

# Chem 241

## Lecture 24

# Announcement

Mistake we have class on the 3<sup>rd</sup> not 4<sup>th</sup>

Exam 3

Originally scheduled April 23<sup>rd</sup> (Friday)

What about April 26<sup>th</sup> (Next Monday)?

## APRIL/MAY

M	T	W	T	F	S	S
19	20	21	22	23	24	25
26	27	28	29	30	1	2
3	4	5	6	7	8	9
10	11	12	13			



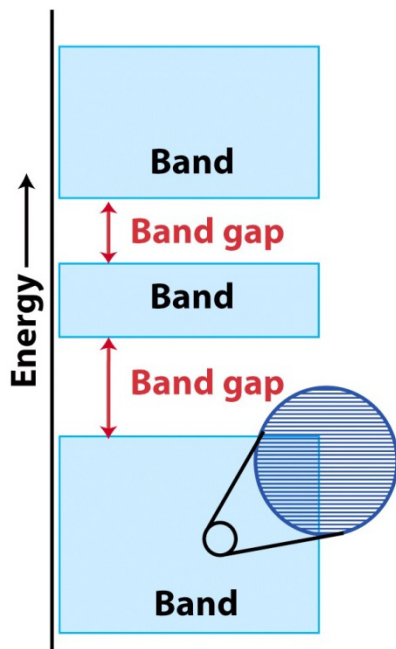
# Recap

Went over Exam 2  
Conductivity



# Inorganic solids

Treat solid like a very large molecule that has a nearly infinite number of atoms, and thus a nearly infinite number of bonding MOs and antibonding MOs. (Tight binding approximation).



- Atomic orbitals of the same type, say the valence s orbitals, give rise to a large number of molecular orbitals that are very similar in energy. This is a **band** of energy levels, the example would be an s-band.
- The bands are separated from each other by **band gaps**, which are ranges of energies that contain no MOs.
- Building up bands ( $\#$  of atoms =  $\#$  of orbitals)

# Inorganic solids

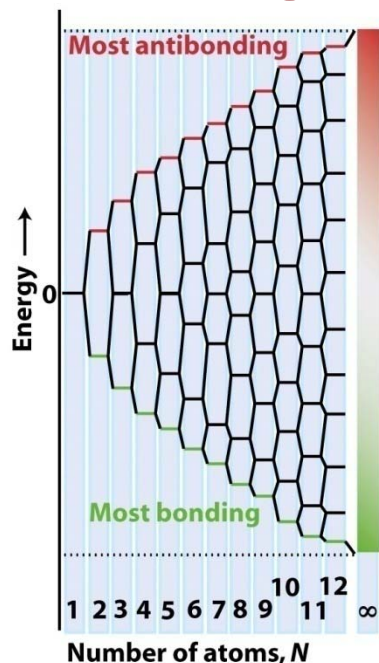


Figure 3-50  
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d. The result of carrying this out for a large number of atoms, is a large number of orbitals

e. The upper and lower energy of the band is constrained by the energy of the most bonding and most antibonding level.

f. since the energy range is finite, and the number of orbitals is the number of atoms in the solid, the band consists of a near-continuum of energy levels.

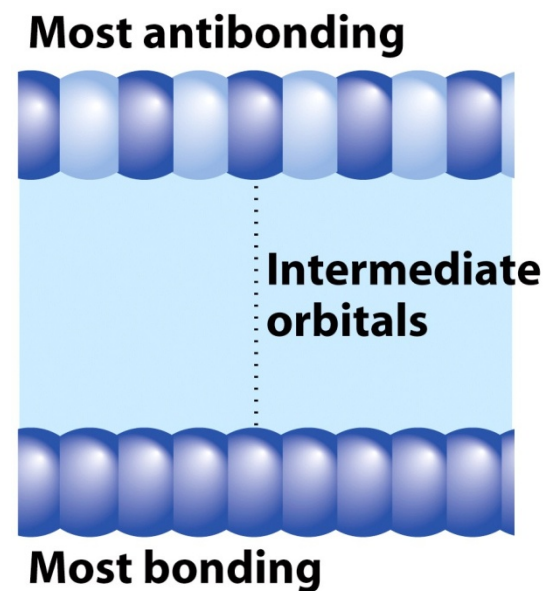
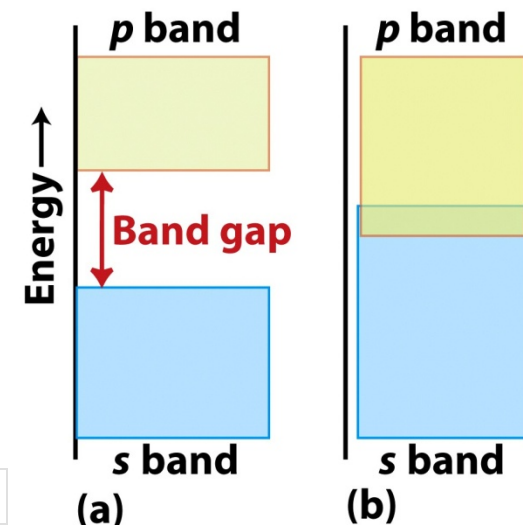
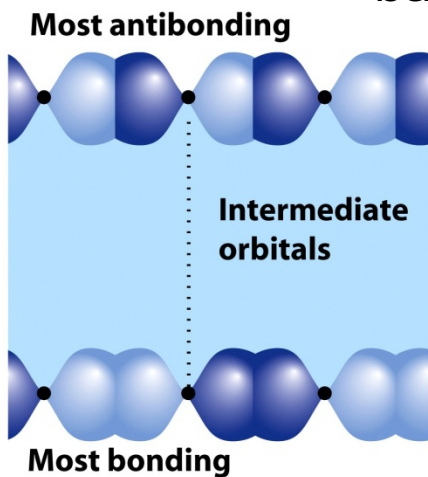


