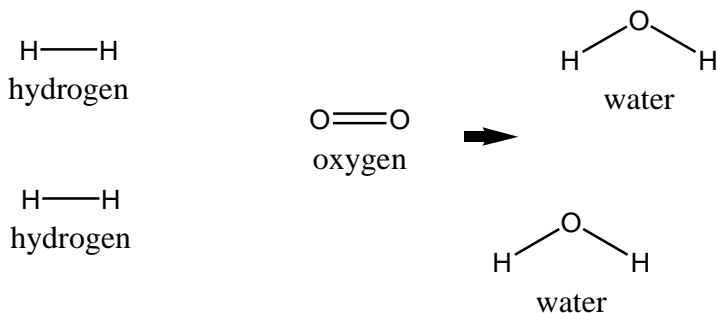
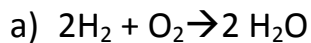
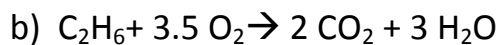


Exercises

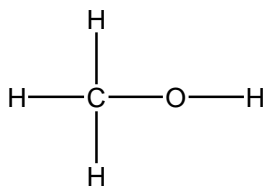
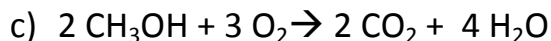
1. Estimate ΔH for each of the following using the given table of bond energies:



ΔH_{bb}	ΔH_{bf}	ΔH $= \Delta H_{\text{bb}} + \Delta H_{\text{bf}}$
$2(436)+498 = 1370 \text{ kJ}$	$2(2)(-464) = -1856$	$1370 \text{ kJ} - 1856 \text{ kJ} = -490 \text{ kJ}$



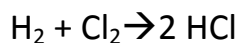
ΔH_{bb}	ΔH_{bf}	$\Delta H = \Delta H_{\text{bb}} + \Delta H_{\text{bf}}$
$6(414)+1(347) +3.5(499)=$	$2(2)(-745)+3(2)(-464)=$	$= - 1204 \text{ kJ}$



ΔH_{bb}	ΔH_{bf}	$\Delta H = \Delta H_{\text{bb}} + \Delta H_{\text{bf}}$
$2(3)\text{C}-\text{H} + 2(\text{O}-\text{C}) + 2(\text{O}-\text{H}) + 3 \text{O}=\text{O}$ $=2(3)(414) + 2(335) + 2(464) + 3(494)$	$2(2)(-745) + 4(2)(-464) =$	-1128kJ

2. a) Without consulting a table of bond energies, find the bond energy of

H-Cl on a per mole basis if H_2 's and Cl_2 's bond energies are 436 kJ/mole and 243 kJ/mole, respectively, and the ΔH for the following reaction is -185 kJ/mole of H_2



$$\Delta H = \Delta H_{\text{bb}} + \Delta H_{\text{bf}} = 436 + 243 + 2(-x) = -185, \text{ where } x = \text{bond energy of 1 mole of HCl}$$

$$x = 432 \text{ kJ/mole HCl}$$

b) Convert H_2 's bond energy to kJ/g

$$436 \text{ kJ/mole } \text{H}_2, \text{ but 1 mole of } \text{H}_2 = 2.02 \text{ g} = 436 \text{ kJ}/2.02 \text{ g} = 216 \text{ kJ/g}$$

c) Show an energy- reaction profile (energy diagram) for $\text{H}_2 + \text{Cl}_2 \rightarrow 2\text{HCl}$ in terms of ΔH_{bb} and ΔH_{bf}

3. Why is H---Cl 's bond energy greater than that of H---Br?

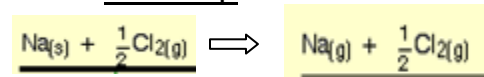
Hint: periodic trends.

Since Cl is more electronegative than Br, it will pull electrons from the bonded H closer to itself, which lowers its potential energy. Just like it takes more energy to raise something from a relatively lower shelf(compared to HBr) to a higher shelf, it will take more energy to then separate the H from the Cl. Hence HCl's bond energy is higher then HBr's.

