

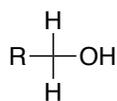
**Alcohol and Ether Nomenclature**

Alcohol

Ether

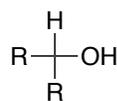
R-OH

R-O-R



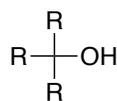
Primary

1°



Secondary

2°



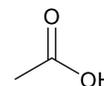
Tertiary

3°

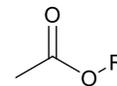
R ≠



Carbonyl



Acid



Ester

Alcohols are classified as primary (1°), secondary (2°), or tertiary (3°), depending on the number of organic groups bonded to the hydroxyl bearing carbon.

**Note:** -OH is called hydroxyl, hydroxy or alcohol

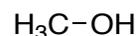
-OR is called Alkoxy group

**Naming:**

1. Find the longest chain, with the maximum number of OH groups.
2. Number in such a way to give the **first OH** the lowest number
3. Drop the “e” of the alkane name, add “ol”

**Note:** the alcohol (-OH) takes priority over ethers, multiple bonds, and halogens

**Examples:**



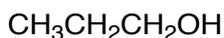
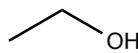
methanol

-toxic  
wood alcohol



ethanol

grain alcohol

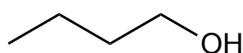
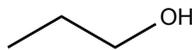


1-propanol

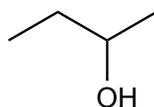
n-propanol

propan-1-ol

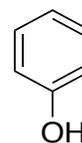
propyl alcohol



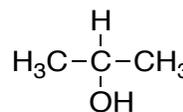
1-butanol  
n-butanol



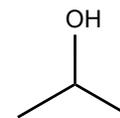
2-butanol



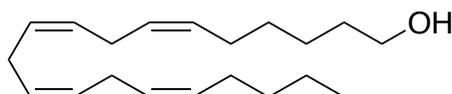
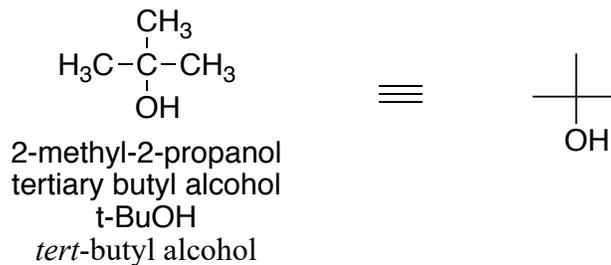
Hydroxybenzene  
Phenol  
C<sub>6</sub>H<sub>6</sub>O



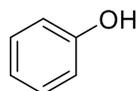
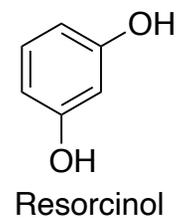
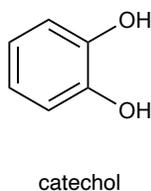
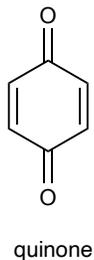
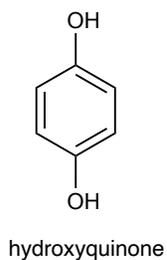
2-propanol  
propan-2-ol  
Isopropyl alcohol  
isopropanol



2° alcohol



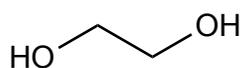
**Eicosa-6Z,9Z,12Z,15Z-tetraen-1-ol**



Phenol

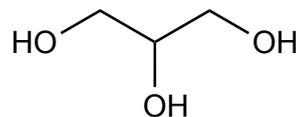
Phenol is very common in nature, phenol based structures occurred in lignin

Catechol: part of some neural transmitter



ethan-1,2-diol  
 ethylene glycol

1,2-ethane-diol

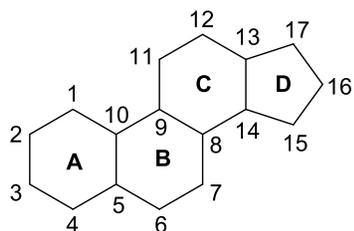


1,2,3-propantriol  
 glycerol

glycerine

**Note:** most alcohols are flammable, however, as the chain gets longer, the molecules would tend to stick together (i.e., intermolecular forces) and would be less flammable.

