

**Formal Charge**

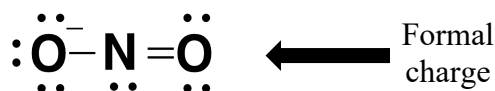
- Convention to keep track of charges
- $\Sigma$  (sum of) of formal charges on all atoms in a molecule = the overall charge on the molecule

**Rules for calculating formal charge**

- Add the number of protons (the atomic number) in the nucleus
- Subtract the number of inner shell electrons
- Subtract the number of unshared electrons
- Subtract  $\frac{1}{2}$  of the number of shared outer shell electrons

**Examples:**

1. **NaNO<sub>2</sub> (sodium nitrite; food preservative)**  
**Nitrite anion**



Overall charge on the nitrite anion is = **-1**

Single bonded oxygen:

+8 (number of protons)

-2 (1s electrons)

-6 (unshared electrons)

$\frac{1}{2} \times 2 = -1$  (1/2 of shared electrons)  
**-1**

Central N:

+7 (number of protons)

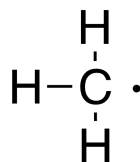
-2 (1s e<sup>-</sup>)

-2 (unshared e<sup>-</sup>)

-3 (1/2 shared e<sup>-</sup>)

= **0**

2. **Methyl radical (sp<sup>3</sup>, tetrahedral)**



Overall charge on the methyl anion is = **0**

Very unstable since it doesn't have an inert gas configuration

Formal Charge on Carbon

+6 (number of protons)

-2 (1s electrons)

-1 (unshared electrons)

$\frac{1}{2} \times 6 = -3$  (1/2 of shared electrons)  
**0**

**3. Methyl cation (carbocation,  $sp^2$ , planar)**

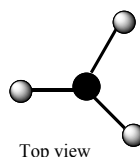
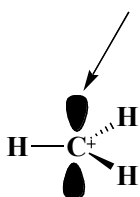
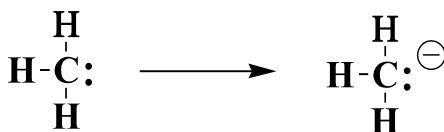
- ( $sp^2$  hybridized carbon, planar shape)
- can be reactive intermediate in principle

Overall charge on the methyl anion is = +1

Formal Charge on Carbon

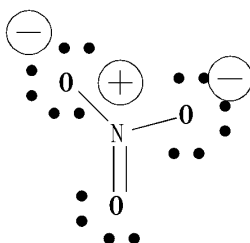
$$\begin{aligned}
 &+6 \text{ (number of protons)} \\
 &-2 \text{ (1s electrons)} \\
 &0 \text{ (unshared electrons)} \\
 &\frac{1}{2} \times 6 = \underline{-3} \text{ (1/2 of shared electrons)} \\
 &\quad \quad \quad +1
 \end{aligned}$$

Empty p orbital

**4. Methyl anion ( $sp^3$ , tetrahedral)**Formal Charge on Carbon

$$\begin{aligned}
 &+6 \text{ (number of protons)} \\
 &-2 \text{ (1s electrons)} \\
 &-2 \text{ (unshared electrons)} \\
 &\frac{1}{2} \times 6 = \underline{-3} \text{ (1/2 of shared electrons)} \\
 &\quad \quad \quad -1
 \end{aligned}$$


Overall charge on the methyl anion is = -1


**5. Sodium Nitrate ( $NaNO_3$ )**Formal Charge on Nitrogen

$$\begin{aligned}
 &+7 \text{ (number of protons)} \\
 &-2 \text{ (1s electrons)} \\
 &0 \text{ (unshared electrons)} \\
 &\frac{1}{2} \times 8 = \underline{-4} \text{ (1/2 of shared electrons)} \\
 &\quad \quad \quad +1
 \end{aligned}$$