4. a) Explain why the first 3 steps of the salt-producing reaction are endothermic.
 b) Why are the last two steps are exothermic?
 c) Use algebra to show that the overall reaction is indeed $\text{Na}_{(s)} + \frac{1}{2} \text{Cl}_{2(g)} \rightarrow \text{NaCl}_{(s)}$

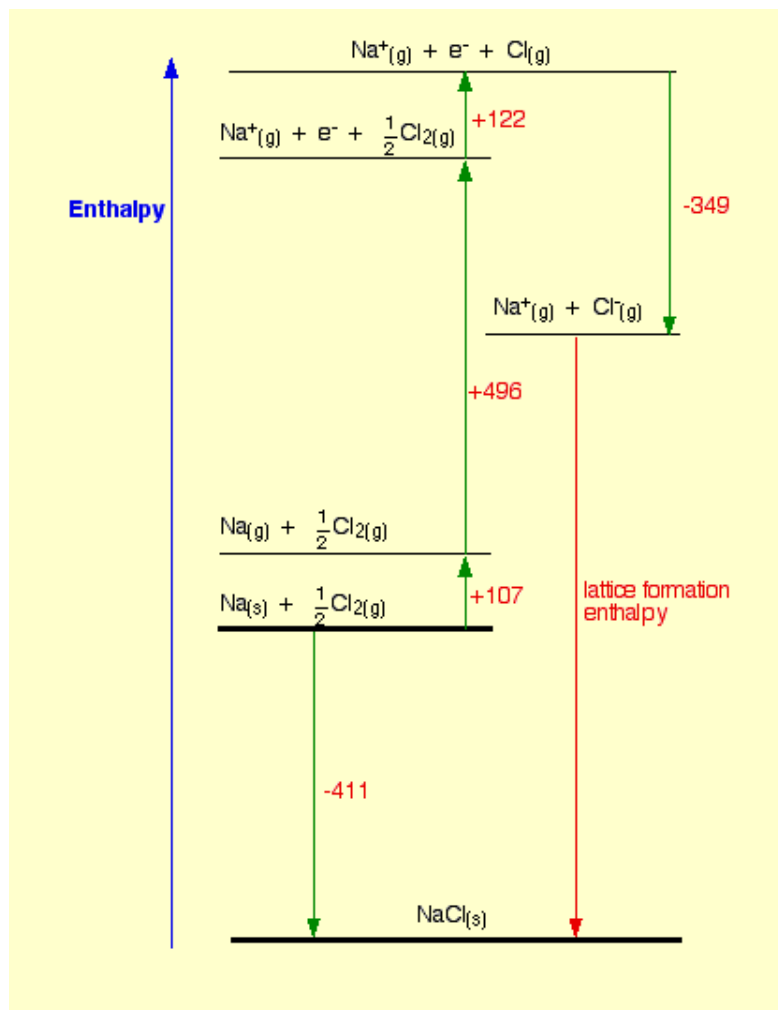
A)

• first step



The Na is sublimating; endo

- The second step: the sodium gas is getting ionized. The electron is being separated from the atom. (like ionization energy)

