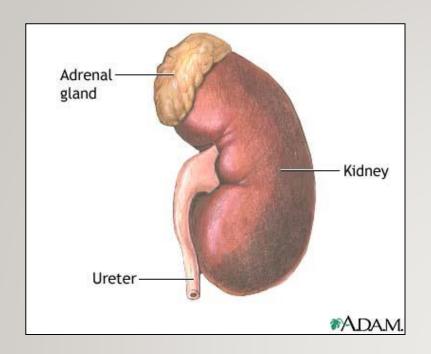
## BIOCHEMISTRY OF CUSHING SYNDROME

**ENDOCRINE BLOCK** 

## **OBJECTIVES**

- To identify physiological and biochemical characteristics of Cortisol.
- To understand the diagnostic algorithm for Cushing's Syndrome.
- To understand the interpretation of laboratory and radiological investigations for diagnosis of Cushing's Syndrome.

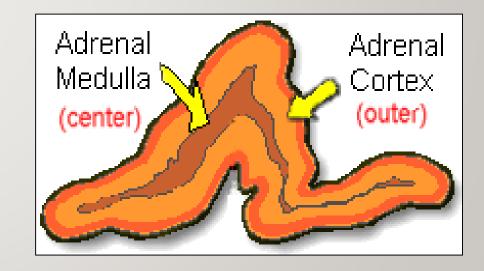


#### **Anatomically:**

The adrenal gland is situated on the anteriosuperior aspect of the kidney

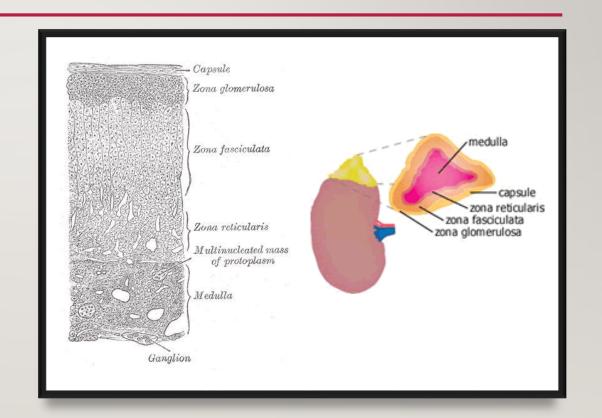
#### Histologically:

The adrenal gland consists of two distinct tissues of different embryological origin, the outer cortex and inner medulla.



## THE ADRENAL CORTEX

- Based on cell type and function, the adrenal cortex comprises three zones:
  - Zona <u>G</u>lumerulosa (the outmost zone) → aldosterone, the principle mineralocorticoid.
  - Zona <u>F</u>asciculata →
    glococorticoids, mainly cortisol
    (95%).
  - Zona Reticularis  $\rightarrow$  sex hormones.



#### Cholesterol (27C) STEROID Pregnenolone (21C) 3-β-Hydroxysteroid dehydrogenase HORMONE **Progesterone (21C) SYNTHESIS** 17-α-Hydroxylase 17-α-Hydroxyprogesterone (21C) 21-α-Hydroxylase 11-Deoxycorticosterone (21C) **Androstenedione (19C)** Peripheral tissues 11-Deoxycortisol (21C) Testosterone (19C) 11- β -Hydroxylase Corticosterone **Estradiol (18C)** Cortisol (21C) Aldosterone (21C)

## HYPOTHALAMIC-PITUITARY-ADRENAL (HPA) AXIS

• The <u>hypothalamus</u> secretes corticotropin-releasing hormone (CRH) which stimulates the <u>anterior pituitary gland</u> to synthesis and release ACTH.

• ACTH acts on the zona fasiculata cells  $\rightarrow$  release of glucocorticoids (*Cortisol*).

## GLUCOCORTICOID FUNCTIONS

- Glucocorticoids have widespread metabolic effects on carbohydrate, fat and protein metabolism.
- Upon binding to its target, <u>CORTISOL</u> enhances metabolism in several ways:
  - In the liver, Cortisol is an insulin antagonist and has a weak mineralocorticoid action:
    - $\uparrow \uparrow$  Gluconeogenesis  $\rightarrow$  production of glucose from newly-released amino acids and lipids
    - ↑↑ Amino acid uptake and degradation
    - \hftarrow
       \hftarrow
       Ketogenesis.
  - In the adipose tissue: Cortisol  $\rightarrow \uparrow \uparrow$  Lipolysis through breakdown of fat.
  - In the muscles: Cortisol  $\rightarrow \uparrow \uparrow$  proteolysis and amino acid release.
  - Conserving glucose: by inhibiting uptake into muscle and fat cells.