**Resonance Structures:** Different drawings (or pictures) of the same molecule made by moving electrons but not atoms

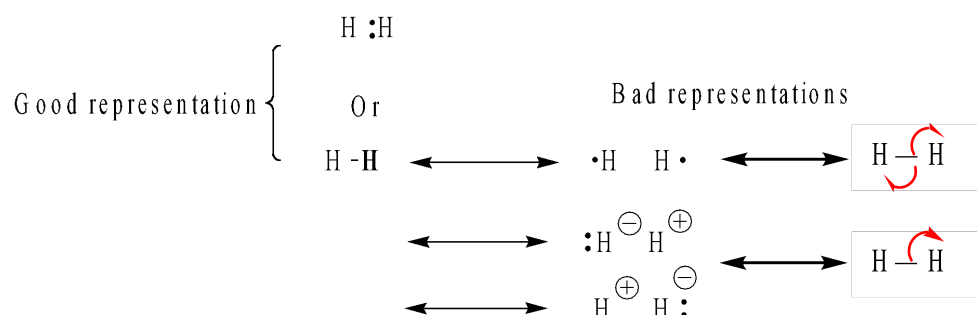
- Move the electrons, keeping the position of the atoms the same
- Good resonance structures:
  - o Maintain inert gas configuration around each atom
  - o Avoid the separation of charges
- Avoid like-charges on adjacent atoms
- Double-headed arrow ( $\longleftrightarrow$ ) is used to indicate resonance forms. Fish Hook and double-headed arrows are used to show electron movement

  
Double Headed Arrow  
Show movement of  $2e^-$

  
Fish Hook Arrow  
Show movement of  $1e^-$

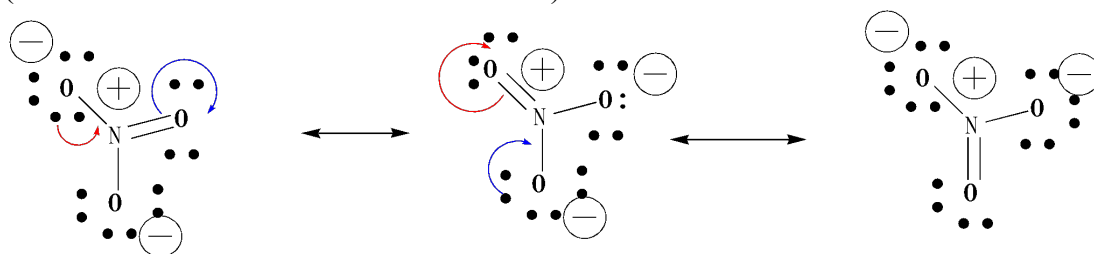
## Examples

### 1. Hydrogen gas, $H_2$



In the bad representations, non- inert gas configuration and extra charges have been created

### 2. Sodium Nitrate, $NaNO_3$ , $Na^+ NO_3^-$ (Nitrate has 3 resonance forms shown here)

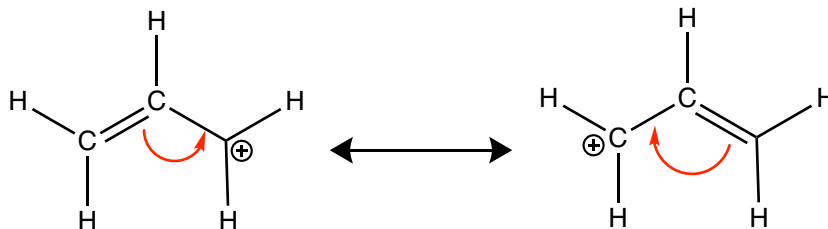


**No inert gas configuration disrupted**

**No extra charge created**

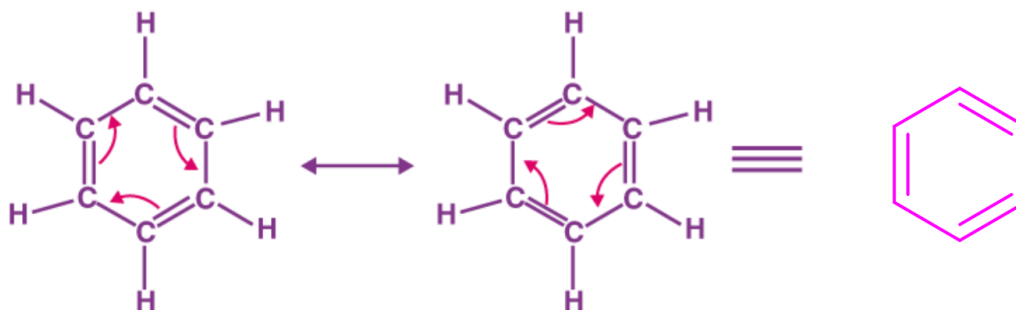
- The O atoms contain partial single and double bond characteristics (each O has  $-2/3$  charge)