## Steroids

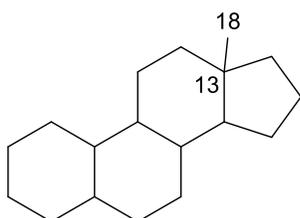


**Steroid Skeleton**

**Groups above:**  $\beta$  (beta)

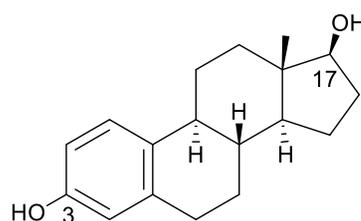
**Groups below:**  $\alpha$  (alpha)

## Types of Steroids

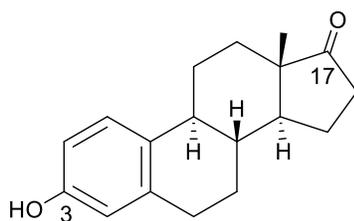


**Estrane**

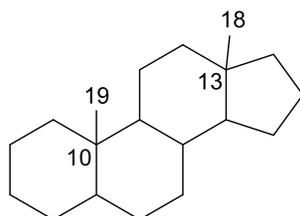
Characterized by a methyl group in C13



**Estradiol**

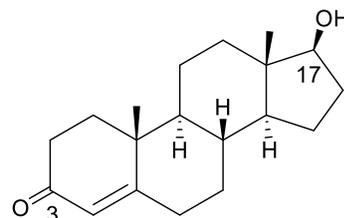


**Estrone**

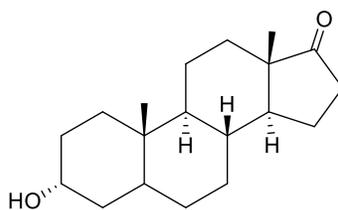
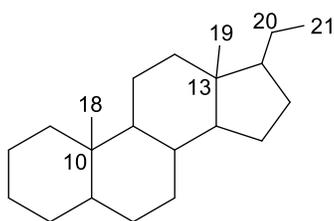


**Androstane**

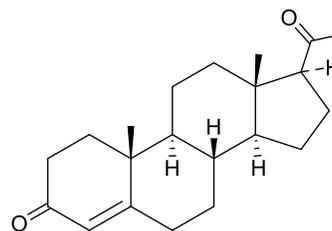
Characterized by the presence of  $-\text{CH}_3$  groups in C10 and C13



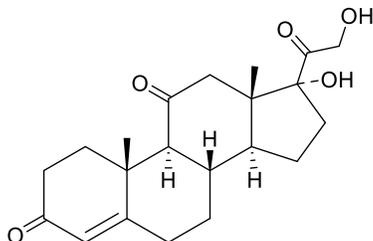
**Testosterone**

**Androsterone****Pregnane**

- Characterized by two -CH<sub>3</sub> groups in C10 and C13, and a substituent in C17.
- Not biologically active

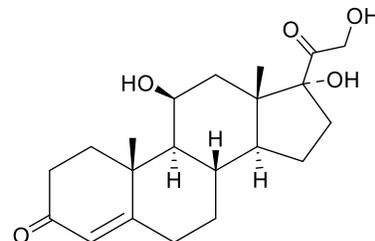
**Progesterone**

Pregnancy Hormone

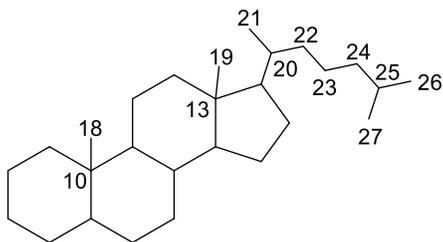
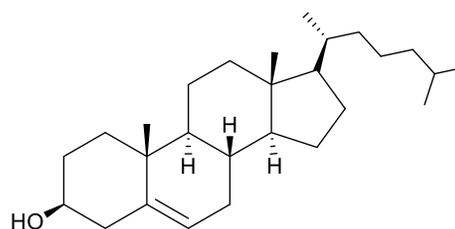
**Cortisone**

