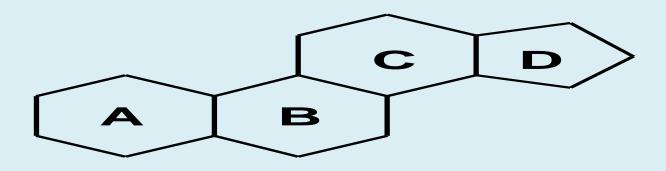
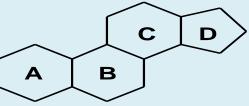
# **Steroidogenesis 1**



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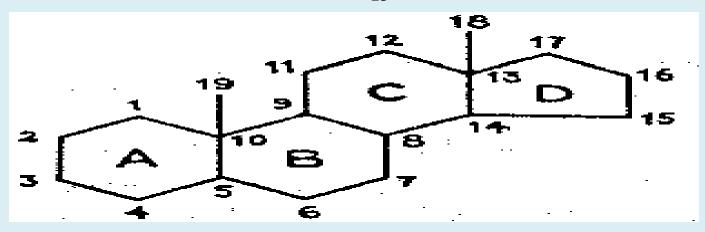
# **Steroids**

• Steroids are group of plant and animal lipids that have a similar tetracyclic nucleus.

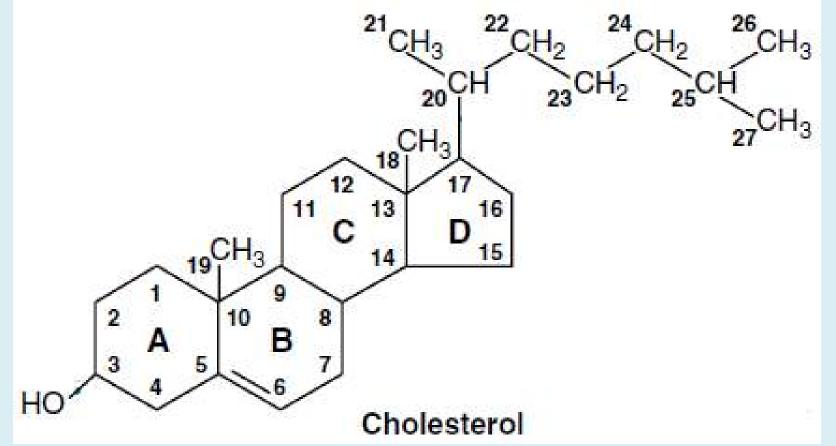


#### **Steroid nucleus:**

- So these rings are composed of 17 carbon atoms besides two methyl groups (C<sub>18</sub>, C<sub>19</sub>).
- There is a methyl group at C<sub>10</sub> (it makes C 19).
- And there is another methyl group at  $C_{13}$  (it makes C18).



- <u>Cholesterol consists of</u>
- Four fused hydrocarbon rings (A, B, C, and D, called the "steroid nucleus")
- Eight-carbon branched hydrocarbon chain attached to C17 of the D ring.
- Ring A has a hydroxyl group at C-3, and ring B has a double bond between C-5 and C-6.



#### **Cholesterol Sources**

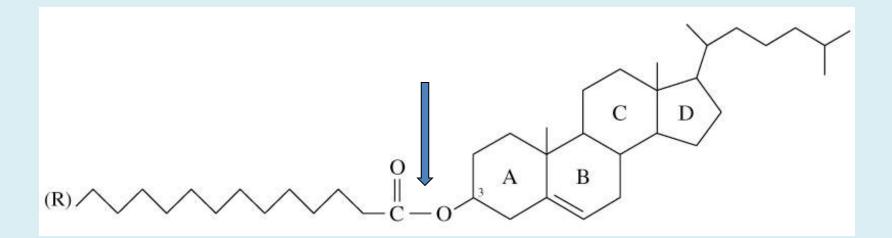
- 1. It is formed in the body from acetyl CoA. Most of the cholesterol is synthesized by the liver.
- 2. It is present in diet: egg yolk, meat, liver and brain.

#### **Biomedical importance:**

- 1- It is the main sterol in human body (Nervous tissue, brain, suprarenal gland, and in bile, ,,).
- 2- It is present in blood (normal level 150-200 mg / dl).
- 3- It is often found as cholesterol ester (in combination with fatty acids). The fatty acid is attached to the hydroxyl group (at C3)
- e.g. Cholesteryl oleate or linoleate.

