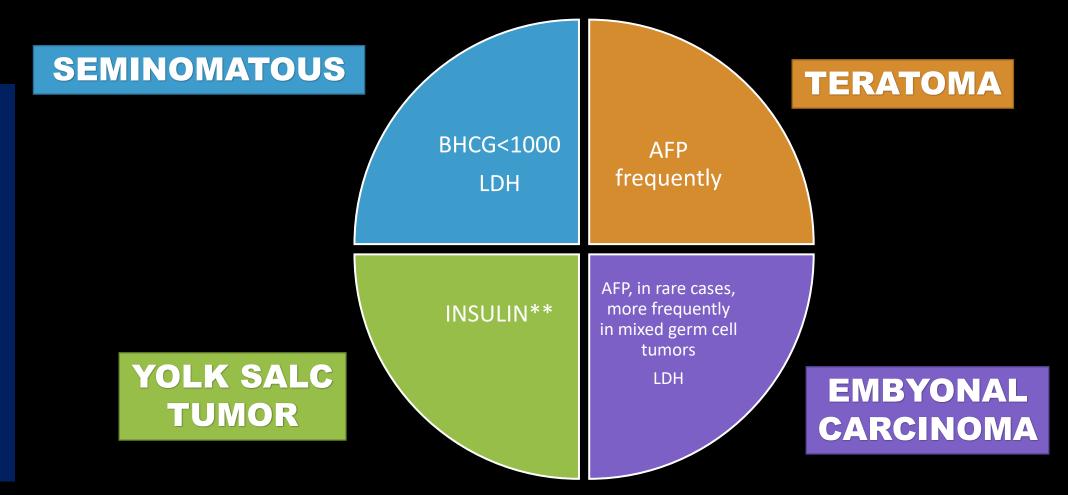
Most germ cell tumors are diagnosed in YOUNG MEN

M:F = 50: 1 Age of Dx: 18-29 (48.5%)



**Mixed 13%** 





<sup>\*</sup> BHCG> 1000 is consistent with choriocarcinoma

<sup>\*\*</sup> in rare cases may contain pancreatic endocrine cells, which produce insulin and hypoglycemia

### **SEMINOMATOUS**

Homogeneous mass Low enhancement No invasiva

#### **Clinical Symptoms:**

Symptomatic due to mass efect

#### Prognosis:

- Survival at 5 years: 88-100%
- Correlates with high LDH levels

# **TERATOMA**

#### Tumour with different tissue types

Mature: well differentiated, mature cell lines Immature poorly differentiated, embryonic/fetal tissue

Mass with different type of tissues (fat, calcifications). Cystic component, helps to differentiate from other tumors.

#### Prognosis:

Good prognosis after surgical resection

#### **NO SEMINOMATOUS**

### YOLK SALC TUMOR

Heterogeneous mass

High enhancement

Invasive

Tipically without calcifications

MRI: Cystic áreas with significant difusión restriction

With pulmonary metastases at the time of diagnosis

#### Prognosis:

Poor, survival up to 40%

\*\*< 5 years and female, think almost exclusively of this tumor.



#### **EMBYONAL CARCINOMA**

Largr mas WITHOUT ENHANCEMENT Compresses and invades large vessels

Pulmonary, lymph node and hematologic metastasis

#### Prognosis:

Poor, immunotherapy treatment (under investigation)

Uncommon tumor, almost exclusive to men over 30



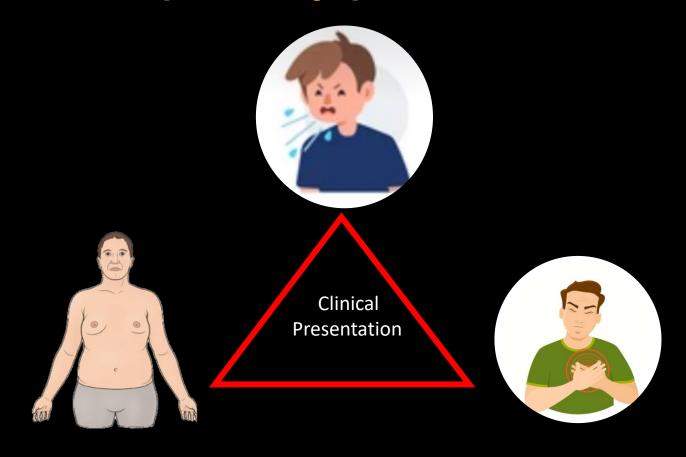
"Mediastinal choriocarcinoma without a detectable primary in the gonads or metastasic desease in the retroperitoneal lymph nodes"