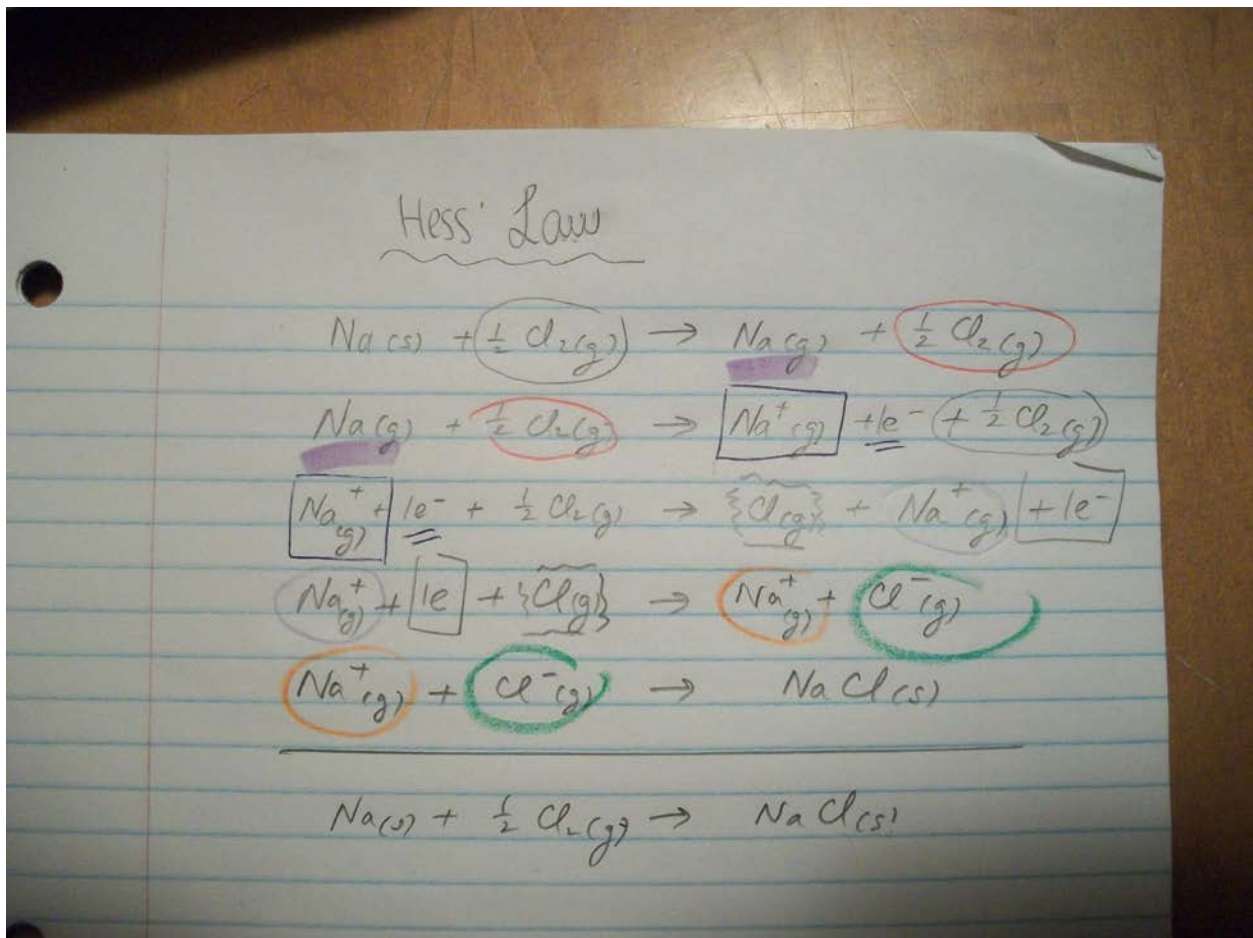


- Third step: Cl_2 's bond is being broken.

b)

- Fourth step: electron is being captured by chlorine (similar to bonding and opposite of ionization)
- Fifth step: the sodium and chloride ions are coming closer together and bonding

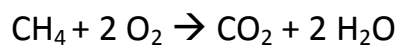
c)



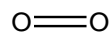
Average bond energies, kJ/mol:

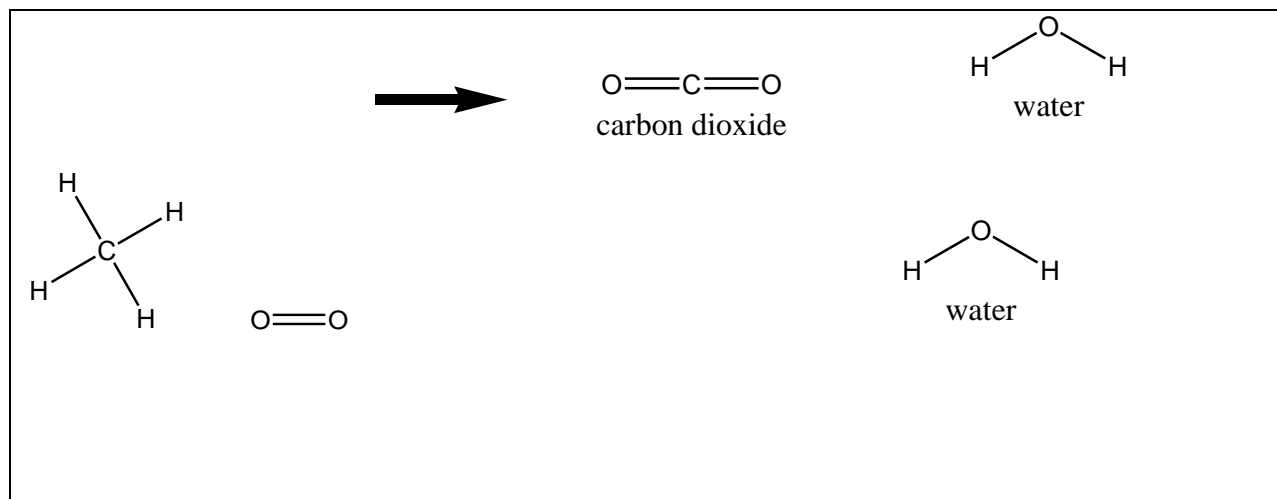
H—H	436	C—H	414	N—N	159	P—H	318
H—F	570	C—O	335	N=N	418	P—Cl	326
H—Cl	432	C=O	745	N≡N	941	S—H	339
H—Br	366	C≡O	1075	N—F	270	Cl—Cl	243
H—I	298	C—F	485	N—Cl	201	Br—Br	193
C—C	347	C—Cl	326	O—O	138	Br—Cl	218
C=C	619	C—Br	285	O=O	494	I—I	151
C≡C	812	C—I	239	O—H	464		
C—N	293	N—H	389	F—F	159		
C=N	515	N—O	175	F—Cl	256		
C≡N	891	N=O	590				

Example 2: Estimate the ΔH for the following reaction. Again consult the bond energies table on the previous page:



Lewis Structures





ΔH_{bb}	ΔH_{bf}	ΔH = $\Delta H_{\text{bb}} + \Delta H_{\text{bf}}$
$4(413) + 2(498) = 2648 \text{ kJ}$	$2(-745) + 4(-460) =$ -3330 kJ	$2648 \text{ kJ} - 3330$ $\text{kJ} = -680 \text{ kJ}$

Example 3 Show an energy- reaction profile(energy diagram) for an endothermic reaction in terms of ΔH_{bb} and ΔH_{bf}

Dalton's Law of Partial Pressures

If there are two or more gases in a container, the total pressure is the sum of the individual pressures of the various gases:

$$P_T = P_1 + P_2 + \dots$$

Consider two gases A and B in a container at the same temperature.

$P_T = P_A + P_B$ where P_A is the partial pressure of gas A (partial pressure is the gas' individual contribution to the total pressure)

If we know n_A and n_B , is there a way we could calculate P_A from P_T , n_A and n_B ?

$$P_A = (n_A/n_T)(P_T)$$

Use $PV = nRT$ to derive this formula.

$$P_A V = n_A RT$$

$$P_T V = n_T RT$$

Divide these two equations, simplify and isolate P_A :

$$\frac{P_A V}{P_T V} = \frac{n_A RT}{n_T RT}$$

$$\frac{P_A}{P_T} = \frac{n_A}{n_T}$$

$$P_A = \frac{n_A}{n_T} P_T$$

Example 1 While kept at a constant temperature, a gas mixture contains the following:

n	gas
0.340	H ₂
5.55	He
2.10	Ar

The manometer attached to the container containing the mixture reads 233 kPa. Find the partial pressure of each gas.