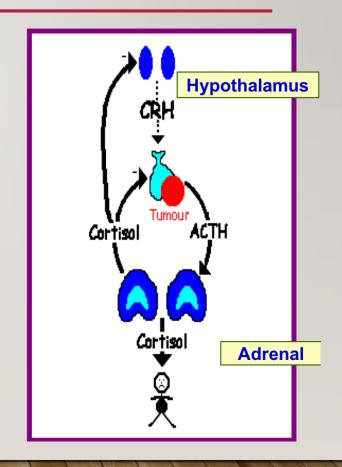
## REGULATION OF ACTH AND CORTISOL SECRETION

#### I. Negative feedback control:

- ACTH release from the anterior pituitary is stimulated by hypothalamic secretion of corticotrophin releasing hormone (CRH).
- CRH  $\rightarrow \uparrow$  ACTH  $\rightarrow \uparrow$  [Cortisol]
- ↑[Cortisol] or synthetic steroid suppress CRH & ACTH secretion

#### 2. Stress:

(e.g. major surgery, emotional stress) Stress  $\rightarrow \uparrow \uparrow$  CRH & ACTH  $\rightarrow \uparrow \uparrow$  Cortisol

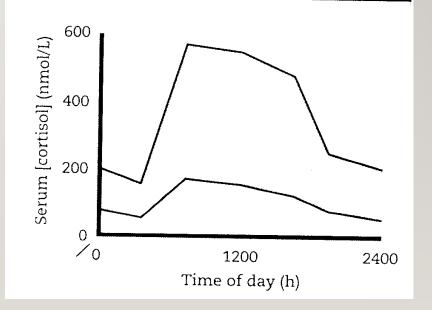


#### 3. The diurnal rhythm of serum cortisol:

- Highest Cortisol level in the morning (8 9 AM).
- Lowest Cortisol level in the late afternoon and evening (8 - 9 PM).

The diurnal rhythm of cortisol secretion; the area between the curves represents values that lie within the reference range

## DIURNAL RHYTHM OF CORTISOL SECRETION



## PLASMA CORTISOL-BINDING GLOBULIN (CBG)

- In the circulation, glucocorticoids are mainly protein-bound (about 90%), chiefly to CBG (transcortin).
  - 1 in pregnancy and with estrogen treatment (e.g. oral contraceptives).
  - $\downarrow \downarrow$  in hypoproteinemic states (e.g. nephrotic syndrome).

The biologically active fraction of cortisol in plasma is the free (unbound) component.

## CORTISOL AND ACTH MEASUREMENTS

#### Serum (Cortisol) and plasma (ACTH):

- Samples must be collected (without venous stasis) between 8 a.m. and 9 a.m. and between 10 p.m. and 12 p.m. because of the diurnal rhythm.
- Temporary  $\uparrow\uparrow$  in these hormones may be observed as a response to emotional stress.

#### **Urinary Cortisol excretion:**

- Cortisol is removed from plasma by the liver  $\rightarrow$  metabolically inactive compounds  $\rightarrow$  excreted in urine mainly as conjugated metabolites (e.g. glucuronides).
- A small amount of cortisol is excreted unchanged in the urine (UFC).
- In normal individuals:
  - Urinary free cortisol (UFC) is < 250 nmol/24 h.</li>

## CAUSES OF ELEVATED SERUM CORTISOL

#### **Increased cortisol secretion:**

- Cushing's syndrome
- Exercise
- Stress, Anxiety, Depression
- Obesity
- Alcohol abuse
- Chronic renal failure

#### **Increased CBG:**

- Congenital
- Estrogen therapy
- Pregnancy

## SYMPTOMS OF CUSHING'S SYNDROME

- Weight gain, central obesity.
- Buffalo's hump.
- Moon face —
- Excessive sweating
- Atrophy of the skin and mucous membranes
- Purple striae on the trunk and legs
- Proximal muscle weakness (hips, shoulders)
- Hirsutism
- The excess cortisol may also affect other endocrine systems  $\rightarrow \downarrow$  libido, amenorrhoea and infertility
- Patients frequently suffer various psychological disturbances ranging from euphoria to frank psychosis.