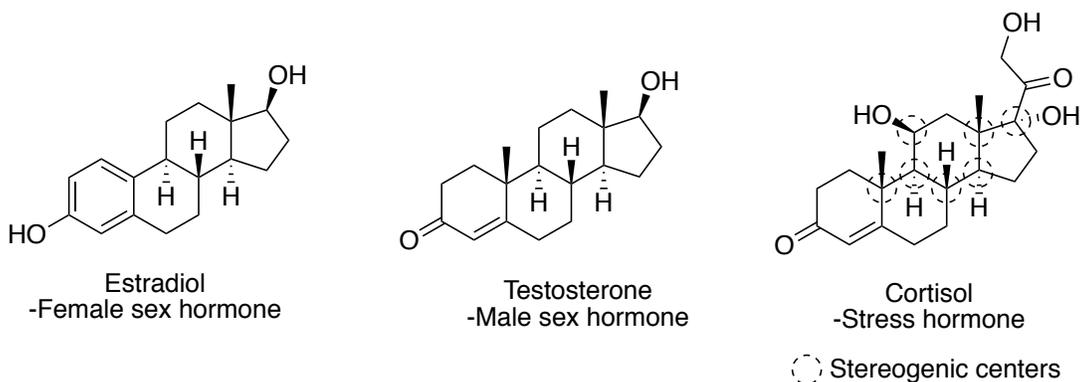
(Adrenocorticoid)

Characterized by a carbonyl at C11 and pregnane skeleton

**Cortisol**

Stress Hormone

**Cholestane****Cholesterol**



### Polyols:

If more than one hydroxyl group is present, a prefix is added to the “ol” :

2 OH's	diol (glycol)
3 OH's	triol
4 OH's	tetraol
5 OH's	pentaol

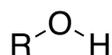
### Physical Properties of Alcohols

- The hydroxyl group is a very polar group. This allows small alcohols (methanol, ethanol, propanols) to be miscible with water (if the number of C < 4) as they are good hydrogen bond donors *and* acceptors.
- Soluble in H<sub>2</sub>O, as the #C increases, the solubility decreases.
- Alcohol densities are usually  $\rho < 1.0 \text{ g/cm}^3$ .
- They have high boiling and melting points, again due to their hydrogen bonding capabilities.

Look at the following comparisons:

<u>Name</u>	<u>Methanol</u>	<u>Ethane</u>	<u>Ethanol</u>
<i>Formula</i>	CH <sub>3</sub> OH	CH <sub>3</sub> CH <sub>3</sub>	CH <sub>3</sub> CH <sub>2</sub> OH
<i>Molecular Weight (g/mol)</i>	32	30	46
<i>Boiling Point (° C)</i>	65	-89	78.5
<i>State (at room temp)</i>	liquid	gas	liquid

