بسم الله الرحمن الرحيم

ENDOCRINOLOGY

The Adrenal Gland

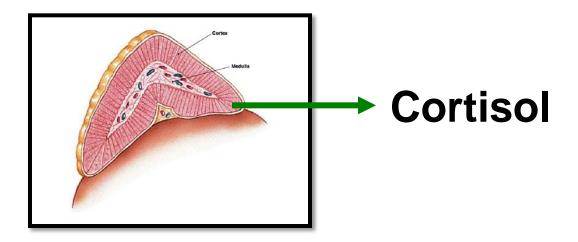
Glucocorticoids

Dr. Abeer Al-Ghumlas

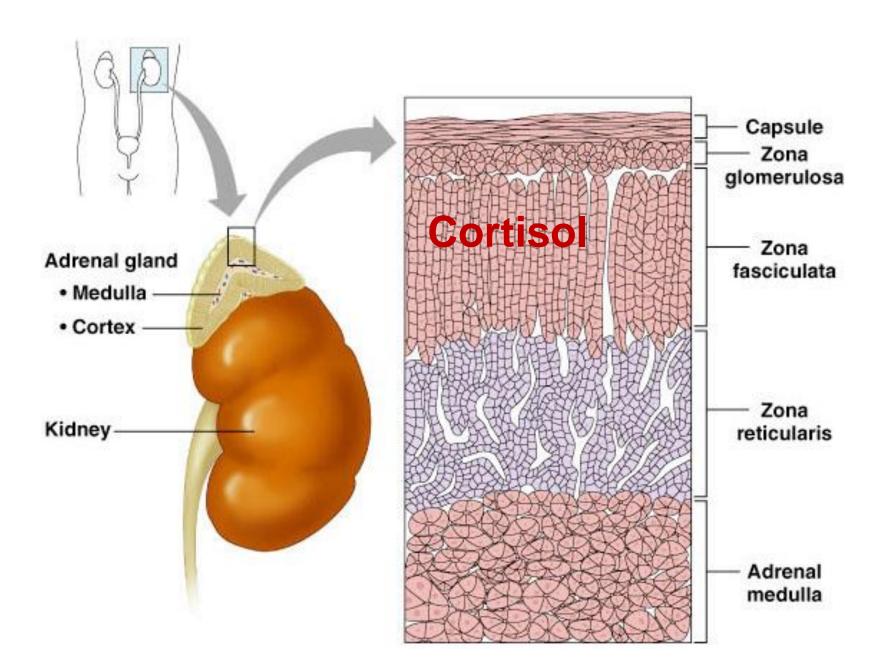
Objectives

At the end of this lecture student should be able to:

- Describe the metabolism and physiological effects of glucocorticoids.
- Describe the mechanisms that regulate secretion of glucocorticoids
- Describe the main features of the diseases caused by excess or deficiency of each of the hormones of the adrenal gland.



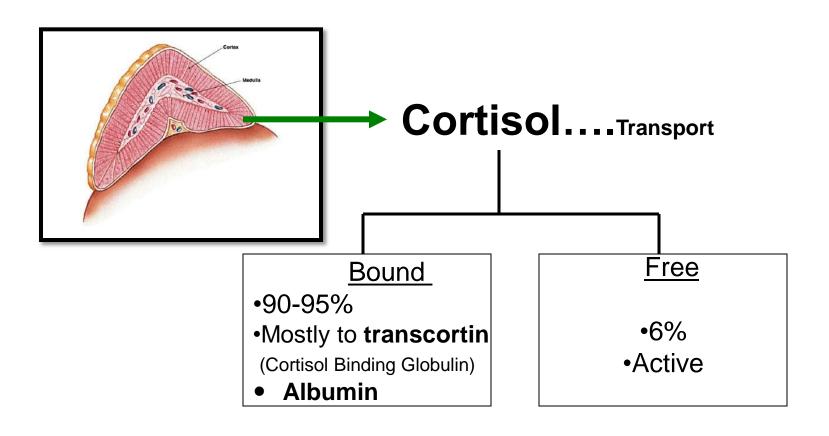
Adrenal Cortex



Glucocorticoids

Main glucocorticoids in humans:

- > Cortisol:
 - > very potent
 - Account for 95% of glucocorticoid activity
- > Corticosterone:
 - account for about 4% of total glucocorticoid activity
 - Less potent than cortisol
- ➤ Cortisol:corticosterone produced in humans in a ratio of 10:1



- Half life =60-90 minutes
- Metabolized in liver by reductases & conjugated to glucuronides and excreted via kidney
- Free cortisol is excreted into urine.