Figure 3-50

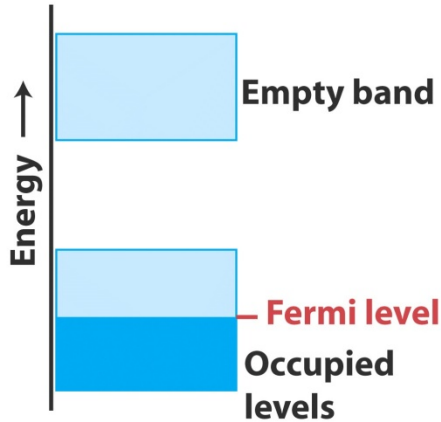
# Bands

- We can do the same for p-orbitals, d-orbitals, etc. to form p- and d-bands.
  - a. since atomic p-orbitals are higher in energy than s-orbitals, the p-band will lie above the s-band in energy by an amount that is determined by the difference in energy of the atomic orbitals.
  - b. the width (in energy) of a band is determined by the strength of the interaction between the orbitals. If the interaction is strong, the band will be wide. Sometimes the width of the band is sufficient for bands to overlap.



# Filling them up

Now we need to put electrons into the bands.

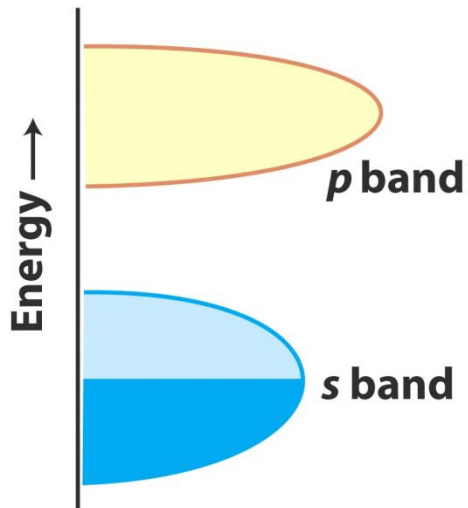


a. The highest level occupied in a band at  $T = 0$  K is the **Fermi level**. In a case where the diatomic would fill the bonding, but not the antibonding, the Fermi level is in the middle of the band.

b. **Conduction band** explains metallic conductors.

c. The ability of the electrons to move in the solid is dependent on the uniformity of the material (purity of the material all atoms the same).

d. the number of energy levels in a given energy range is the density of states,  $\rho$ , and is high in the middle of a band, and low at the edges. (0 in a band gap)



# Semimetals

In some cases the highest energy level of a filled band is the same as the lowest energy of an empty band (there is no band gap), but since the density of states at the top edge of the lower band and the lower edge of the upper band is very low. Such solids are semimetals (*e.g.*, graphite).

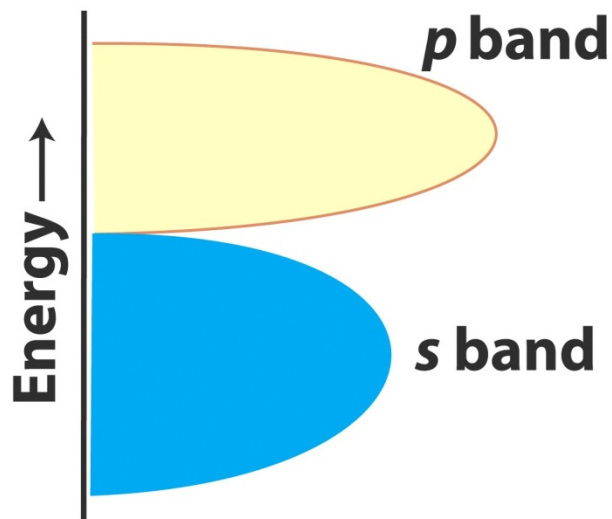


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# Insulators

- **Insulators** are materials where a filled band is separated in energy from an empty band by a large band gap.
- **Semiconductors** are materials that have small band gaps, where empty bands can be populated through thermal excitation of electrons from the filled band to the empty band.