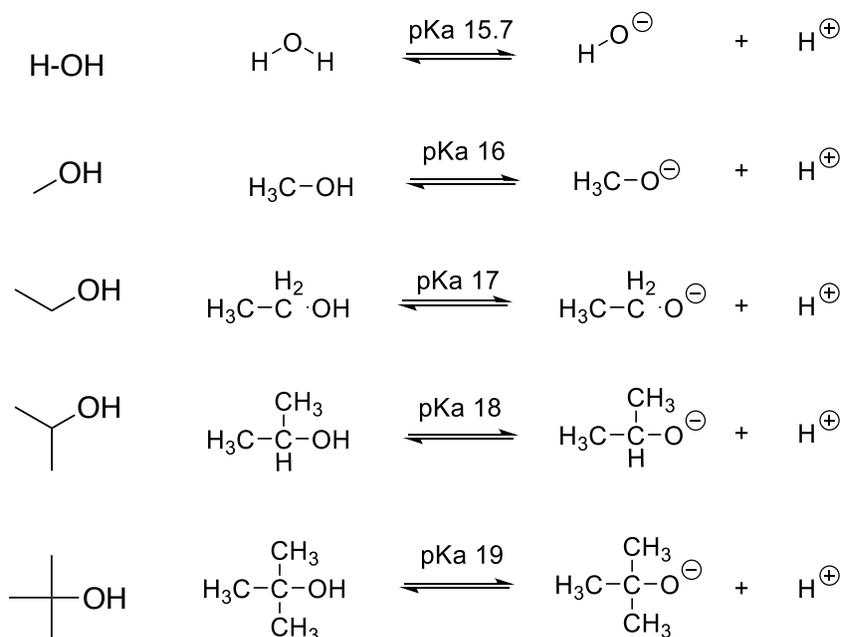
Ethane has almost the same molecular weight as methanol. However, the boiling point is much lower than methanol. Methanol molecules like to stick together via H-bonding while ethane molecules interact with each other via hydrophobic interactions.



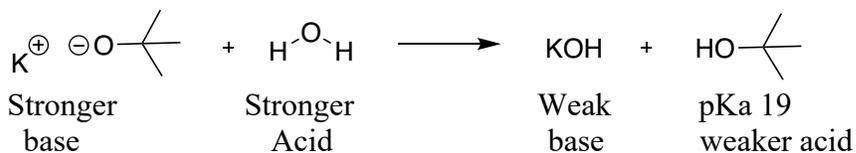
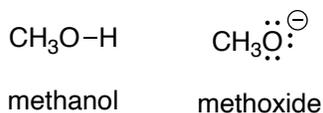
O-H bond - easy to break

R-O bond - hard to break; always needs a strong acid to break and can proceed either S<sub>N</sub>-1 / S<sub>N</sub>-2, or E1 / E2

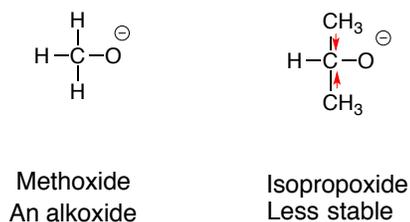
### Acidity of R-OH



- pKa depends on conjugated base stability
- Harder to make a t-butoxide than methoxide. The alkyl group donates electron density to the C-O bond and destabilizes the negative charge (less favorable).



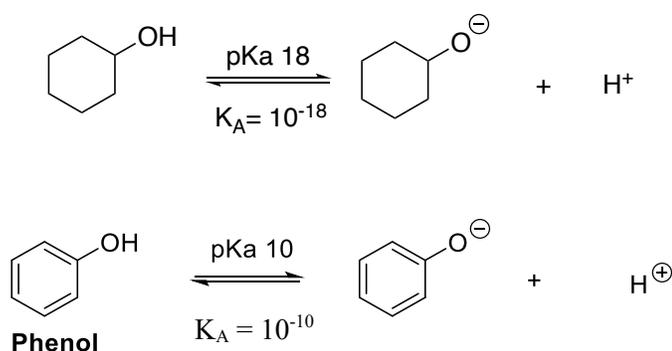
### Methoxide vs. Isopropoxide:



Alkyl groups donate electrons through single bonds, destabilizing the negative charge (Inductive Effect), therefore, isopropoxide is less stable than methoxide