Circadian rhythm of cortisol secretion

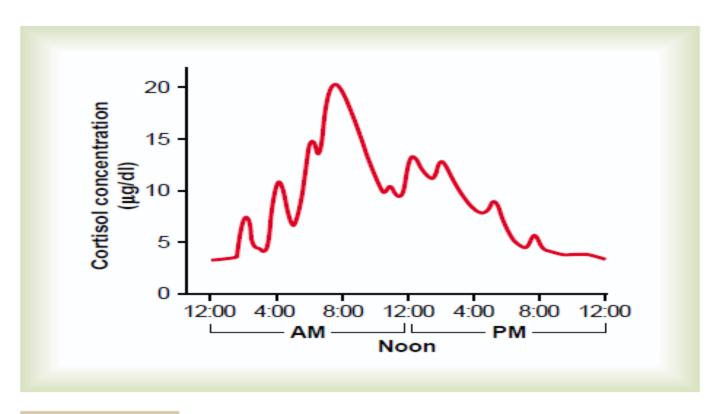


Figure 77-7

Typical pattern of cortisol concentration during the day. Note the oscillations in secretion as well as a daily secretory surge an hour or so after awaking in the morning.

Circadian rhythm of cortisol secretion

- The secretory rates of CRF, ACTH, and cortisol:
 - o high in the early morning: the plasma cortisol level ranges between a high of about 20 μ g/dl an hour before arising in the morning
 - o low in the late evening: low of about 5 μ g/dl around midnight.
- ▶ This effect results from a 24-hour cyclical alteration in the signals from the hypothalamus that cause cortisol secretion.
- When a person changes daily sleeping habits, the cycle changes correspondingly. Therefore, measurements of blood cortisol levels are meaningful only when expressed in terms of the time in the cycle at which the measurements are made.

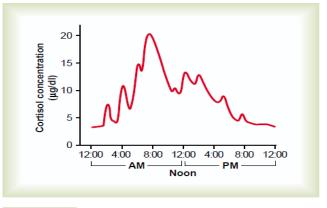


Figure 77-7

Typical pattern of cortisol concentration during the day. Note the oscillations in secretion as well as a daily secretory surge an hour or so after awaking in the morning.

Physiological actions of cortisol

A. On metabolism

On carbohydrate metabolism:

- ↑↑Glucose level in the blood:
- Stimulates gluconeogenesis and glycogen synthesis by the liver.
- ↓↓Glucose utilization by the cells.
 - "Adrenal Diabetes".



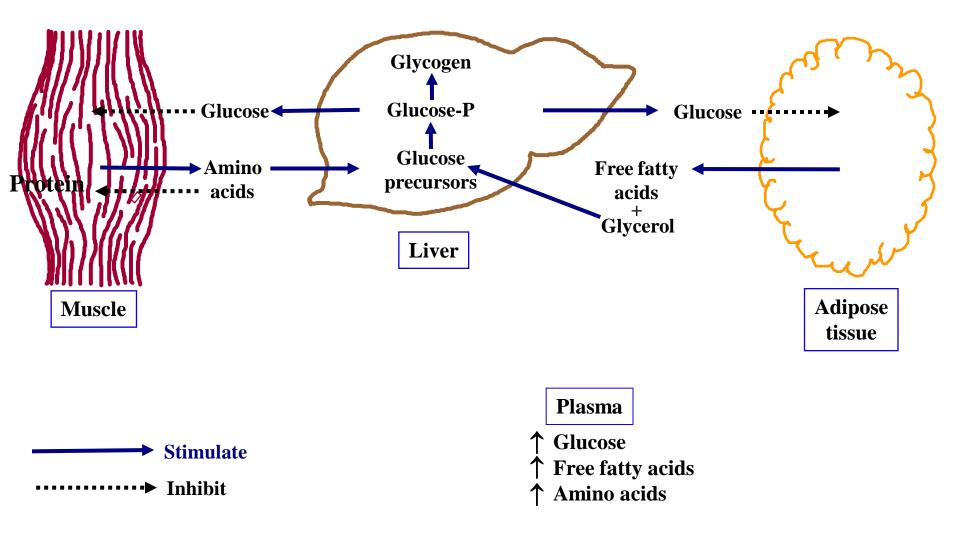
On protein metabolism:

