

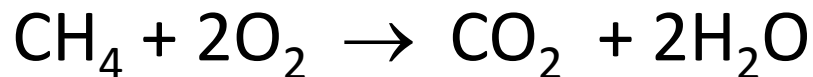
# Standard Enthalpy, Hess, and Free Energy

# Standard Enthalpy

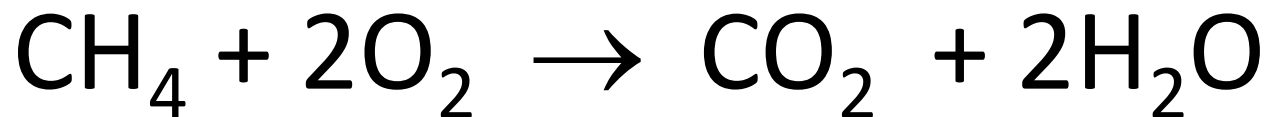
- In order to compare one thing to another, there must be a “standard”
- Standard temperature, 25°C.
- Enthalpy changes during a reaction at 25°C are considered the standard enthalpy.
- $\Delta H_f^\circ$  = standard enthalpy of formation  
(found in book)
- Standard enthalpies of formation for elements in their common form at 25°C is 0kJ/mol
- Moles (mol) is just an amount for now (like puffs).

# Standard Enthalpy of Formation

- Look up the enthalpies on a table
- $\Delta H = H_{\text{Products}} - H_{\text{reactants}}$



	Substance	# of Moles	$\Delta H_f$ kJ/mole
Reactants	CH <sub>4</sub>	1	-74.8
	O <sub>2</sub>	2	0
Products	CO <sub>2</sub>	1	-393.5
	H <sub>2</sub> O	2	-285.8



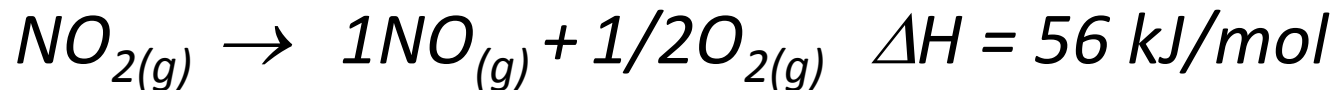
	Substance	# of Moles	$\Delta H_f$ kJ/mole
Reactants	$\text{CH}_4$	1	-74.8
	$\text{O}_2$	2	0
Products	$\text{CO}_2$	1	-393.5
	$\text{H}_2\text{O}$	2	-285.8

$$\begin{aligned}\Delta H_f &= [-393.5 + 2(-285.8)] - [-74.8 + 2(0)] \\ &= -890.3 \text{ kJ/mol}\end{aligned}$$

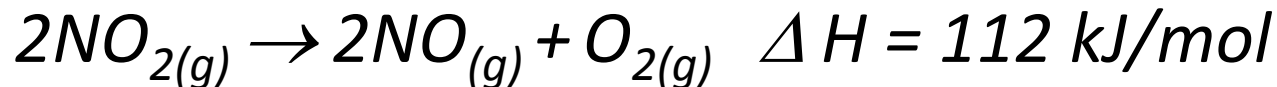
# Hess's Law

- Enthalpy is a “state” function
  - It doesn't matter how you get there, the energies are the same
- Reaction can be broken down into a series of steps.
  - We can add, subtract, multiply, reverse reaction enthalpies for each step
  - To find the overall enthalpy change