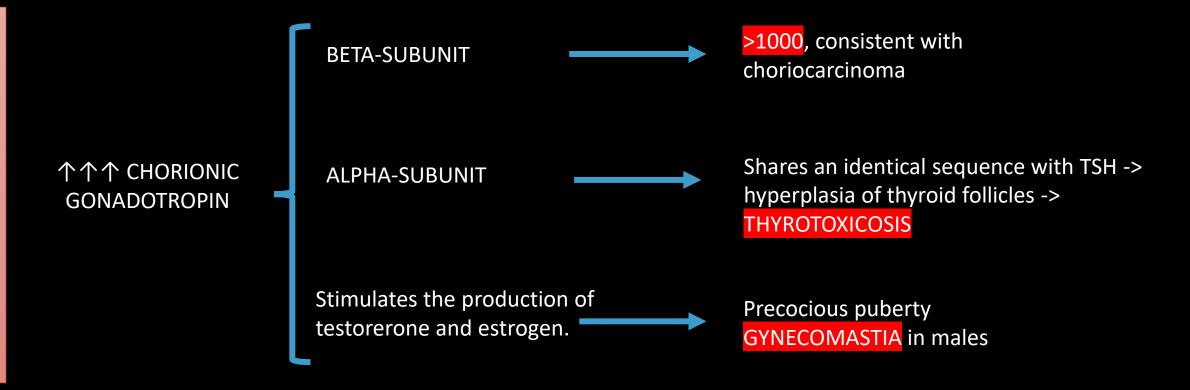
Extremely rare, less than 50 cases reported to date

Nearly exclusive of young men.

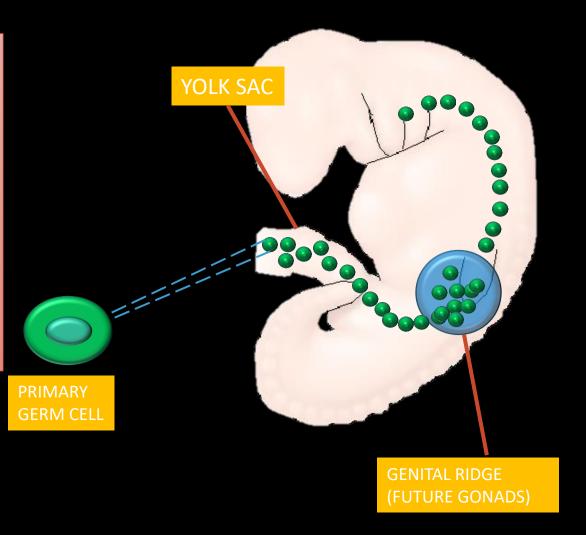
Associated with Klinefelter Syndrome (10 times more frecuent)

Usually asintomatic. in advanced cases presents with the triad of dyspnea, cough and gynecomastia





<u>DIAGNOSIS CERTAINTY:</u> Increased serum levels of *B- hCG* and presence of *cytotrophoblast* intermixed with syncytriotrophoblast at patology exam



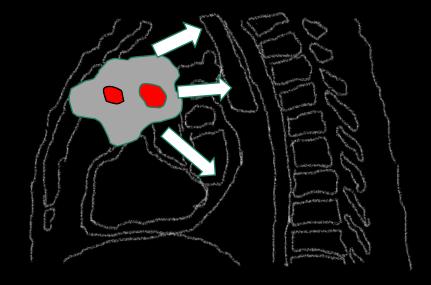
They normally migrate along side the genital ridge to localice at the future gonads

An arrest anywhere in this path may lead to a primary mediastinal Choriocarcinoma

These cell can remain dormant unitl puberty or later sexual life, when some stimulus can causa them to mature into a tumor mass

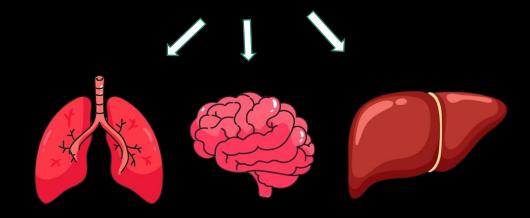
Large solid mediastinal mass mostly in the prevascular mediastinum with poor defined margins

#### HIGHLY LOCALLY INVASIVE



Great vessels: SVC obstruction
Pericardium: effusion --> tamponade
Pleura: effusion

#### **HIGHLY METASTASIC**



#### **LUNG "CANNONBALL METASTASES"**

LIVER LYMPH NODES KIDNEY BRAIN



Reduced number of cases → lack for protocols and adecuate consens of management

Inespecificity in symtoms usually → late diagnosis, usually when metastasis is already present.



Not usually a viable option due to the aggresiveness and the promptly with witch it generates metastasis

THIS LEAD TO A POORER PROGNOSIS THAN ITS TESTICULAR COUNTERPART

# CONCLUSIONS

- ✓ Germ cell tumors and most importantly primary mediastinal choriocarcinoma is an infrequent entity, which leads to a late diagnosis and a lack of established treatment.
- ✓ This, together with its aggressive behavior and frequent presence of metastasis at diagnosis, makes it a highly lethal condition.
- ✓ Knowledge of their characteristics and suspicion in pediatric patients with hormonal alterations play a fundamental role for a timely treatment and a better prognosis.