Has a catabolic effect on proteins.

- ↓↓<u>Protein</u> stores in all body (except the liver).
- 2. ↑Liver and plasma proteins
- a.a: ↑↑ Amino acid level in the blood.
 - -↓ a.a transport into extra hepatic cells
 - †a.a transport into hepatic cells

On Fat metabolism (lipolytic):

- Mobilization of fatty acid from adipose tissue
- ↑↑ The concentration of free fatty acids in the blood.
- 3. ↑↑ Their utilization for energy.



2. Cortisol is important in resisting stress and inflammation.

On Stress:

- Stress include (trauma, infection, surgery, any debilitating disease, increase heat or cold).
- Cortisol causes rapid mobilization of amino acids and FFA from their cellular stores, making them immediately available both for energy & synthesis of other compounds, including glucose, needed by the different tissues in the body.
- □ ↑ BP,↑ glycogen, prevents stress induced reaction from becoming excessive.

Glucocorticoids and Stress:

- Without Glucocorticoids, the body cannot cope with even mild stressors
- Fat & glucose metabolism (Redistribution of Fat, increase of glucose)
- Maintenance of the vascular response to norepinephrine
- Effects on CNS

3. On inflammation

- Damage to the tissues by trauma/infection almost always leads to inflammation.
- Inflammation can be more damaging than the trauma or disease itself.
- Cortisol has anti-inflammatory effects....How?
- Cause stabilization of the intracellular lysozomal membranes → more difficult for these membranes to rupture → less release of proteolytic enzymes that cause inflammation.
- Reduces all aspects of the inflammatory process.

3. On inflammation:

Anti-inflammatory:

Reduces all aspects of the inflammatory process.

- (1)Block the early stages of the inflammation process before inflammation even begin
- (2) If inflammation begun: It cause rapid resolution of the inflammation and increase rapidity of healing
- (3) Resolution of inflammation

Anti-inflammatory:

- Stabilizes lysosomal membrane.
- Reduces degree of vasodilatation.
- Decreases permeability of capillaries.
- Decreases migration of white blood cells.
- Suppresses immune system.

4. Cortisol blocks the inflammatory response to Allergic reactions

Anaphylactic shock



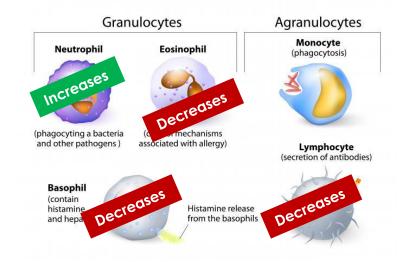
Cortisol prevent shock and death

5. On blood cells:

- Increases RBC count, platelets, neutrophils
- Decreases lymphocyte, eosinophils, basophils count
- Suppresses lymphoid tissue systemically therefore decrease T cell and antibody production decreasing immunity.
- ▶ This effect is useful in transplantation surgery in reducing organ rejection.



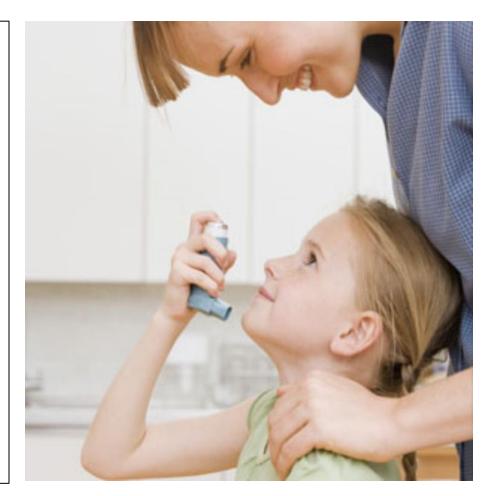
WHITE BLOOD CELL



6. anti-allergic effects

(In pharmacological doses):