### 3. Allyl Cation

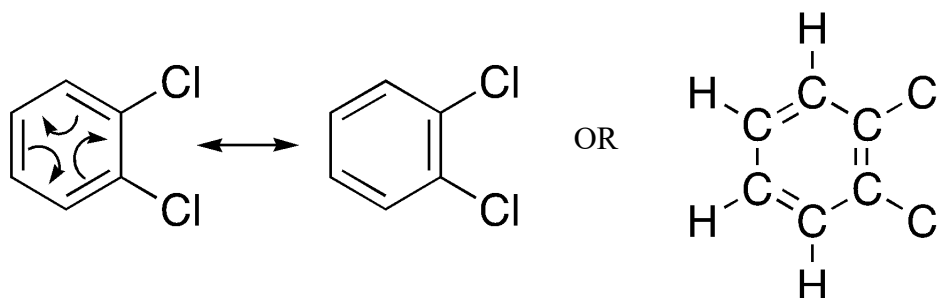


-electrons are delocalized between the two carbons on both side of the central C and C atoms has  $-1/2$  charge and contains partial double and single bond character.

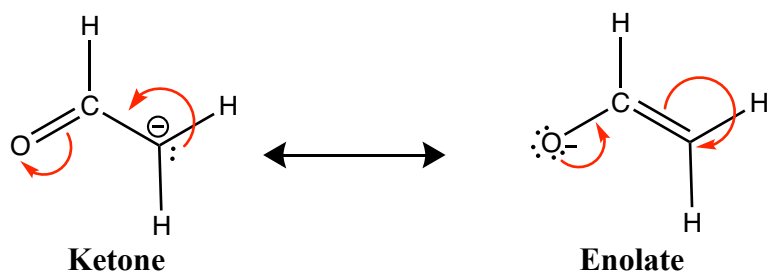
### 4. Benzene, $C_6H_6 = \Phi$



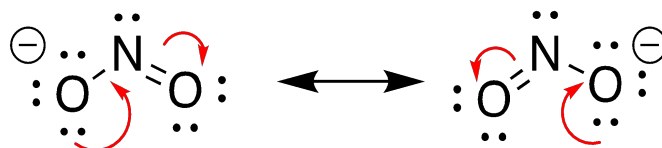
### 4. 1,2-Dichlorobenzene



### 5. Keto-Enol

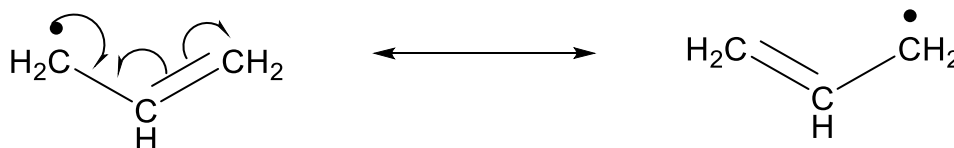


### 6. Sodium Nitrite, $\text{NaNO}_2$



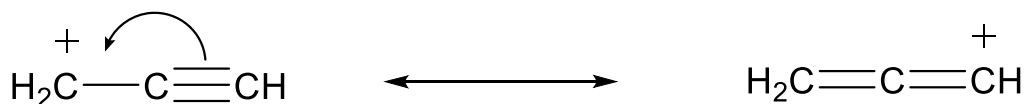
Nitrite anion is reactive in both O atoms. Electrons are delocalized in more than one atom – both O atoms has  $-1/2$  charge and contains partial double and single bond character.

### 7. Allyl Radical



The radical is relatively stable due to resonance.

### 8. Propyne cation



**Intermolecular Forces:** forces present between molecules, governed by electronegativity

- Attractive intermolecular forces:

- i) **Hydrogen bonding** – strongest on per atom basis (e.g. base recognition in forming DNA helix) (also in RNA)
  - *Linus Pauling - development of H bonding*
- ii) **Dipole-dipole interaction** (Intermediate strength)
- iii) **London forces** (temporary dipole; hydrophobic bonding) – weakest on per atom basis – distortion of inner shells.