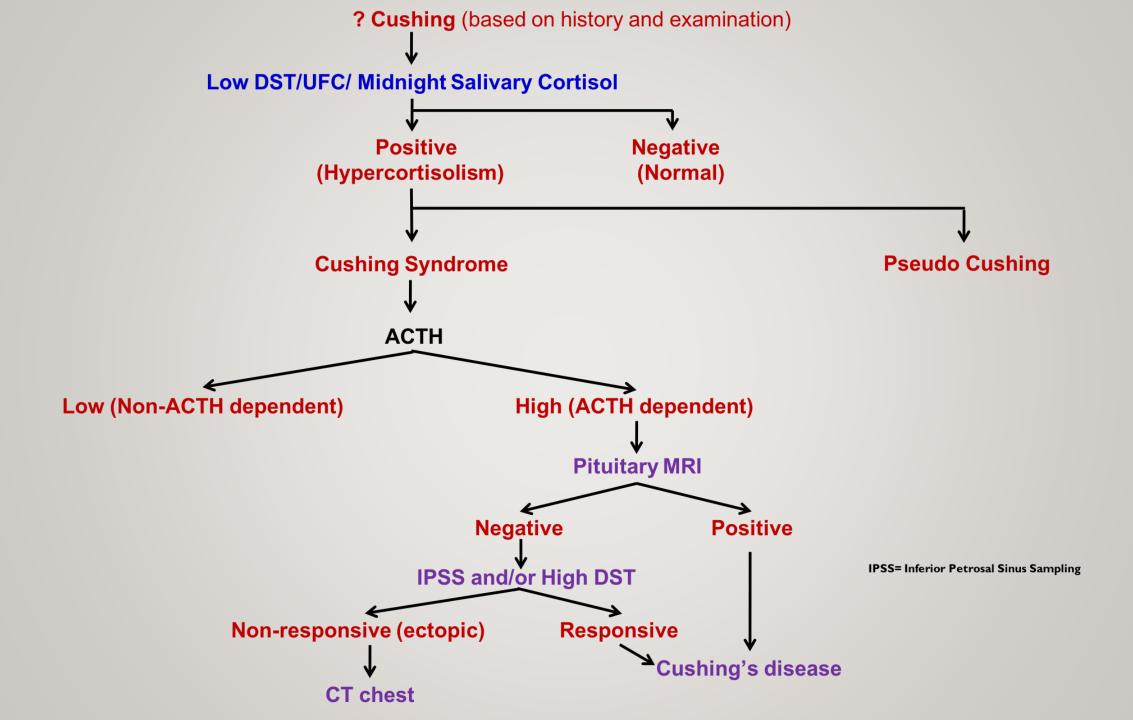


## **SIGNS**

- Loss of diurnal rhythm of cortisol and ACTH.
- Hypertension (due to the aldosterone like effects)
- Hyperglycemia or diabetes due to insulin resistance.
- Hypokalemic alkalosis
- ↑ protein metabolism.
- Impaired immunity.

## INVESTIGATIONS OF SUSPECTED ADRENOCORTICAL HYPERFUNCTION

- A. Screening and confirmatory tests: to assess the clinical diagnosis of adrenocortical hyperfunction.
- **B. Tests to determine the cause:** to ascertain:
  - (a) The site of the pathological lesion (adrenal cortex, pituitary or elsewhere?)
  - (b) The nature of the pathological lesion.



### **SCREENING TESTS**

#### I. Low-dose DST:

#### **Procedure:**

I mg dexamethasone (DXM) administered at II-I2 PM the night before attending the clinic. Serum cortisol is measured at 8-9 AM.

#### **Result:**

Cortisol < 50 nmol/L (suppression) → exclude hypercortisolnemia (Cushing Syndrome)

#### **Precautions:**

Drugs that induce hepatic microsomal enzymes (Phenobarbitone & phenytoin)  $\to \uparrow$  DXM metabolism and  $\downarrow$  DXM blood level to achieve CRH suppression (false diagnosis of Cushing)

#### 2. 24- hour urinary free cortisol:

**Result:** Cortisol < 250 nmol/day  $\rightarrow$  exclude Cushing Syndrome.

**Disadvantage:** incomplete collection of urine  $\rightarrow$  a false-negative result.

#### 3. Midnight Salivary Cortisol.

**Result:** Cortisol < 100 ng/dL  $\rightarrow$  exclude Cushing Syndrome.

## **CONFIRMATORY TESTS**

- Positive results of at least two screening tests would confirm the clinical diagnosis.
- Further investigations are required.

## TESTS USED TO DETERMINE THE CAUSE OF CUSHING'S SYNDROME

- To differentiate ACTH-dependent from ACTH-independent:
   Plasma ACTH (Diurnal rhythm)
- 2. To distinguish between ACTH-dependent causes (Pitutary Vs Lung):
  - a) High-dose DST.
  - b) Inferior Petrosal Sinus Sampling.
- 3. Radiological tests: MRI of pituitary and ultrasound or CT of adrenals.