- It decreases fibroblastic activity and local swelling
- phospholipase A2
- Stabalizes lysosomal membrane
- Inhibits <u>collagenase</u> from breaking down proteins
- Inhibits histamine release (antiallergic)



7. Effect on circulation

Excrete water load:

Cortisol levels vary with water intake

Mineral metabolism:

Cortisol has mineralcorticoid effect:

Na+ reabsorption and K+ secretion

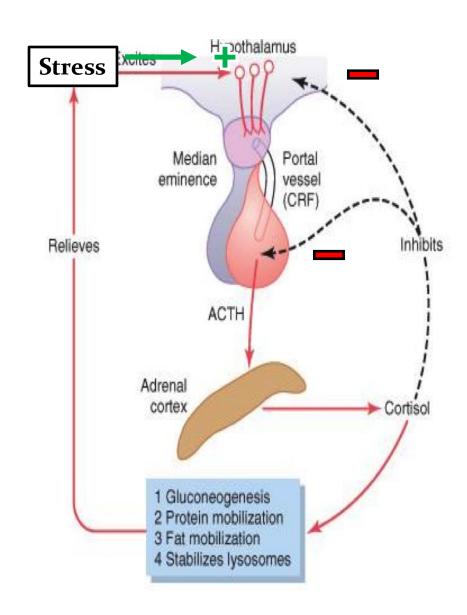
Not as potent as aldosterone.

- ➤ Maintains body fluid volumes & vascular integrity
- ➤ BP regulation & cardiovascular function: Sensitizes arterioles to action of noradrenaline (Permissive effect).
- Decreased capillary permeability
- Maintains normal renal function

Regulation of Cortisol Secretion

 Stress stimulates CRH secretion by the hypothalamus.

Cortisol has a direct negative feedback effect on both the hypothalamus and ant. pituitary.



Abnormalities

(Increase secretion of corticosteroid)



Cushing's Syndrome

(Decrease secretion of glucocorticoids and mineralocorticoids)





Addison's disease:



- Note the generalised skin pigmentation (in a Caucasion patient) but especially the deposition in the palmer skin creases, nails and gums.

- She was treated many years ago for pulmonary TB. What are the other causes of this condition?

(Addison's disease)

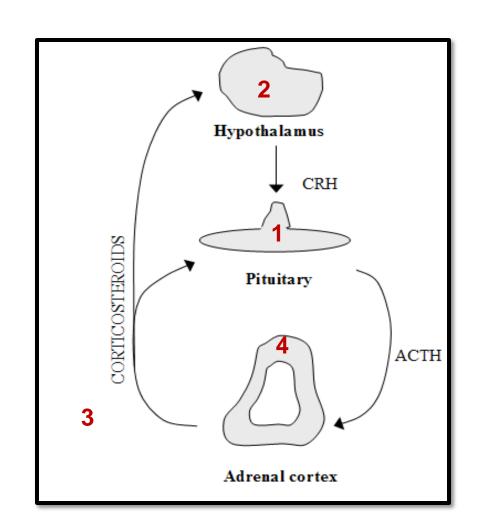
Cushing's Syndrome

Cushing syndrome: (Hypercortisolism)

Causes and types:

- (1) adenomas of the anterior pituitary→↑ ACTH.
- (2) abnormal function of the *hypothalamus* $\rightarrow \uparrow$ CRH.
- (3) "ectopic secretion" of ACTH by a tumor elsewhere in the body, such as an abdominal carcinoma.
- (4) adenomas of the adrenal cortex.

When Cushing's syndrome is secondary to \(\bar{\}\) ACTH by the anterior pituitary = Cushing's disease.



(5) Cushing's syndrome may occur when large amounts of glucocorticoids are administered over prolonged periods for therapeutic purposes.

e.g. patients with chronic inflammation associated with diseases such as rheumatoid arthritis.