# **Cholesterol esters (CE)**

- Cholesterol is converted to cholesteryl esters for cell storage or transport in blood
- Fatty acid is esterified to (C-3) OH of cholesterol
- Cholesterol esters are very water insoluble (hydrophobic) and must be complexed with phospholipids or amphipathic proteins for transport



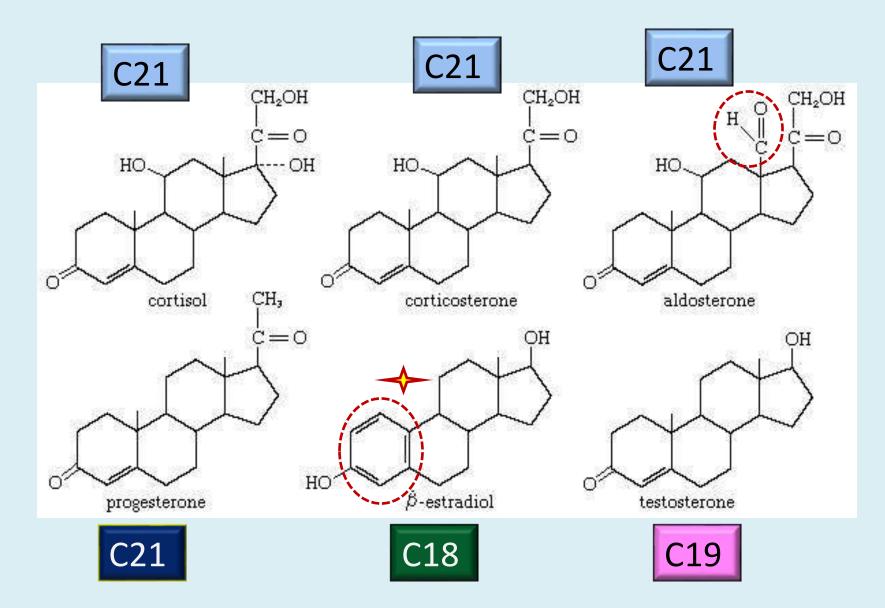
#### **Biomedical importance:**

- 4- It is a major constituent of the plasma membrane.
- The fused ring system makes cholesterol less flexible than most other lipids.
- 5- It is the precursor of:
- Sex hormones
- Cortical hormones
- Vitamin D
- Bile acids.

6- High levels of cholesterol <u>in blood</u> will lead to its precipitation in the wall of blood vessels "atherosclerosis". Also high levels of blood cholesterol may lead to stones in <u>gall bladder</u> (gall stone).

- Cholesterol is the precursor of all five classes of steroid hormones:
- <u>Glucocorticoids</u>: 21 carbons (eg; Cortisol )
- Mineralcorticoids: 21 carbons (eg; Aldosterone )
- **Progestins:** 21 carbons (eg; Progesterone )
- <u>Androgens</u>: 19 carbons (eg; Testosterone )
- **Estrogens:** 18 carbons (eg; Estradiol )

# **Steroid hormones: chemical structure**



## **Steroid hormones: Characteristics**

- Steroid hormones: produced in the adrenal cortex, testis, ovary, and some peripheral tissues (adipose tissue, the brain).
- All steroids are lipid soluble and thus are freely permeable to membranes so are not stored in cells.
- Steroid hormones are not water soluble so have to be carried in the blood complexed to specific binding globulins.
- > Corticosteroid binding globulin carries cortisol
- Sex steroid binding globulin carries testosterone and estradiol.

# How does the synthesis of steroids differ from that of peptide hormones?

- While peptide hormones are encoded by specific genes, steroid hormones are synthesized from the enzymatic modification of <u>cholesterol</u>.
- Thus, there is <u>no gene</u> which encodes ex, aldosterone.
- As a result:
  - There are far fewer different types of steroid hormones than peptide hormones.
  - Steroid structures are the same from species to species.
  - The regulation of steroidogenesis involves control of the enzymes which modify cholesterol into the steroid hormone of interest.
- Steroid hormones are slower acting and have longer half-life than peptide hormones.

# **Steroid hormones: mechanism of action**

 Enzymes which produce steroid hormones from cholesterol are located in mitochondria and smooth endoplasmic reticulum (ER).

