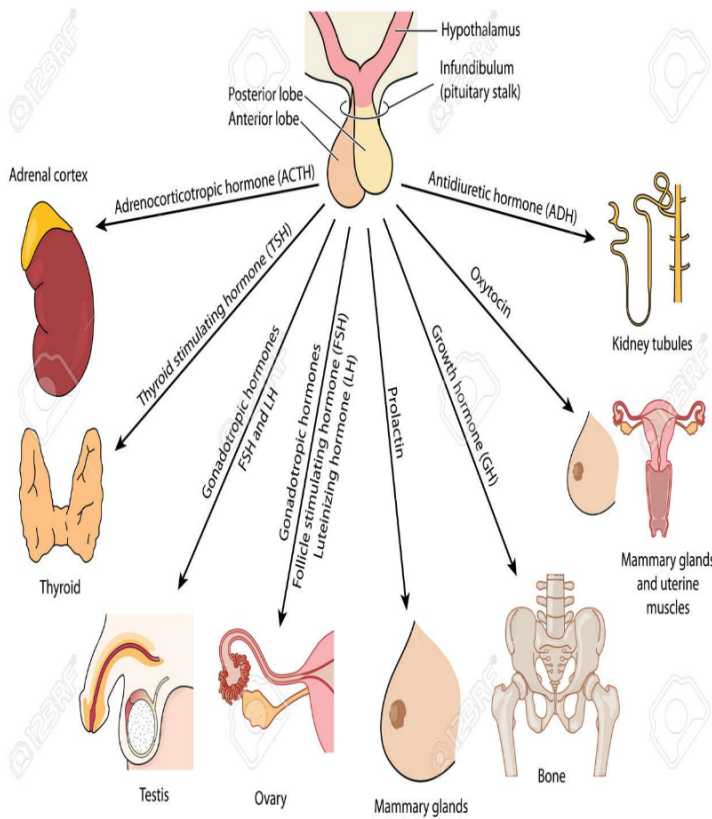
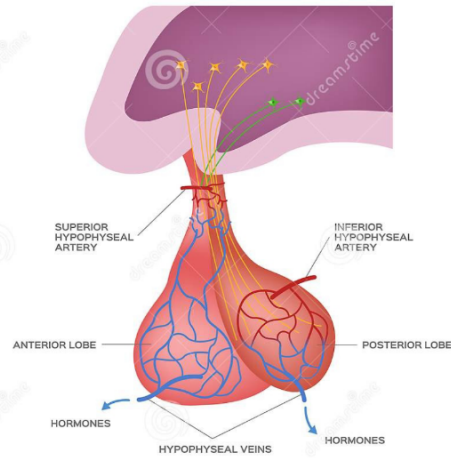
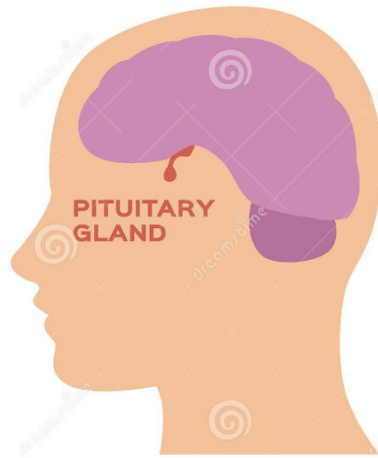


