

الطب والجراحة  
لجنة

# The Adrenal Gland

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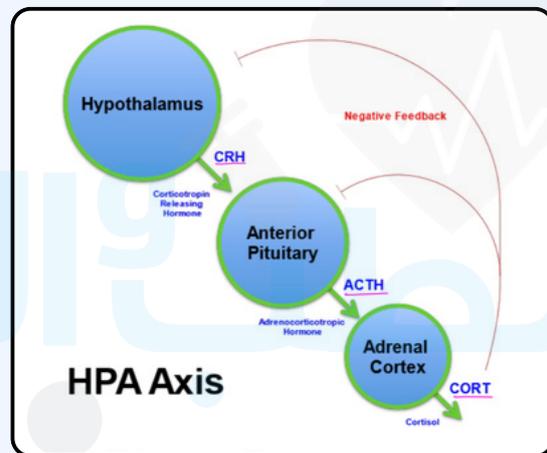
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# THE ADRENAL GLANDS

- Adrenal cortex:
  - Zona glomerulosa... Mineralocorticoids,
  - Zona fasciculata.....Glucocorticoids
  - Zona reticularis.....Sex Hormones
- Adrenal medulla : Adrenaline, Noradrenaline, Dopamine

## Hypothalamic – pituitary – adrenal axis



Adrenal mass: → functioning → secrete hormone  
 → non functioning → doesn't secrete hormones