#### **Functions of steroid hormones:**

- Steroid hormones play important roles in:
  - carbohydrates regulation (glucocorticoids)
  - minerals balance (mineralocorticoids)
  - reproductive functions (gonadal steroids)
- Steroids also play roles in inflammatory responses, stress responses, bone metabolism, cardiovascular fitness, behavior, cognition, and mood.

# **Steroid hormones functions**

Product	Functions	
Progesterone	prepares uterus lining for implantation of ovum	
Glucocorticoids (cortisol) (produced in adrenal cortex)	promote gluconeogenesis; favor breakdown of fat and protein (fuel mobilization); anti-inflammatory	
Mineralocorticoids (aldosterone) (produced in adrenal glands)	maintains blood volume and blood pressure by increasing sodium reabsorption by kidney	
Androgens (testosterone) (produced in testis primarily)	development of male secondary sex characteristics; prevents bone resorption	
Estrogen (produced in ovaries primarily but also in adipose cells of males and females)	development of female secondary sex characteristics; prevents bone resorption; increase HDL & dec.LDL	

## **Steroidogenic enzymes:**

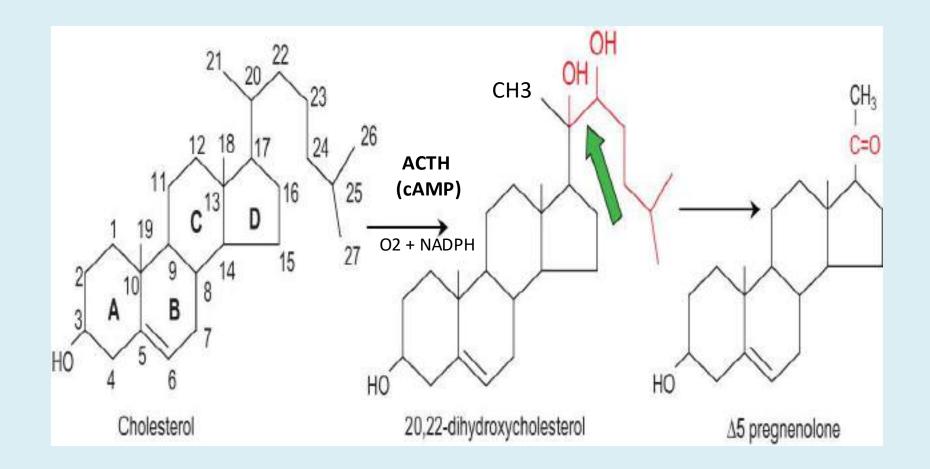
Common name	"Old" name	<b>Current name</b>
Side-chain cleavage enzyme; desmolase	P450 <sub>scc</sub>	CYP11A1
3 beta-hydroxysteroid dehydrogenase	<b>3</b> β-HSD	<b>3</b> β-HSD
17 alpha-hydroxylase/17,20 lyase	P450 <sub>C17</sub>	CYP17
21-hydroxylase	P450 <sub>C21</sub>	CYP21A2
11 beta-hydroxylase	P450 <sub>C11</sub>	CYP11B1
Aldosterone synthase	P450 <sub>C11AS</sub>	CYP11B2
Aromatase	P450 <sub>aro</sub>	CYP19

# **Steroidogenesis:**

- All mammalian steroid hormones are formed from Cholesterol via Pregnenolone.
- <u>The first step</u> is the conversion of cholesterol to pregnenolone, which occurs in the *mitochondria*.
- This reaction is carried out by the enzyme, cytochrome P450 side chain cleavage (P450scc). (also called desmolase, or CYP11A1). Carbons 20 and 22 are sequentially oxidized followed by oxidative cleavage of the bond between them.
- This is a rate limiting, nonreversible enzymatic step in the initiation of steroid biosynthesis.

- <u>Adrenal Steroidogenesis formation of pregnenolone</u>
- Cholesterol in the adrenal is esterified and stored in cytoplasmic lipid droplets.
- Upon stimulation of the adrenal by <u>adrenocorticotropic hormone</u> (ACTH), an <u>esterase</u> is activated, and cholesterol is transported into the mitochondrion by ACTH-dependent steroidogenic acute regulatory (StAR) protein
- Cytochrome P450 side chain cleavage enzyme (P450scc) converts cholesterol to pregnenolone in the inner mitochondrial membrane by:
- 1-Hydroxylations, first at C22 and then at C20
- 2-Side chain cleavage, removal of the six-carbon fragment isocaproaldehyde, to give the 21-carbon steroid.

