

الطب والجراحة
لجنة

The Adrenal Gland

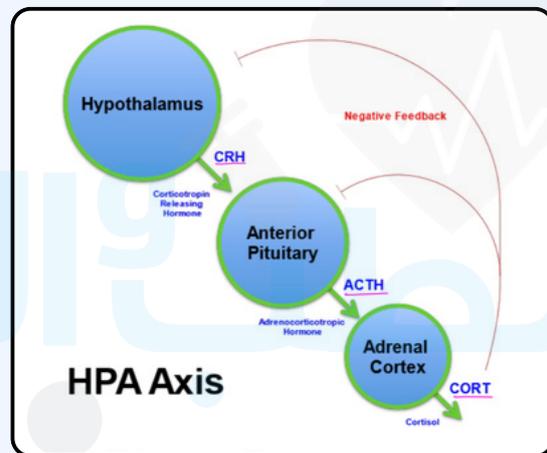
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THE ADRENAL GLANDS

- 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

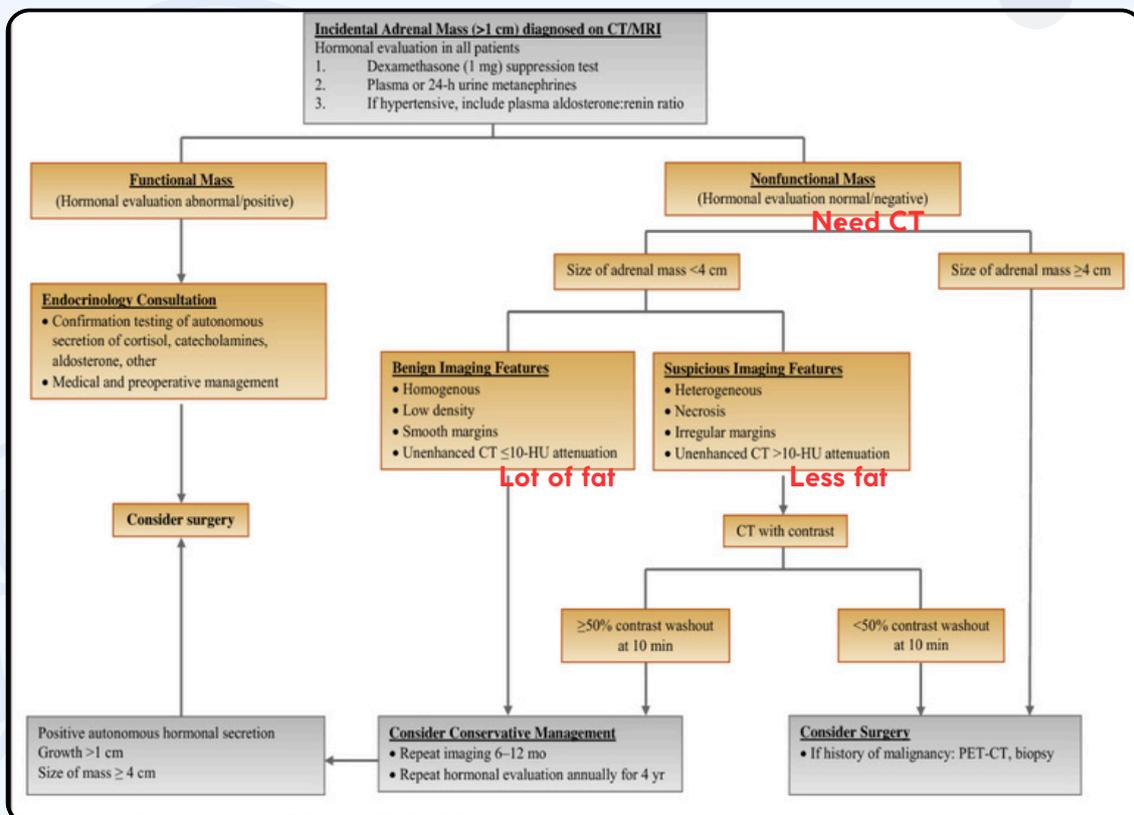
Is called ✓

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 :

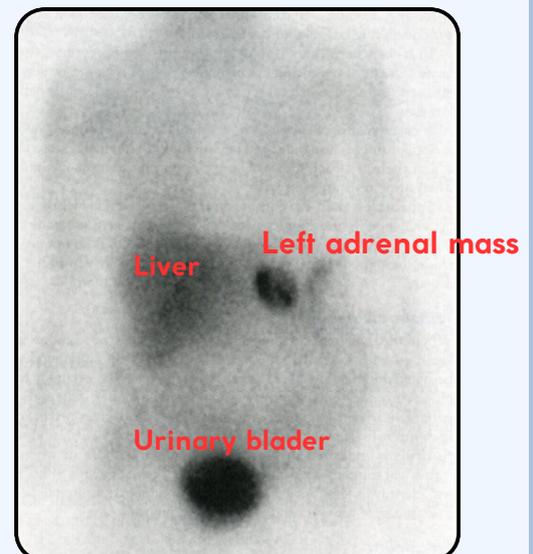
Clinical suspicions should be raised when Hypertension occur with hypokalemia.

- Moderate to severe hypertension
- Hypokalemia
- Muscle weakness
- Malaise
- polydipsia

Investigations

- Blood : Hypokalemia
Plasma aldosterone
- Urine : Increase urinary potassium
- Imaging : US/ CT/ MRI

Iodocholesterol isotope / scan / Adrenal vein sampling **From (Rt and Lt) vein**
Specific for Conn's syndrome



Treatment

- Spironolactone **K⁺ 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