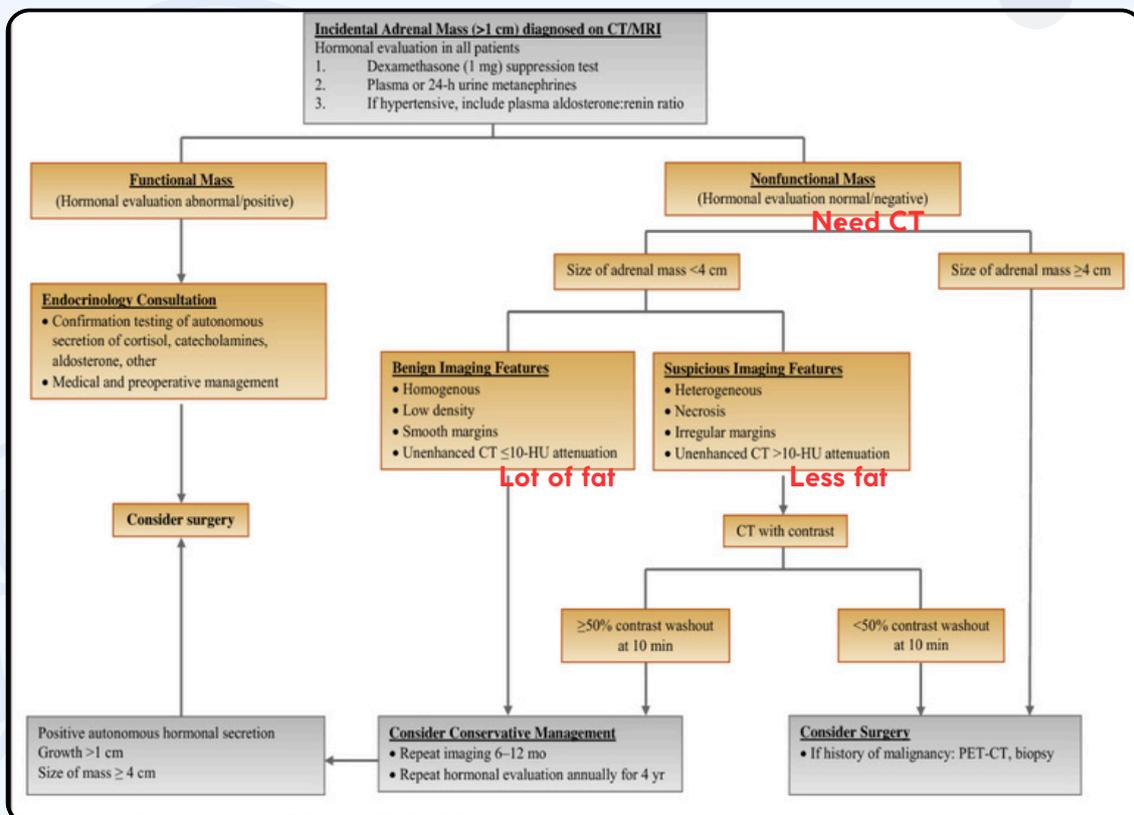
Is called ✓

## Adrenal incidentaloma (non-functional )

Non- functional adrenal mass More than 1 cm / Asymptomatic /

Found incidentally

Diagnosed by radiological imaging



## Functional Adrenal Abnormalities

- Benign or malignant tumors or hyperplasia **Increase size of cells**
- Cortex : Cortical tumors :
- Cortisone secreting tumors- Cushing's Syndrome
- Aldosterone secreting tumors- Conn's Syndrome
- Sex hormone secreting tumors- Virilisation or Feminization.

## Diffuse Hyperplasia

- Primary or a consequence of stimulation by trophic hormones leading to hypercortisism , Conn's disease or Adrenogenital syndrome

## Medulla

- Tumors secreting adrenaline/nor-adrenaline( Phaeochromocytoma )

## Cushing's Syndrome

=Primary adrenal disease:

- Adenoma
- Carcinoma
- Primary adrenal hyperplasia " ACTH independent

=Secondary adrenal disease: **Stimulation from out of the gland**

- **Mainly** Primary pituitary micro-adenoma **ACTH dependent**
- Non pituitary source " Ectopic ACTH syndrome "

## Cushing's Syndrome

- Definition: Excess circulating cortisol that occurs as a result of endogenous steroid hyper secretion, due to:  
ACTH dependent or ACTH\_ independent disease Or exogenous steroid medication.

## ACTH-Dependent

- most common** 1. Pituitary microadenoma.
2. Ectopic ACTH secretion:
  - a) Small cell carcinoma
  - b) Fore gut carcinoid
  - c) Ectopic CRH Syndrome:  
Medullary thyroid tumor/ Pancreatic neuro-endocrine tumors

## ACTH Independent

- Adrenocortical Adenoma **Benign**
- Bilateral nodular hyperplasia
- Adrenal carcinoma **Malignant**

## Cushing's Syndrome

- Physiological and bodily changes caused by excess of circulating cortisol:
- Commonest cause is iatrogenic: administration of steroids for the treatment of other diseases.

## Action of glucocorticoids

- Glucose metabolism
- Peripheral glucose utilization
- Lipid metabolism
- Cells of immune system
- Mediators of inflammation
- Bone and minerals metabolism
- Soft tissue and skeletal growth
- Fluid and electrolytes homeostasis
- CNS system

## Clinical presentation

- Obesity
- Loss of connective tissue
- Hirsutism and Virilism
- Muscle weakness
- Osteoporosis
- Hypertension
- Glucose intolerance
- Psychological changes



## Ectopic ACTH Secretion

- Rapid evolution of the Cushing's;
- Symptoms of the primary disease:
  - Small cell carcinoma of the lung
  - Carcinoid
  - Medullary Ca of Thyroid
  - Other primary carcinomas

### Investigations:

- 1 : Biochemical diagnosis
  - Persistent increase in cortisol concentration.
  - Cortisol suppression by dexamethasone **In Cushing patients cortisol doesn't decrease**
  - Resistancy to insulin administration
- 2 : Establishment of the cause
  - Low ACTH = Adrenal disease
  - High ACTH = Extra- adrenal cause **As small lung cancer**

### Anatomical details **مهم**

- Pituitary: Skull X ray/ CT /MRI
- Adrenals: US/ CT/ MRI
- Scintigraphy : cholesterol scan // NP59 scan **For Cushing patients**
- Search for ectopic ACTH source : CT chest/ Angiography

### Plan of Management

**Adrenal disease need post and pre Op care due to hormones which can't stop it suddenly**

- Pituitary adenoma : Microadenectomy
- Hyperplasia : Bilateral adrenalectomy
- Solitary adenoma: Unilateral adrenalectomy

### Adrenocortical Carcinoma

- Rare
- Any age 4-5th decades
- 60% : no important secretory function
- Benign or Malignant ? Pain / Weight loss/ Weakness/ Fever
- Functional tumors present depending on their type of secretion .