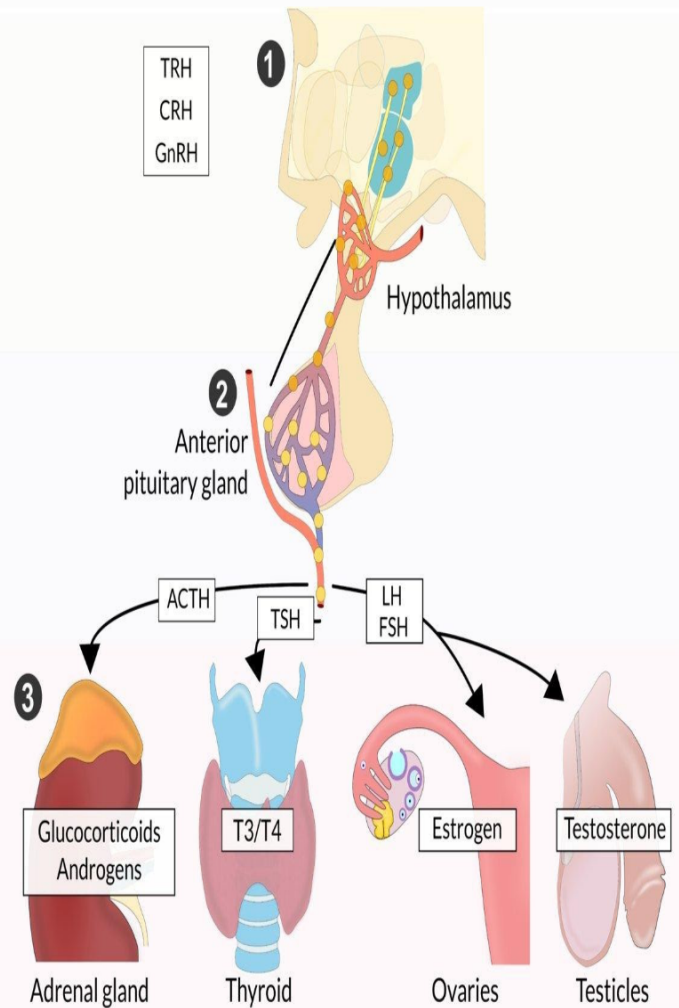
# Diseases of the Pituitary Gland :-



Releasing hormones

Anterior pituitary hormones

Target organ



## (1) \* Pituitary adenomas :-

- 10% of intracranial neoplasms
- almost all are benign.

micro  $\leq$  10 mm    macro  $>$  10 mm

## \* Clinical features :-

1) Hormonal

⇒ 1) hormonal :-

- Prolactin → hyperprolactinemia
- GH → Acromegaly or gigantism.
- ACTH → Cushing disease
- TSH → hyperthyroidism.

- 2) hypopituitarism → Compression of hypothalamic-pituitary stalk (hypogonadotropic, hypogonadism)
- 3) mass effect → headache  
Visual defects - Bitemporal hemianopia

### \* Diagnosis:-

- MRI study of choice
- pituitary hormones level.

### \* treatment:-

- 1) transphenoidal surgery → indicated in most pts  
except pts with prolactinomas
- 2) Radiation & medical therapy adjuncts in most pts.

### (2) ⇒ hyperprolactinemia

#### \* Causes:-

- Prolactinoma → M.C.C  
most common type of pituitary adenoma
- drugs → Psychiatric medications, H<sub>2</sub> blockers, estrogen  
verapamil, metoclopramide
- pregnancy.

Hypothalamus



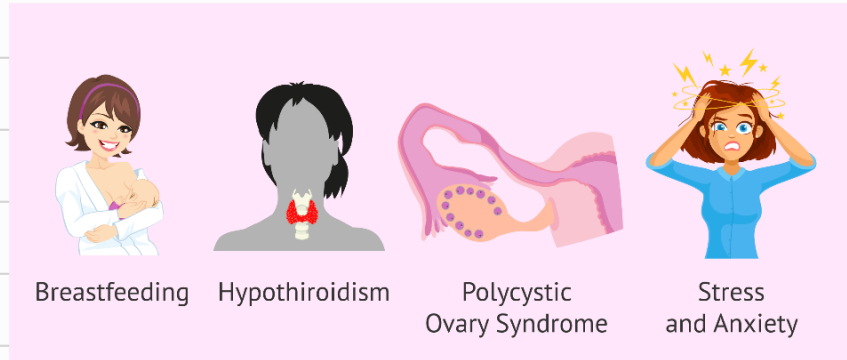
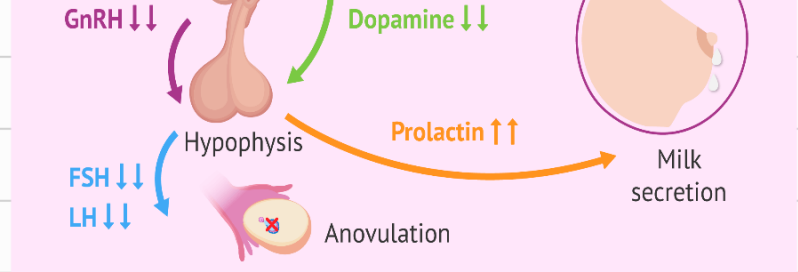
- Renal failure