Thus, insulators are just semiconductors with large band gaps.

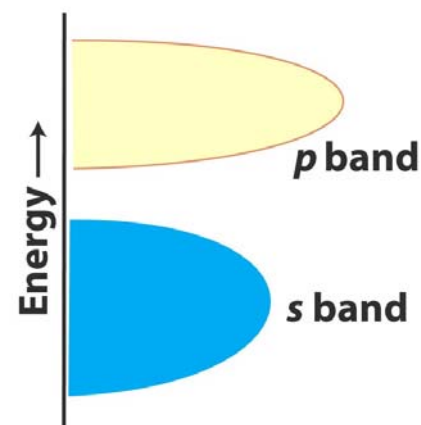


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# Intrinsic semiconductors

Bands in the pure material are close enough in energy to be thermally populated.

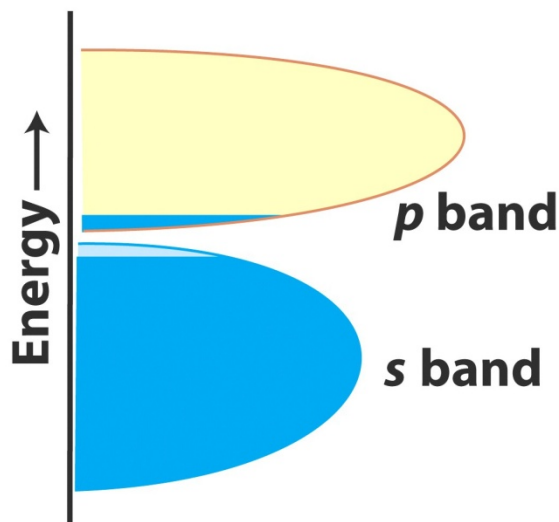


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- The promotion of electrons from the filled band into the empty band, introduces electrons into the empty band, and holes in the filled band.

- The population of the upper band has Boltzmann-like temperature dependence, and thus the conductivity shows an exponential dependence with temperature.

$$\sigma = \sigma_0 e^{-E_{\text{gap}}/2kT}$$

- This predicts an activation energy equal to  $\frac{1}{2}$  the band gap.

# Extrinsic Semiconductors

These are compounds where the semiconductor properties are the result of introducing impurities that produce either occupied donor bands (n-type) or empty acceptor bands (p-type) in the band gap of the pure material.

-the process is called doping

-very low levels of dopant (one atom per  $10^9$ ) are required.

- Ex1. If you dope a Si crystal ( $[\text{Ne}]3s^23p^2$ ) with As ( $[\text{Ar}]4s^24p^3$ ), you introduce one additional valence  $e^-$  per As. **n-type**

- Ex2. If you dope a Si crystal ( $[\text{Ne}]3s^23p^2$ ) with Ga ( $[\text{Ar}]4s^24p^1$ ), you introduce one additional valence  $e^-$  per As. **p-type**

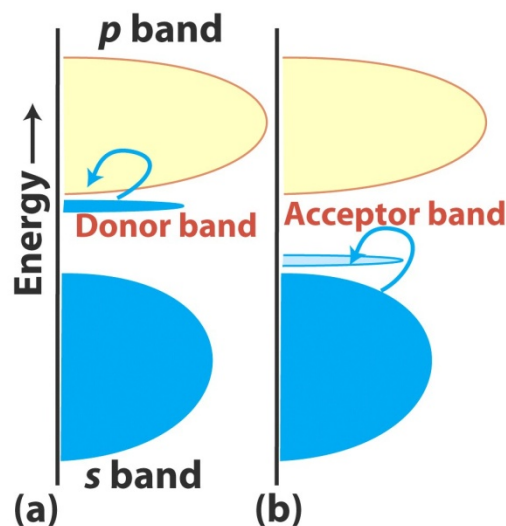


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# Homework

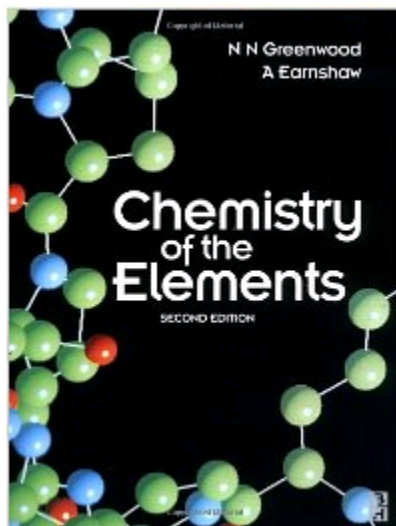
Chapter 3:

Exercises: 2, 4, 7, 9, 16, 17

Start reading chapter 9.