## Treatment

- When possible Surgical resection
- Radiotherapy
- Chemotherapy

## Aldosteronism <Conn's Syndrome>

• Primary due to :

1) tumor( Adenoma ) 2) nodularity 3) hyperplasia

-Secondary due to:Excess stimulation by Angiotensin

Commonest cause is :

“Aldosterone producing Adenoma “

Incidence: Females more than males

30–60 years of age

1% of patients investigated for hypertension

## Pathophysiology

• Aldosterone :

Promotes sodium absorption / Promotes water retention

Increase potassium secretion **Hypokalemia not hyponatremia**

## Clinical features :

Clinical suspicions should be raised when Hypertension occur with hypokalemia.

•Moderate to severe hypertension

•Hypokalemia

•Muscle weakness

•Malaise

•polydipsia

## Investigations

• Blood : Hypokalemia

Plasma aldosterone

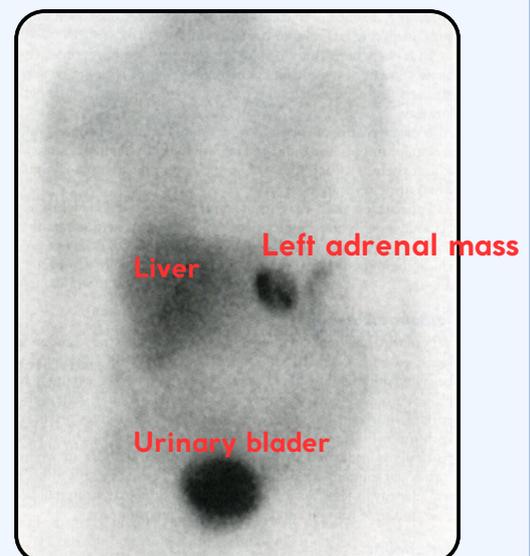
• Urine : Increase urinary potassium

• Imaging : US/ CT/ MRI

Iodocholesrerol isotope / scan / Adrenal vein sampling

**Specific for Conn's syndrome**

**From (Rt and Lt) vein**



## Treatment

- Spironolactone **K<sup>+</sup> sparing**
- Adrenalectomy

## Phaeochromocytoma :

### Phaeochromocytoma

### Neuroblastoma

### Paraganglioma

### Ganglioneuroma

Are derived from the neural crest → Arise more than tumor from it

- 90% ---solitary – adrenal
- 5 –10% bilateral
- 10%---Exrta-adrenal
- 0.1% of patients investigated for hypertension
- Average size is 5 cm
- Discovered early because of catecholamines effects
- 10% are malignant
- Mostly secretes adrenaline

Pheochromocytoma---> Disease of 10%:  
-multible  
-bilateral  
-extraadrenal  
-malignant

## Symptomatology

- Palpitation
- Hypertension
- Sweating and pallor
- Anxiety
- Chest pain & weakness 50%
- Attacks often occur spontaneously but may be precipitated by vigorous exercise, Alcohol, tobacco and drugs : Anesthesia, phenothiazines & tricyclic antidepressants.

## Clinical associations

- \* Multiple endocrine neoplasia type 2 Phaeo , medullary thyroid ca, hyperparathyroidism
- \* Neurofibromatosis . 10% of patients with neurofibromatosis may develop phaeochromocytoma

## Investigations

Adrenaline is broken in urin

• A- 24 hours urinary vanil mandilic acid (VMA) 60% sensitive.

• Urinary catecholamines . 90% sensitive

• Localization:

CT scan

MRI

MIBG , isotope scan

## Managemant:

=Adrenalectomy

• Preoperative management

• Operative management **Can't stop hormone abruptly**

• Post operative management

**You should control HTN**

إِنَّ الْمُؤْمِنَ الْعَارِفَ بِاللَّهِ حَقًّا لَا يَزَالُ يَطْلُبُ مِنَ اللَّهِ حَاجَتَهُ، وَلَوْ كَانَتْ  
السُّنَنُ الطَّبِيعِيَّةُ كُلُّهَا تُعْبَرُ عَنْ اسْتِحَالَةِ الْوُقُوعِ!  
-د. فريد الأنصاري -رحمه الله