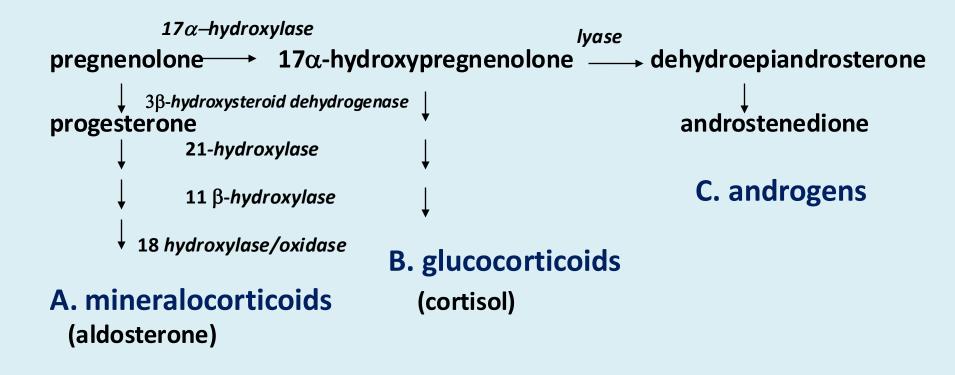
#### **Conversion of cholesterol to pregnenolone**



+ isocaproaldehyde (6C)

# **Adrenal Steroidogenesis**

 Next, pregnenolone can be converted (in smooth ER) into three different pathways, depending upon whether the cell wants to make mineralcorticoids, glucocorticoids, or androgens.



### What determines which pathway is taken?

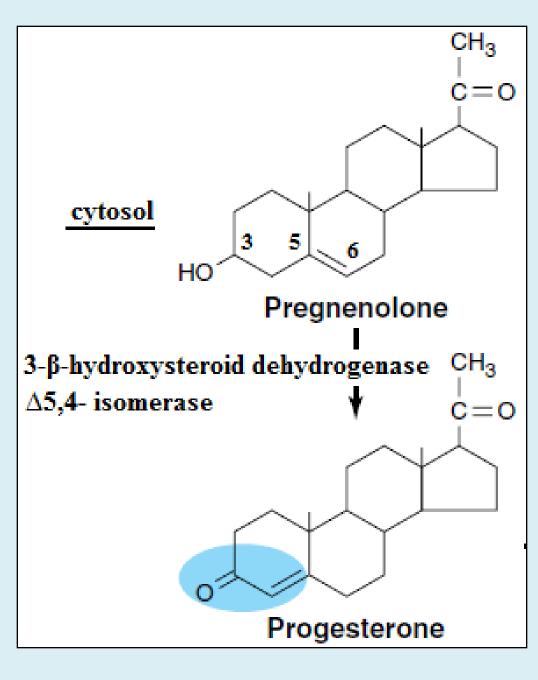
- Each step of the pathway is regulated by a specific enzyme. For ex., 18- hydroxylase and hydroxysteroid dehydrogenases, which are required for aldosterone synthesis are found only in the <u>zona glomerulosa cells</u>, so that the biosynthesis of this mineralocorticoid is confined to this region.
- Different zones of the adrenal cortex have *different relative activities of enzymes,* resulting in different chemical reactions taking place.
- Adrenal steroid biosynthesis involves the shuttling of precursors between mitochondria and the ER

# **1. Mineralocorticoids biosynthesis**

- Occurs in Zona glomerulosa [ZG].
- All mammalian steroid hormones are formed from cholesterol via pregnenolone
- Pathway:
- <u>Pregnenolone</u> is converted to <u>progesterone</u> by  $3\beta$ hydroxysteroid DH ( $3\beta$  -OHSD) and  $\Delta^{5,4}$ - isomerase
- <u>Progesterone</u> is *hydroxylated at C21* to form <u>11-</u> <u>deoxycorticosterone</u> (DOC). DOC is an active (Na+retaining) mineralocorticoid
- Next hydroxylation is at C11 producing corticosterone
- 18-Hydroxylase acts on corticosterone to form aldosterone