- Sellar mass lesions

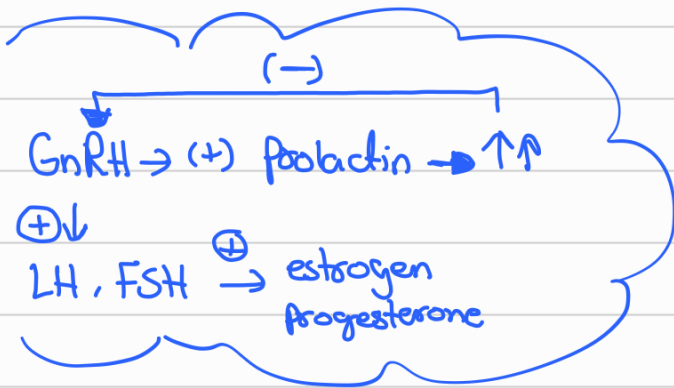
- hypothyroidism

- Idiopathic.

### \* Clinical Features:-



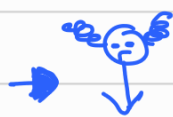
- prolactin →  Perimenopausal → menstrual irregularities



oligomenorrhea or amenorrhea  
 anovulation  
 infertility  
 ↓ libido  
 Vaginal dryness  
 risk of osteoporosis  
 dyspareunia

Postmenopausal →

Pituitary signs & symptoms  
 "visual field defects & headache"



- hypogonadism, ↓ libido, infertility  
impotence.

- galactorrhea or gynecomastia 'uncommon'

➔ **Diagnosis** :- \* ↑ serum prolactin

\* order pregnancy test, TSH level

\* CT, MRI to any mass lesions.

→ treatment :-

- treat the underlying cause

- if prolactinoma → symptomatic → dopamine agonist  
"Cabergoline, bromocriptine"  
→ surgical intervention if symptoms progress despite appropriate medical treatment.

(3) \* Acromegaly :-

- broadening of the skeleton
- GH-secreting adenoma 10%

\* Clinical features:

- Growth promotion
- Soft tissue & skeleton overgrowth
- Coarsening of facial features
- abnormally large hand & foot size
- Organomegaly
- hypertrophic cardiomyopathy ⇒ M.I.C. of death.
- Arthralgia
- enlarged jaw

→ metabolic disturbances

hyperhidrosis

Glucose intolerance & DM in 25% of pts.

→ Parasellar manifestations.

- headache

Superior → bitemporal hemianopia

lateral → cavernous sinus compression



inferior  $\Rightarrow$  sphenoid sinus inversion

HTN, sleep apnea.