Cushing's Syndrome

On carbohydrate metabolism:

- ↑ blood <u>glucose</u> level
- (↑ gluconeogenesis and
- ↓ glucose utilization by the tissues).

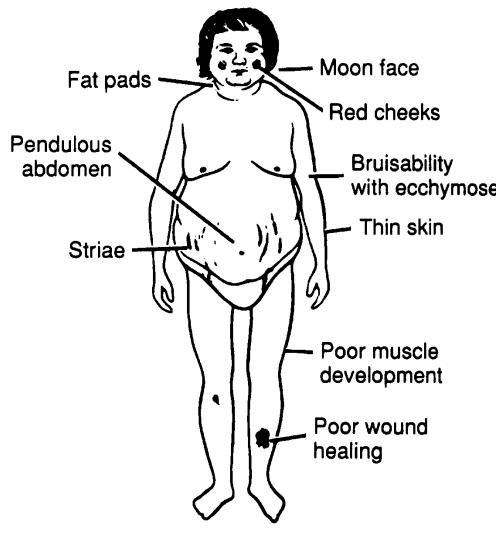
Abnormal fat redistribution

- mobilization of <u>fat</u> from the lower part of the body, with concomitant extra deposition of fat in the thoracic and upper abdominal regions, giving rise to a buffalo torso (truncal obesity).
- The appearance of the face described as a "moon face"

On protein metabolism:

- ↓ tissue proteins almost everywhere in the body (except liver).
- Protein loss from the muscles in particular causes severe weakness.
- protein collagen fibers in the s.c.(loss of CT)
- Thining of the skin
- severely ↓ protein deposition in bones → severe osteoporosis.
- suppressed immune system.

•80% of patients have hypertension (because of the mineralocorticoid effects of cortisol).



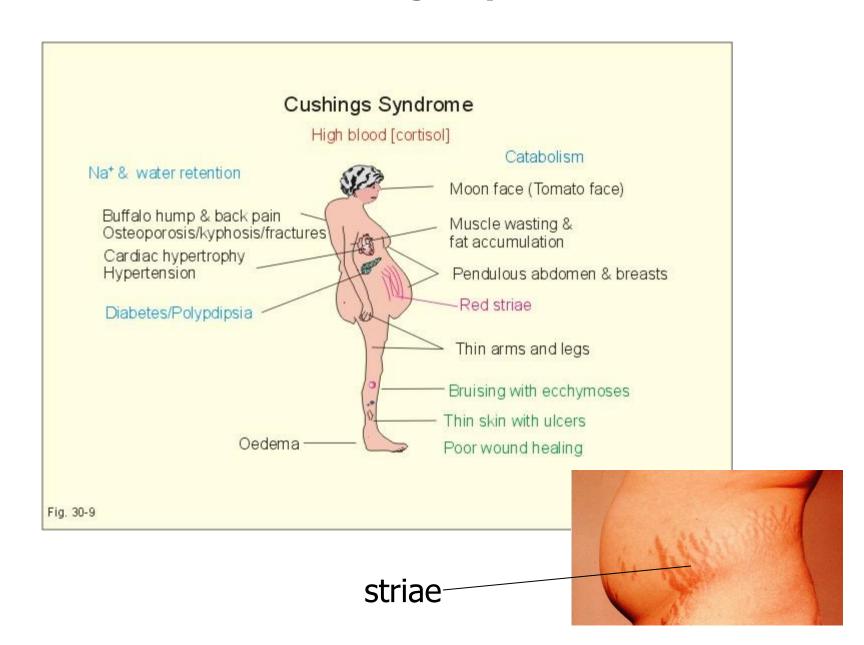
CUSHING'S SYNDROME

Cushing's Syndrome

Many people with excess cortisol secretion develop a peculiar type of obesity, with excess deposition of fat in the chest and head regions of the body, giving a buffalo-like torso and a rounded face, a "moonface".