#### **Progesterone**

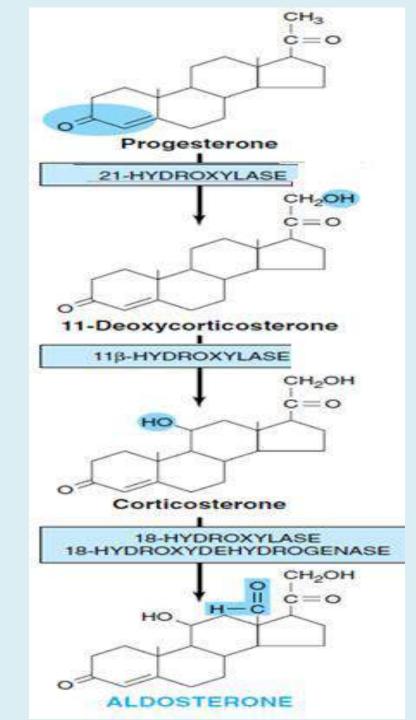
- is synthesized in two steps:
- The 3-hydroxyl group of pregnenolone is oxidized to 3- keto group by 3-βhydroxysteroid dehydrogenase.
- 2- The  $\Delta^5$  double bond is isomerized to a  $\Delta^4$ double bond by  $\Delta^{5,4}$ isomerase.



#### Aldosterone (mineralocorticoid)

- 1- Hydroxylation of progesterone at C-21 by 21-hydroxylase.
- 2- The resulting deoxycorticosterone is hydroxylated at C-11 by 11βhydroxylase.
- 3- The <u>oxidation</u> of the C-18 to an aldehyde then yields aldosterone by:

18-hydroxylase and 18-hydroxysteroid dehydrogenase.



# **2. Glucocorticoids biosynthesis**

- Occurs in the Zona fasiculata [ZF] and Zona reticularis [ZR] of adrenal cortex.
- All mammalian steroid hormones are formed from cholesterol via pregnenolone

#### **Biosynthesis of glucocorticoids (A & B pathways):**

#### \* <u>A. Pathway:</u>

- Pregnenolone is converted to progesterone by 3βhydroxysteroid DH (3 β -OHSD) and Δ<sup>5,4</sup>- isomerase
- Progesterone is hydroxylated at C21 to form <u>11-</u> deoxycorticosterone (DOC).
- Next hydroxylation is at C11 producing <u>Corticosterone</u>. (it has glucocorticoid & weak mineralocorticoid action).

#### **B. Pathway:**

> 17 $\alpha$ -Hydroxylase acts upon either progesterone or pregnenolone to form <u>17  $\alpha$  - hydroxyprogesterone</u> and/or <u>17  $\alpha$  -hydroxypregnenolone</u>.

><u>17  $\alpha$  -hydroxyprogesterone</u> is *hydroxylated at C21* to form <u>11-deoxycortisol</u>.

<u>11-deoxycortisol</u> is then *hydroxylated at C11* to form <u>Cortisol</u>.

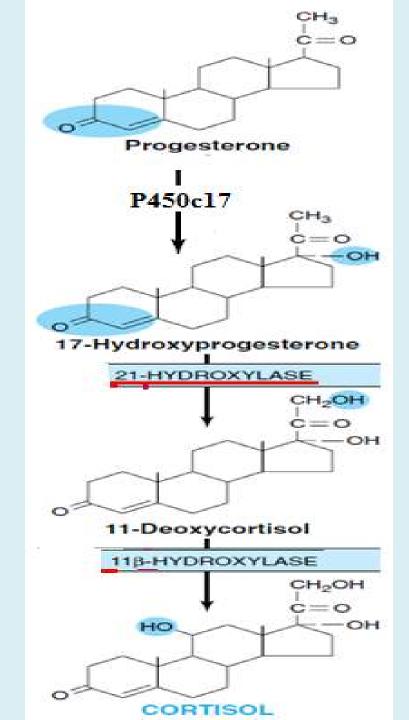
### **Glucocorticoid biosynthesis**

- Cortisol is the most potent natural glucocorticoid hormone in humans.
- Cortisol release is primarily under ACTH.
- <u>Three hydroxylases are needed for cortisol synthesis</u>
  <u>on position:</u>
- <u>C17</u> by 17alpha-hydroxylase, (acts rapidly; present in ER)
- > <u>C21</u> by 21-hydroxylase, (acts rapidly; present in ER)
- <u>C11</u> by 11-hydroxylase, (acts slowly; present in mitochondria).
- If C 11 is hydroxylated first, the action of 17alphahydroxylase is impeded and mineralocorticoid pathway is followed (forming corticosterone or aldosterone depending on the cell type).