# CAUSES OF ADRENOCORTICAL HYPERFUNCTION (CUSHING'S SYNDROME)

#### ACTH - dependent:

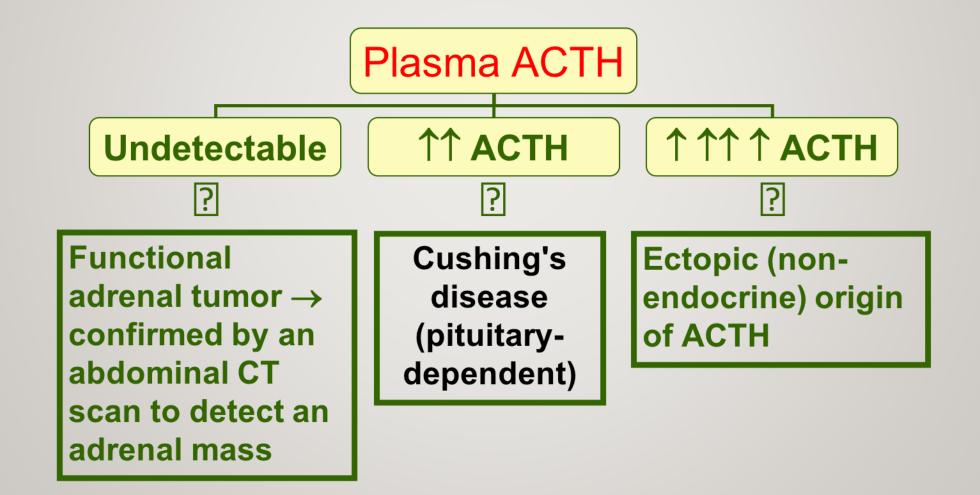
- I. ↑ Pituitary ACTH >90% (Cushing's disease).
- 2. Ectopic ACTH by neoplasms <10%.

#### ACTH - independent:

- I. Glucocorticoid therapy.
- 2. Adrenal tumor <20% (adenoma or carcinoma).

### PLASMA ACTH

It should be measured on blood specimens collected at 8-9 a.m. and 8-9 p.m.



## HIGH-DOSE DST

#### It is used to distinguish Cushing's disease from ectopic ACTH secretion.

- 2 mg DXM six-hourly for 48 hours to suppress cortisol secretion.
- Basal (pre-DXM) serum cortisol or 24-hour urine free cortisol is compared with the results at the end of the 48-hour period.
- Suppression is defined as a fall to less than 50 % of basal value.
- About 90 % of patients with Cushing's disease show suppression of cortisol output.
- In contrast, only 10% of patients with ectopic ACTH production (or with adrenal tumors) show suppression.

## OTHER BLOOD TEST

The following blood tests are commonly performed for patients suspected to have Cushing's syndrome:

- Full blood count
- Blood glucose
- Blood electrolytes and pH
- Renal function tests
- Liver function tests

## CASE STUDY

58 years old man was admitted with weight loss and respiratory distress. He had increased pigmentation and BP was 140/80.

#### Lab tests

8.6	(2.5-7 mmol/L)
144	(135-145 mmol/L)
2.0	(3.5-4.5 mmol/L)
1650	(150-550 nmol/L)
1530	(<50nmol/L)
	144 2.0 1650

## Further investigation revealed the following

DMX suppression test	Basal	after 48 h	after 48h 2.0 mg qid	
		0.5 mg qid		
Serum cortisol	1350	1420	1100	No suppression
	8 am	22.00 pm		
Plasma ACTH (ng/L)	220	180		Ref. range: 7-51

### TAKE HOME MESSAGES

- Initial screening for Cushing by 24 h urine free cortisol, low-dose dexamethasone suppression test or midnight Salivary Cortisol.
- Confirmatory tests for Cushing by getting positive results of at least two of the screening tests.
- Tests to determine the cause of Cushing: Plasma ACTH, high-dose dexamethasone suppression test, Inferior Petrosal Sinus Sampling and radiological investigations.
- ACTH-dependent Cushing: due to pituitary causes (Cushing's disease) and due to ectopic production of ACTH.
- ACTH-independent Cushing: due to adrenal adenoma or carcinoma and due to steroid therapy (iatrogenic).

### REFERENCES

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- Clinical Chemistry, Principles, Procedures, Correlations, Lippincott Williams & Wilkins, 7<sup>th</sup> edition, 2013, chapter 21, page 453-471.
- Lippincott's Illustrated Reviews: Biochemistry 6<sup>th</sup> edition, Unit III, Chapter 18, Pages 219-244.
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