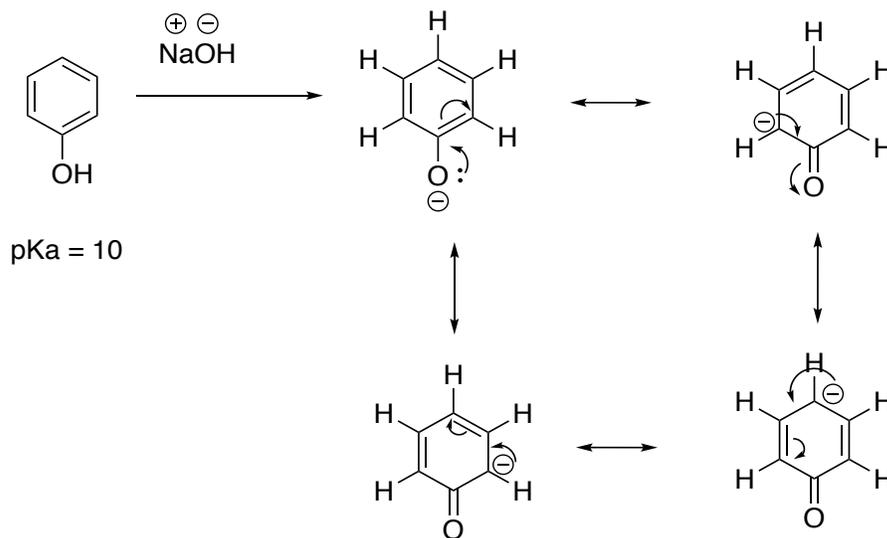
- Inductive effect – donation or withdrawal through single bonds

### Conjugated/Aromatic R-OH

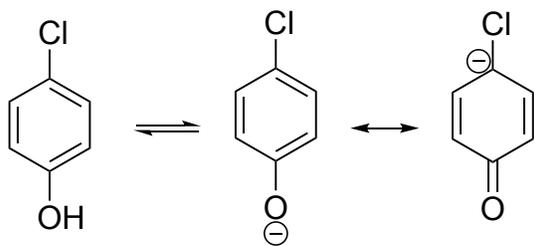


- More acidic than H<sub>2</sub>O
- Resonance (resonance effect) takes electron density away from the O atom, resulting in stabilization of the negative charge.
- Resonance effect is strong through  $\pi$  system