### Electronegativity:

- An atom's desire for electrons (negative charge).
- On the periodic table, electronegativity increases as you go from left to right (up to inert gases, which are not electronegative) and as you go from down to up
- Halogens (F, Cl, Br, I) are highly electronegative
  - o i.e. Fluorine is the most electronegative atom (wants to gain the inert gas configuration of Ne) and is small (has few electrons)
- It influences acidity of H's attached, as well as the intermolecular forces between molecules.

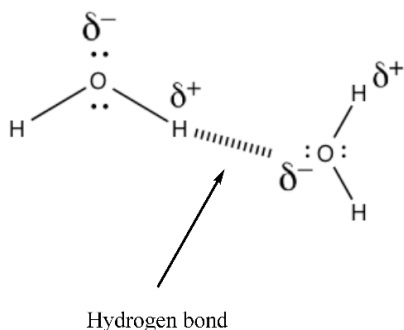
### Hydrogen Bonding:

- Strongest intermolecular attractive force
- Need H directly attached to a very electronegative atom (N, O, F, Cl, Br, I)
  - o Known as **donors**
- Very electronegative atom needs a lone pair of electrons (N, O, F, Cl, Br, I)
  - o Known as **acceptors**

### Examples:

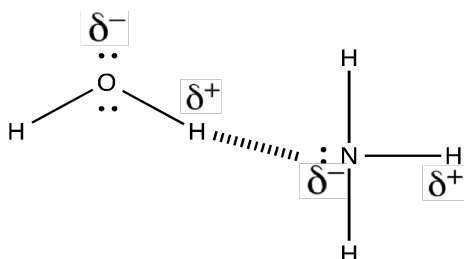
**Methane:** (CH<sub>4</sub>) Incapable of hydrogen bonding, has a low boiling point because the intermolecular forces are weak.

### H-O-H (water):



- Oxygen is electronegative and it is sp<sup>3</sup> hybridized
- The partial positive charge on H and the partial negative charge on O lead to their attraction
- Results in high boiling point (100 C) and high melting point by self-association
- HF, H<sub>2</sub>O and NH<sub>3</sub> form hydrogen bonds

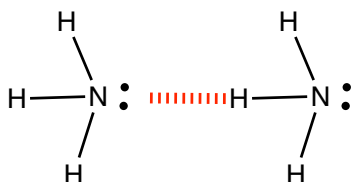
Water can form “temporary” bonds, resulting in a large number of intermolecular bonds.



- Water is a liquid at RT while ammonia is a gas
- Oxygen is more e-neg than nitrogen, so the protons on water have a higher positive partial charge than the protons on ammonia
- In an ammonia solution, water would be the hydrogen bond donor and ammonia would be the acceptor
- Water dissolves ammonia very well – up to 18M

### Ammonia:

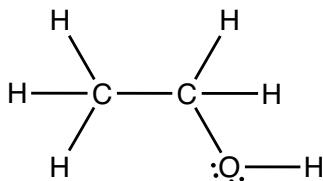
- both H-bond acceptor and donor
- H-bond is weaker than the H-bond of water because N is less electronegative than O
- BP: -33°C (much higher than methane)

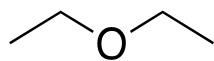


Acceptor (lone pair). Donor (H available)

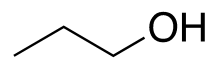
### Ethanol

- both H-bond acceptor and donor
- BP: 78.5 °C



**Hydrogen bonding in mixtures:****Diethyl Ether**

- Cannot hydrogen bond to itself
  - Has no H directly attached to oxygen (No donor)
  - Can H-bond to water because it has an acceptor
- Has a low boiling point
- Will not dissolve in water very well (although a little bit will be dissolved)

***n*-Propanol  
(1-Propanol)**

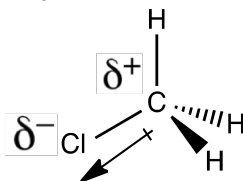
- Can hydrogen bond to itself
  - Has H directly attached to oxygen
- Has a high boiling points relative to its size due to hydrogen bonding
- Can dissolve in water very well

**Dipole-Dipole Interactions:**

Dipole drawing convention:

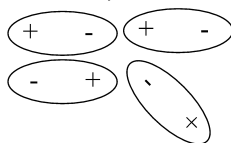


Partial positive charge is the “plus” end, partial negative charge is the arrow head.