$$P_A = \frac{n_A}{n_T} P_T$$

$$P_{H_2} = \frac{n_{H_2}}{n_T} P_T$$

$$P_{H_2} = \left[\frac{0.34}{0.34 + 5.55 + 2.10} \right] 233 \text{ kPa}$$

$$= 9.9 \text{ kPa}$$

Repeat for other gases; $P_{He} = 162 \text{ kPa}$; $P_{Ar} = 61 \text{ kPa}$

Example 2 Since equal volumes of ideal gases contain the same number of moles under the same conditions of P and T, % volumes of gases are directly proportional to mole fractions.

With this in mind, find the partial pressure of oxygen in air at STP.

Air is 21% O₂ by volume.

$$n_{O_2} = 0.21$$

$$P_T = 101.3 \text{ kPa at STP}$$

$$P_A = \frac{n_A}{n_T} P_T$$

$$P_{O_2} = \frac{n_{O_2}}{n_T} P_T = \frac{0.21}{1} 101.3 = 21 \text{ kPa}$$

Exercises

1. A balloon contains 0.100 moles of oxygen and 0.400 moles of nitrogen. If the balloon is at standard temperature and pressure, what is the partial pressure of the nitrogen?

$$P_{N_2} = \frac{n_{N_2}}{n_T} P_T = \frac{0.400}{0.100 + 0.400} 101.3 = 81.0 \text{ kPa}$$

2. The pressure of a mixture of nitrogen, carbon dioxide, and oxygen is 150.0 kPa. What is the partial pressure of oxygen if the partial pressures of the nitrogen and carbon dioxide are 100.0 kPa and 24.0 kPa, respectively?

$$\begin{aligned} P_T &= P_{N_2} + P_{O_2} + P_{CO_2} \\ 150.0 &= 100.0 + 24.0 + P_{CO_2} \\ P_{CO_2} &= 26.0 \text{ kPa} \end{aligned}$$

3. A gaseous mixture made from 10.0 g of oxygen and 15.0 g of argon is placed in a 8.00 L vessel at 25.2°C. What is the partial pressure of each gas, and what is the total pressure in the vessel?

$$10.0 \text{ g O}_2 / (32.0 \text{ g/mole}) = 0.3125 \text{ mole O}_2$$

$$15.0 \text{ g Ar} / (39.9 \text{ g/mole}) = 0.376 \text{ mole Ar}$$

$$P_T V = n_T R T$$

$$P_T (8.00) = (0.3125 + 0.376) * 8.31 (273 + 25.2)$$

$$P_T = 213 \text{ kPa}$$

$$P_{O_2} = \frac{0.3125}{0.376} 213.26 = 96.8 \text{ kPa}$$

$$P_{Ar} = P_T - P_{O_2} = 213 - 96.8 = 116 \text{ kPa}$$

4. a) 0.888 L of “wet” oxygen (this implies that it’s mixed with water vapour) are collected at a temperature of 25.0 °C. The total pressure of the gases is 99.8 kPa. What is the partial pressure of the dry O₂? The partial pressure of water at 25.0° C is 3.17 kPa.

b) How many grams of water are in the 0.888 L mixture?

$$a) P_{O_2} = P_T - P_{H_2O} = 99.8 - 3.17 = 96.6 \text{ kPa}$$

$$b) P_T V = n_T R T$$

$$99.8 (0.888) = n_T (8.31) (25 + 273)$$

$$n_T = 0.03578 \text{ moles}$$

$$P_{H_2O} = \frac{n_{H_2O}}{n_T} P_T$$

$$3.17 = \frac{n_{H_2O}}{0.03578} 99.8$$

$$n = 0.001136 \text{ moles of water} (18 \text{ g/mole}) = 0.0204 \text{ g of water}$$