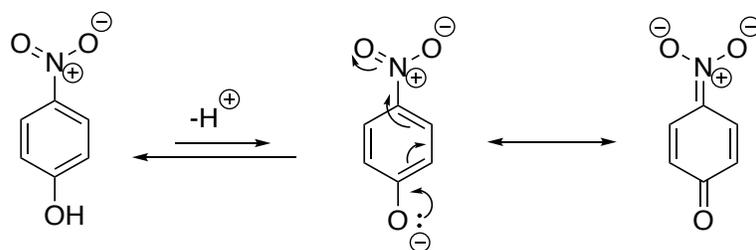
### Example 1: Phenol



The alkoxide of phenol is a conjugated anion and is therefore much more acidic

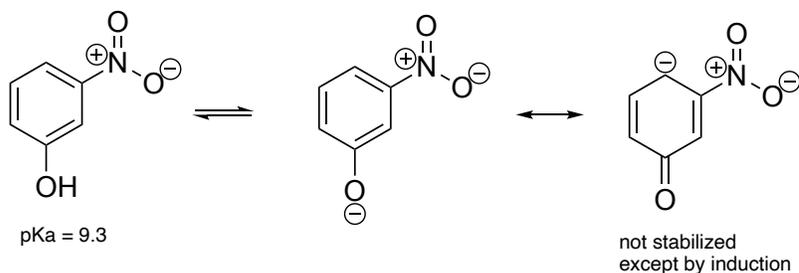
**Example 2: *p*-Chlorophenol**

pKa = 9.3

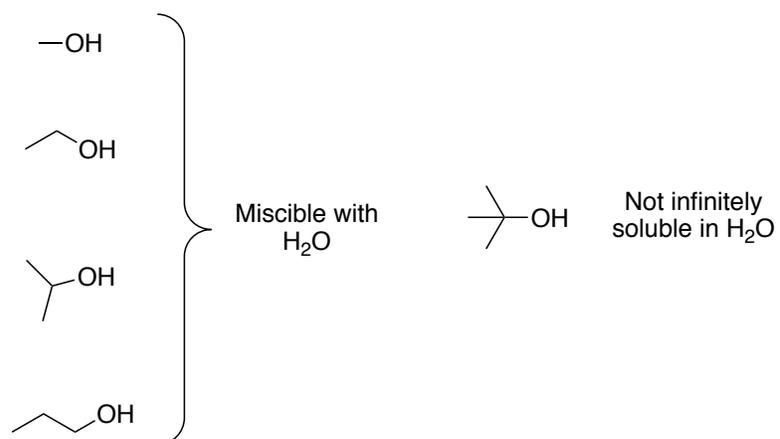
**Example 3: *p*-Nitrophenol**

pKa = 7.3

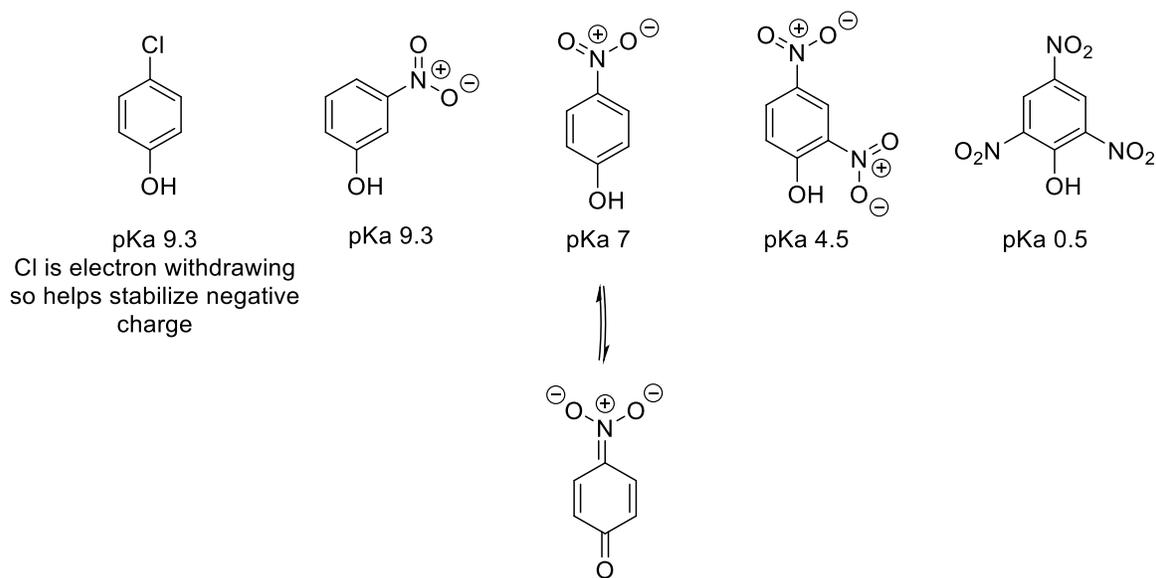
*p*-Nitrophenol is more acidic than phenol because on top of the resonance forms that phenol contains, *p*-nitrophenol also contains the above extra resonance form, making the proton on the alcohol even more acidic.

**Example 4: *m*-Nitrophenol**

Less acidic than *p*-nitrophenol because there is less stabilization of the negative charge. The negative charge is not conjugated with the nitro group double bond.

**Miscibility of Alcohols with Water**

\*butanol is soluble in  $\text{H}_2\text{O}$  but not miscible

**More examples:**

- As you get more resonance possibilities, the negative charge is more spread out across the molecules, and is more stabilized, resulting in lower pKa (more acidic).