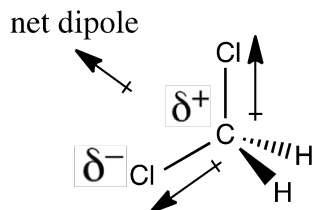
**Chloromethane**, methyl chloride;  $\text{CH}_3\text{Cl}$ 

- H and C have similar electronegativity values (non-polar bond)
- Cl is very electronegative due to the fact that it only needs one electron to get inert gas configuration.
- Electron density is pulled toward the chlorine atom, creating a net dipole toward chlorine atom. A net dipole is the vector sum of individual bond dipoles.
- Has a higher MP and BP than methane

Dipoles in different molecules tend to line-up temporarily with each other (partial positive / negative charge on the molecule) – causes molecules to “stick” to each other



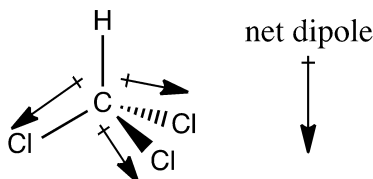
**Dichloromethane**, methylene chloride;  $\text{CH}_2\text{Cl}_2$ . (Methylene =  $\text{CH}_2$  group)



- Liquid at room temperature BP  $40^\circ\text{C}$  MP  $-95^\circ\text{C}$
- More polar than chloromethane
- Not miscible with water

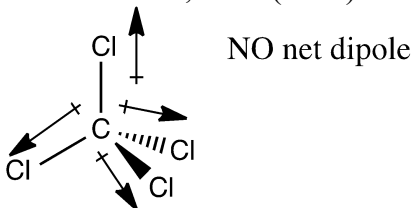
Net dipole: overall vector sum of all the bond dipoles.

**Trichloromethane**, chloroform;  $\text{CHCl}_3$



- More polar than methylene chloride BP  $61^\circ\text{C}$  MP  $-64^\circ\text{C}$
- Higher than dichloromethane due to dipole dipole interaction

**Tetrachloromethane**, carbon tetrachloride;  $\text{CCl}_4$  (toxic)



- Non-polar molecule (net-zero dipole)
- Has temporary dipoles since chlorine is polarizable (see below), BP  $\sim 77^\circ\text{C}$
- Historically used as a dry-cleaning fluid

### London Forces:

- Also known as dispersion forces, temporary dipoles or Van der Waals forces (less good)
- Weakest attractive force
- Distortion of filled outer shell electrons
- Principal effect in hydrophobic interactions

### Atoms

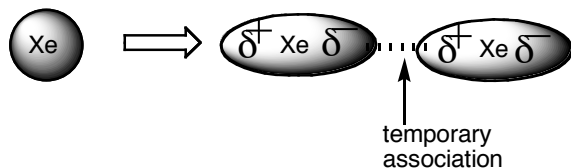
### Boiling Point

He	-269 $^\circ\text{C}$	Small atom/ Low polarizability ↓
Ne	-246 $^\circ\text{C}$	
Ar	-186 $^\circ\text{C}$	
Kr	-153 $^\circ\text{C}$	

Xe

-108 °C

Large atom/ High polarizability

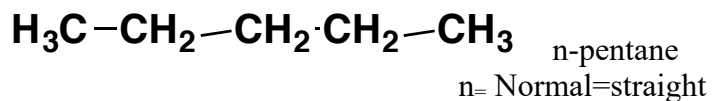


- The larger the atom (expanded electron density), the easier the formation of temporary dipoles.

Steric effect:  
interaction of  
a filled shell  
of electrons.  
Causes  
repulsion.

This is the reason why CH<sub>4</sub> associates with CH<sub>4</sub>, due to London forces

**C<sub>5</sub>H<sub>12</sub> hydrophobic bonding:**



Hydrophobic bonding

n-Pentane has a boiling point of 35 °C; therefore, it is a liquid at room temperature - why is it a liquid? Because its temporary dipoles – it is not miscible in water – water would rather hydrogen bond to itself – like dissolves like.