# The Groups



**Chemistry of the Elements**  
A. Earnshaw, Norman Greenwood  
**ISBN-10: 0750633654**



# Hydrogen

Does not fit neatly in the periodic table

A. Electronic Configuration:  $1s^1$

- Group 1 =  $ns^1$
- Group 17 = needs one more for full shell

3 Useful isotopes

$^1\text{H}$  – protium

$^2\text{H}$  – deuterium

$^3\text{H}$  – tritium



Used in kinetics isotope effect.

NMR – differ in spins

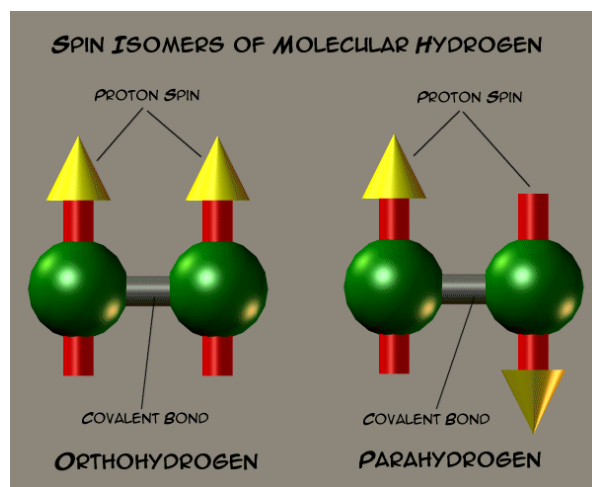


# Ortho-para hydrogen

Dihydrogen can be found in two forms.

*Ortho*-hydrogen spins are parallel

*Para*-hydrogen spins are anti-parallel



# Two ways to look at hydrogen

$H^+$  - proton - Acts like metals (Lewis Acid)

$H^-$  - hydride – Good nucleophile  
- metal hydrides –  $LiAlH_4$ ,  $NaH$ ,

When assign oxidation number if it is acting like a hydride it is -1 ( $AlH_3$ ) and when it is acting like a proton +1 ( $NH_3$ )

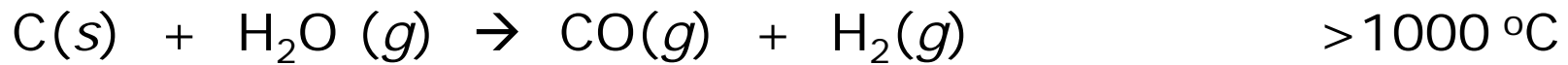
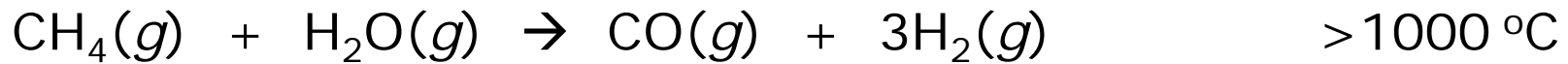
With metals it acts like hydride, more polar bond.

With non-metals it acts more covalent (but polar).

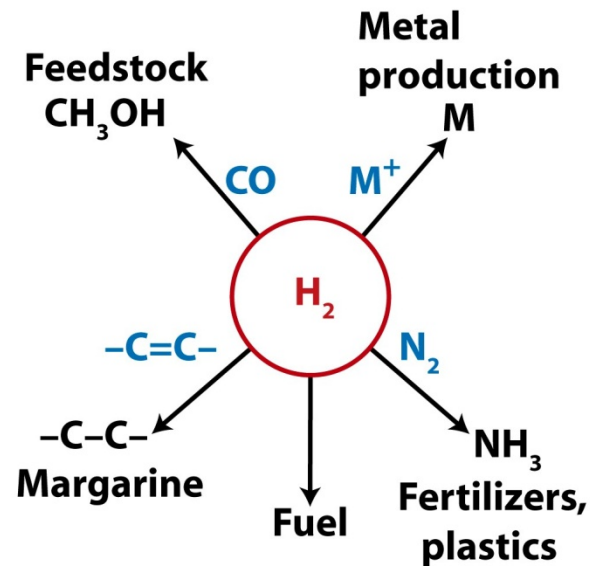
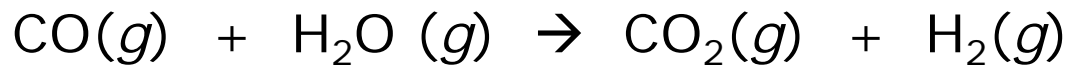




# Production and Use



Water- Gas Shift



# Reactions of Hydrogen

Hydrogen can be homolytically cleaved on the metal surfaces.