## Diagnosis

$\rightarrow$  IGF-1 / Somatomedin C  $\uparrow$

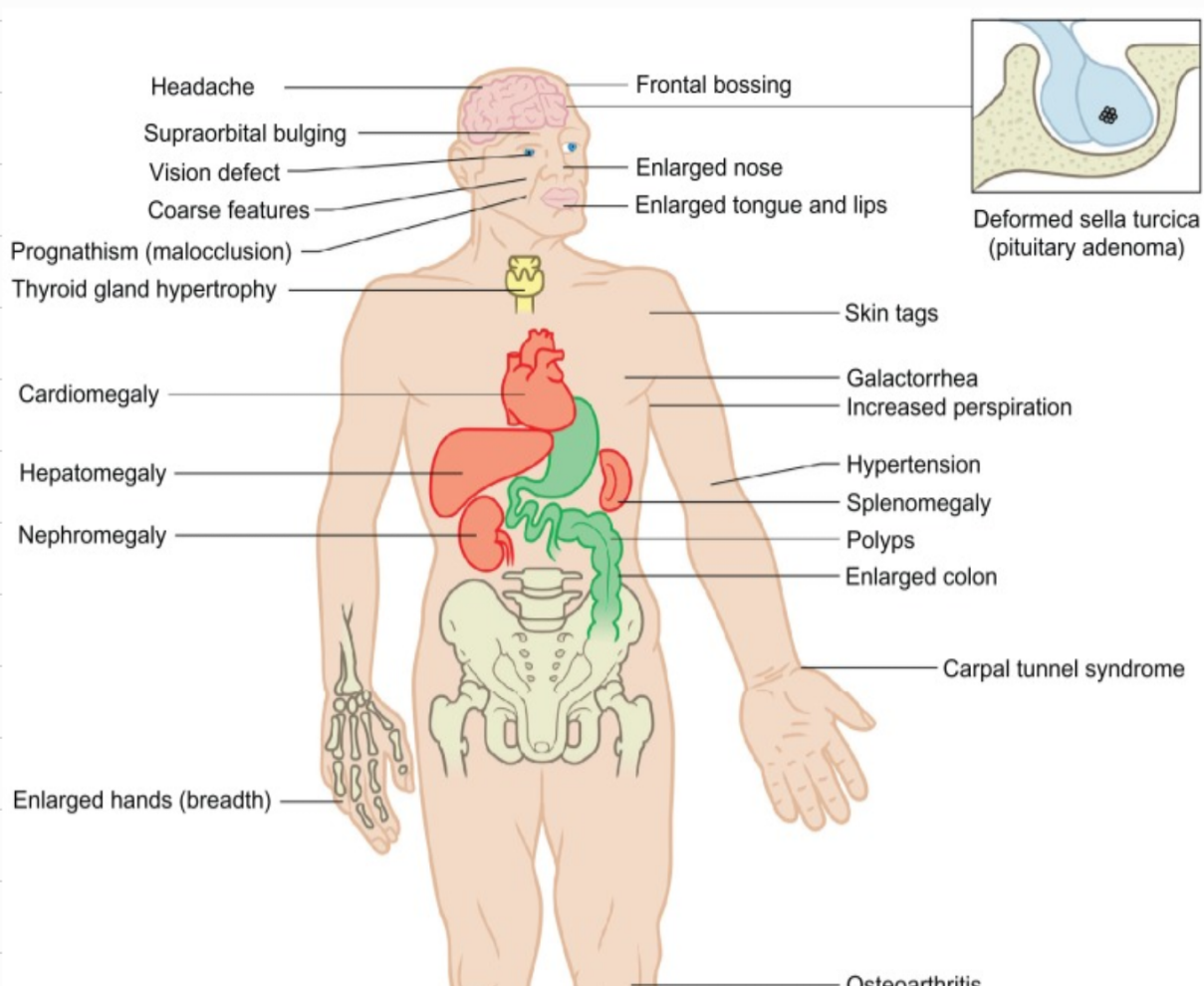
$\rightarrow$  oral glucose suppression test  $\rightarrow$  still  $\uparrow$  IGH

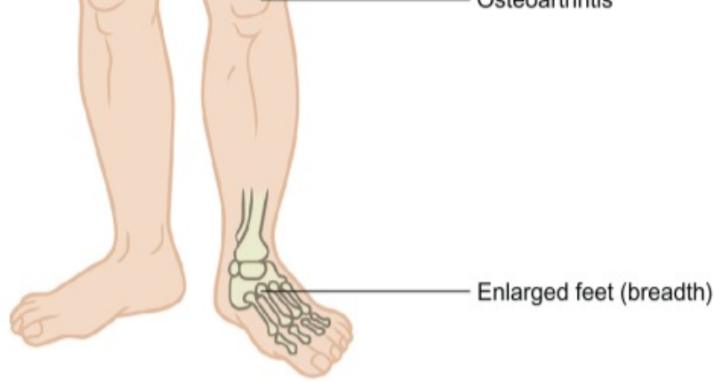
$\rightarrow$  MRI of the pituitary  $\Rightarrow$  elevation in serum glucose, triglycerides, phosphate levels.

## Treatment:-

- transphenoidal resection of pituitary adenoma

. Radiation therapy if IGF-1 elevated after surgery





## (4) Hypopituitarism:-

↓ or absent all hormones of Anterior pituitary  
or some

↓ First in LH, FSH, GH before ACTH, TSH

\* Causes:-

- hypothalamic or pituitary tumor
- Radiation therapy
- head trauma
- surgery
- infiltrative process
  - Sarcoidosis, hemochromatosis
- Cavernous sinus thrombosis

\* Clinical features:-

↓ GH → growth failure, ↑ LDL, ↑ Risk for  $\heartsuit$  disease

↓ Prolactin → failure to lactate.

↓ ACTH → adrenal insufficiency

↓ TSH → hypothyroidism.

↓ LH, FSH → infertility, amenorrhoea, loss of secondary sex  
↓ libido

↓ ADH → diabetes insipidus

↓ MSH → ↓ skin & hair pigmentation

\*diagnosis → low levels of target hormones  
MRI of the brain

\*treatment → Replacement of hormones