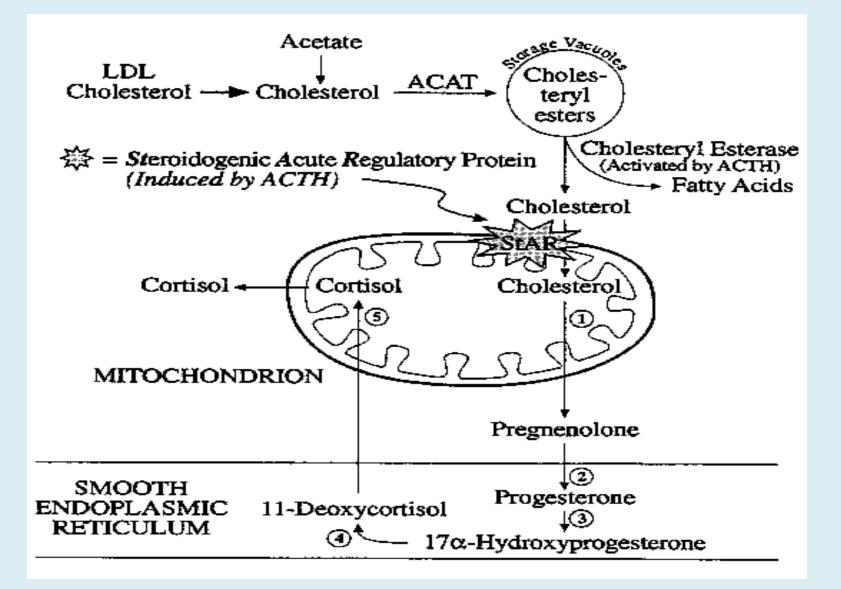
#### **Cortisol**

The major glucocorticoid, is synthesized from progesterone by hydroxylation's of

- <u>C-17</u> by P450c17
- <u>C-21</u> by 21-hydroxylase
- <u>C-11</u> by 11 $\beta$ -hydroxylase



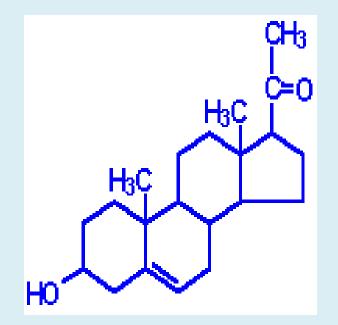
# Adrenal steroid biosynthesis involves the shuttling of precursors between mitochondria and ER

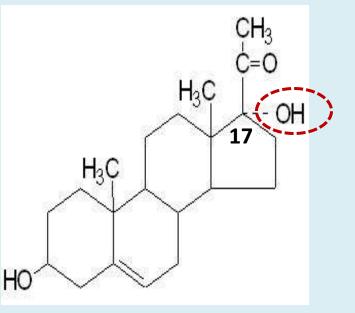


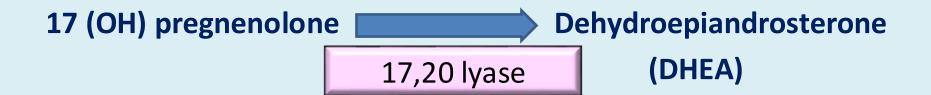
# **3. Adrenal androgens biosynthesis**

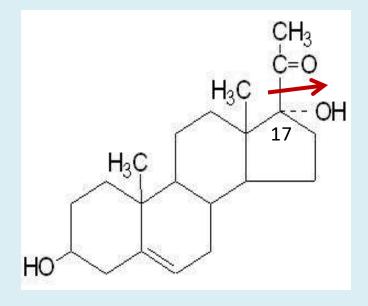
- Produced in the zona reticularis [ZR].
- The major androgen produced by adrenal cortex is dehydroepiandrosterone (DHEA).
- Most of 17-hydroxy pregnenolone follows glucocorticoid pathway.
- Few of 17-hydroxy pregnenolone follows androgen pathway.
- 17alpha-hydroxylase + 17, 20 lyase = dual-function protein.
- <u>Lyase activity</u> is important in both adrenals and gonads. Lyase acts exclusively on 17-hydroxy containing molecules.