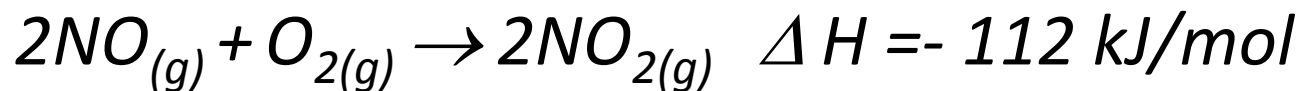
# Hess' Law or State Functions



*Double the reaction = double the enthalpy*

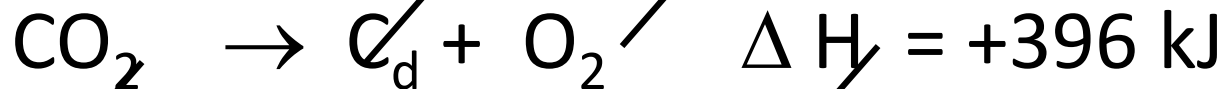
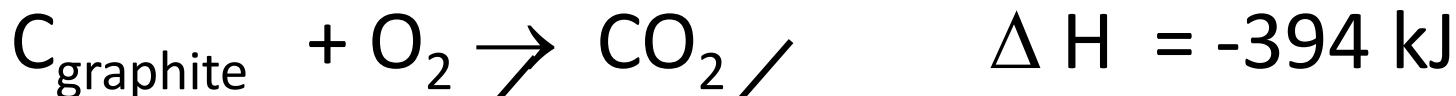
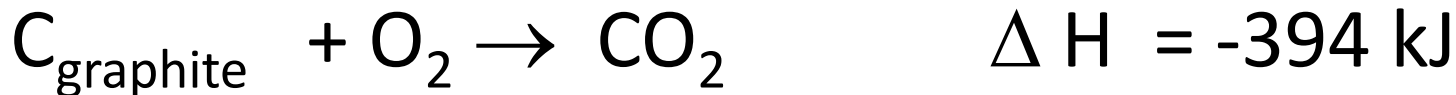
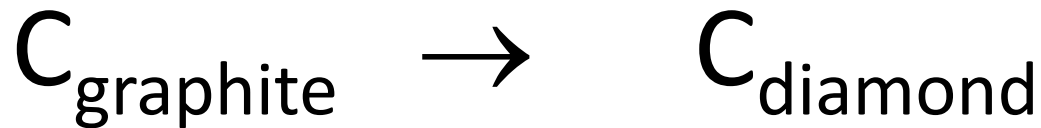


*The reverse of the reaction, "negative"*



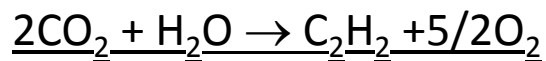
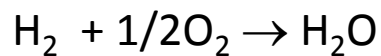
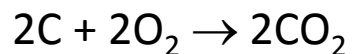
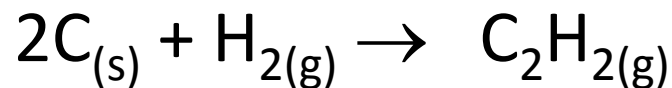
# Hess's Law

- $2\text{H}_2\text{O}_2 \rightarrow 2\text{H}_2\text{O} + \text{O}_2$        $\Delta\text{H} = -190\text{kJ}$
- $4\text{H}_2\text{O}_2 \rightarrow 4\text{H}_2\text{O} + 2\text{O}_2$        $\Delta\text{H} = -380\text{kJ}$
- $2\text{H}_2\text{O} + \text{O}_2 \rightarrow 2\text{H}_2\text{O}_2$        $\Delta\text{H} = +190\text{kJ}$
- $4\text{H}_2\text{O} + 2\text{O}_2 \rightarrow 4\text{H}_2\text{O}_2$        $\Delta\text{H} = +380\text{kJ}$
- $1\text{H}_2\text{O}_2 \rightarrow 1\text{H}_2\text{O} + \frac{1}{2}\text{O}_2$        $\Delta\text{H} = -95\text{kJ}$

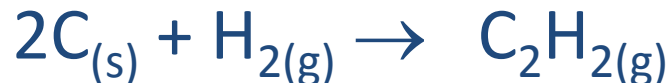




Calculate the  $\Delta H$



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# Spontaneity

- Some reactions happen on presumably on their own under certain conditions.
- These reactions are called spontaneous reactions.
- There are multiple factors that go into whether a reactions will be spontaneous or not.
- These are
  - Change in Enthalpy
  - Temperature
  - Change in Entropy

# Entropy

- Entropy is referring to disorder of the substance.
- The higher the entropy, the more the disorder.
- For example:
  - a gas has a higher entropy than a liquid.
  - Aqueous solutions have more entropy than solid compounds.
- Nature moves toward raising entropy. This is why:
  - A gas will spread out in a room
  - It takes so much energy for your body to stay together.

# Gibb's Free Energy

- Gibb's free energy is a way to measure the spontaneity of a reaction.
- It is denoted by the symbol  $\Delta G$ .
- If  $\Delta G$  is negative, then the reaction is spontaneous.
- If  $\Delta G$  is positive, then the reaction is not spontaneous.
- The equation that relates this is:

$$\Delta G = \Delta H - T \Delta S$$

- $\Delta S$  is the change in entropy,  $T$  is temperature, and  $\Delta H$  is the change in enthalpy.

# Is it Spontaneous?

- If the enthalpy is negative (meaning the bonds in the products are stronger than the reactants) then it is more likely to be spontaneous.
- If the entropy is positive (meaning the reaction moves toward disorder) then it is likely to be spontaneous.
- If the temperature is high, then the reaction is more likely to be spontaneous as long as the entropy is positive.
- The opposite will decrease the probability of a spontaneous reaction.