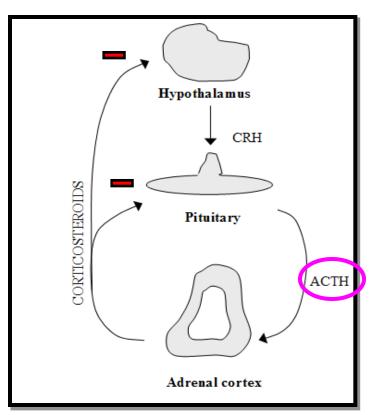
Features of Cushing's Syndrome



How to differentiate between ACTH-dependent and ACTH-Independent Cushing's syndrome?

By administrating large doses of cortisol (dexamethasone).

In patients with ↑ ACTH → no suppression of ACTH secretion.



patients with primary adrenal overproduction of cortisol (ACTH-independent) $\rightarrow \downarrow$ levels of ACTH.

Abnormalities

(Increase secretion of corticosteroid)



Cushing's Syndrome

(Decrease secretion of glucocorticoids and mineralocorticoids)





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(Addison's disease)

Adrenocortical insufficiency

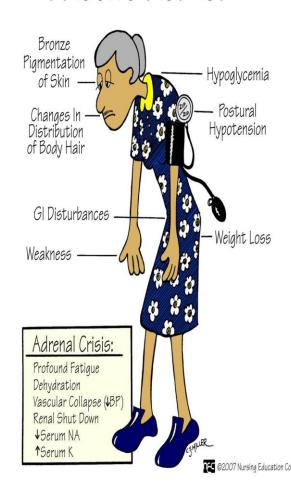
- primary causes, ie. Addison's disease
 - autoimmune disease, tumors, infection, hemorrhage, metabolic failure, ketoconazole (glucocorticoid antagonist activity)
- secondary causes
 - hypopituitarism, suppression by exogenous steroids

<u>Adrenal insufficiency</u>: (<u>Addison's disease</u>)

- Increased excretion of sodium and water.
- Reduction in ECF volume.
- Tendency toward <u>low blood pressure</u>.
- Complete absence of aldosterone, the volume depletion may be severe.
- Hypoglycemia
- Skin pigmentation

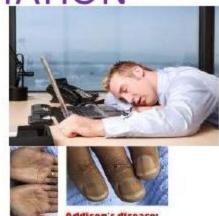
The person is allowed to eat large amounts of salt and drink large amounts of water to balance the increased urine output of salt and water.

ADDISON'S DISEASE



CLINICAL MANIFESTATION

- General weakness and becoming easily tired.
- Darkened areas of skin ('pigmentation').
- Blood pressure is low and falls further when you stand which can make you dizzy.
- · Being off your food and weight loss.
- Feeling sick and vomiting from time to time.
- Abdominal pains which may come and go.
- Diarrhoea or constipation which may come and go.
- · Cramps and pains in muscles.
- Craving for salt, or salty foods and drinks.
- Menstrual periods in women may become irregular, or stop.





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Endocrine Images: Adrenal Insufficiency



This slide of identical twins is from Dr. Hammer's lecture and is meant to emphasize the hyperpigmentation and thin body habitus that is often seen in primary adrenal insufficiency (the woman with adrenal insufficiency is on the right). Hyperpigmentation may also be seen in the extensor surfaces of the limbs (knuckles, elbows, knees), in newly formed scars and in palmar creases and buccal mucosa. (What's the cause?)

COMMON DISORDERS OF ADRENOCORTICAL FUNCTION

Conn's (Primary Hyperaldosteronism)	Aldosterone 个	Hypertension, hypokalemia, alkalosis
Secondary Hyperaldosteronism	Aldosterone 个 Renin 个	Hypertension, hypokalemia, alkalosis
<u>Cushing's</u>	Cortisol ↑ ACTH ↑ or ↓ (depending on aetiology)	Altered appearance, bruising, muscle wasting, hypertension, diabetes, hypokalemia
<u>Addison's</u>	Cortisol ↓ Aldosterone ↓ or ↔	Hypoglycemia, hyponatremia, hyerkalemia and raised urea Acute: abdominal pain, vomiting, dehydration and hypotension Chronic: fatigue, pigmentation, anorexia, weight loss and postural hypotension

Herbs to Lower Cortisol



Ginseng



Holy Basil



Ashwaganda



Astralagus



Licorice root



Rhodiola

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