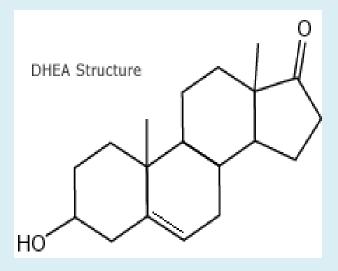
# Pregnenolone17(OH) pregnenolone17α-hydroxylase

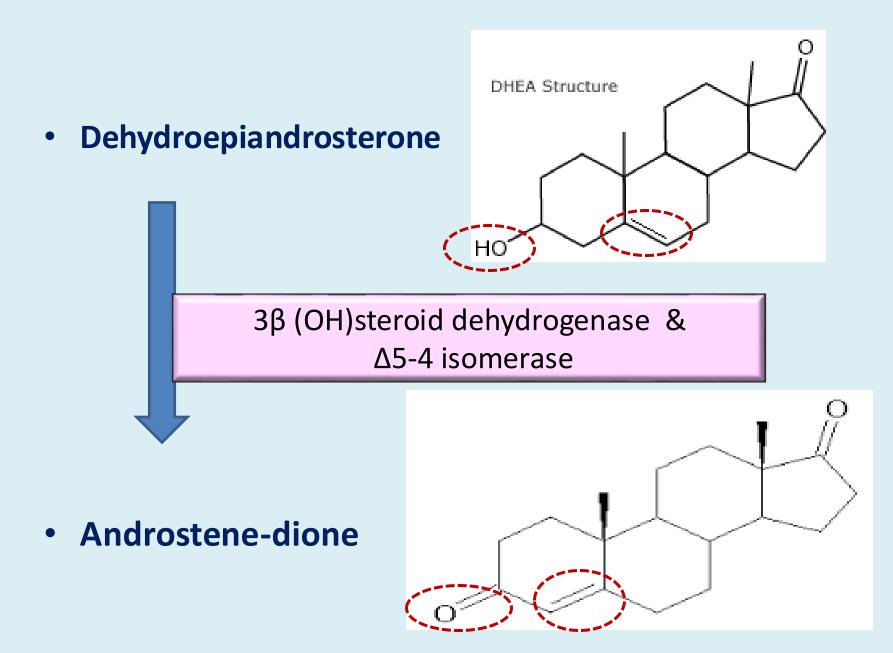




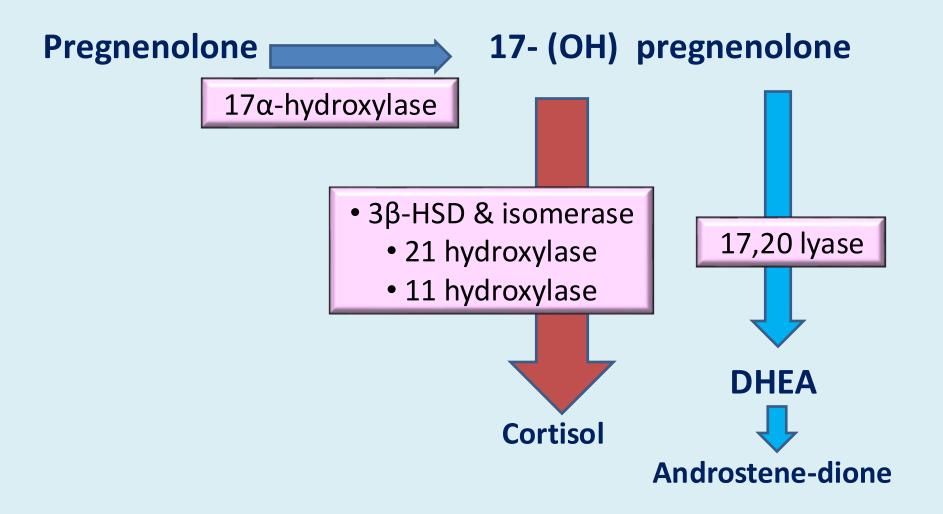








# **Adrenal androgens:**



# **Steroid synthesis: overall**

