CH3 (O CH3 (l) + 402(g) -> 3 (O2(g) + 3H20(l); AH = -1800 KJ

What would be the enthapy change when 25 g of water are produced by the reaction?

- 1 Convert 25 grams water to moles. Use FORMULA WEIGHT.
- 2 Convert moles water to enthalpy change. Use THERMOCHEMICAL EQUATION.

H₂0: H: 2x | .008
0:
$$\frac{1}{18.016}$$
 $\frac{1}{18.016}$ $\frac{1}{18.01$

This is an EXOTHERMIC reaction. 830 kJ are transferred to the SURROUNDINGS. The enthalpy change here is the same as Q, as long as we do a constant pressure process.

A few more terms related to enthalpy:

- Enthalpy of vaporization / heat of vaporization: The enthalpy change on vaporizing one mole of a substance. (from liquid to vapor)
- Enthalpy of fusion / heat of fusion: The enthalpy change when a mole of liquid changes to the solid state.



- A reaction that forms exactly one mole of the specified substance from its elements at their STANDARD STATE at 25C and 1 atm pressure.

(O2(g):
$$(s,graphite) + O2(g) \rightarrow (O2(g);\Delta H = -393.5 kJ)$$

heat of formation of carbon dioxide ΔH_{f}^{o} or ΔH_{f}
"enthal py of formation"
 $(O(g): (s,graphite) + \frac{1}{2}O2(g) \rightarrow (O(g);\Delta H = -110.5 kJ)$

you may see fractional coefficients in these formation reactions, because you MUST form exactly one mole of the product!

- The heat of formation for an element in its standard state at 25C and 1 atm is ZERO.

- What are formation reactions good for?

... finding enthalpies for more interesting reactions!

Let's say we would like to find the enthalpy of reaction for this equation: $2(2H_2(g) + 5O_2(g) \longrightarrow 4(O_1(g) + 2H_2O(g); \Delta H = 1)$ Hess' Law: If you add two reactions to get a new reaction, their enthalpies also add. DH = 226.7 (242: 2((s) + H2(g) -> (2H2(g)) (O2 (g) DH=-393,5 CO2 1 $((s) + O_2(y) \longrightarrow$ DH= -241,8 H20(g) $H_2(g) + \frac{1}{2}O_2(g)$ H20 1 $\rightarrow 2(s) + H_2(s)$ (2H2lg) DH = -226.7 \rightarrow 2((s) + H_2 (g) (2H2lg) DH = -226.7 $((s) + O_2(g))$ $\rightarrow (O_2(g))$ DH = -393.5 ((5) + O2 (g) $\longrightarrow (O_2(q)$ DH = -393,5 $((s) + O_2(g))$ $\longrightarrow (O_2(q)$ DH = -393,5 $\longrightarrow (O_2(q)$ DH = -343.5 $H_2(q) + \frac{1}{2}O_2(q)$ H20 (9) DH= -241,8 $2(q) + \frac{1}{2}O_2(q)$ H20 (9) DH= -241,8

$$2(2H_2(g) + So_2(g) \longrightarrow 4co_2(g) + 2H_2o(e)$$

Hess' Law using enthalpy of formation:

$$\Delta H = \sum_{0} \Delta H_{f,pNJNLts} - \sum_{0} \Delta H_{f,renchnts}$$

$$\frac{226.7}{2(2H_{2}(g) + 50_{2}(g))} \longrightarrow \frac{-393.5}{4(0_{2}(g))} + \frac{241.8}{420(g)}; \Delta H = \frac{2}{5}$$

$$\Delta H = \left[4(-393.5) + 2(-241.6) \right] - \left[2(226.7) + 5(0) \right]$$

$$= -2511 \text{ J}$$

See Appendix C in the textbook for enthalpy of formation data: p A-8 to A-11

* Remember:

- Multiply each enthalpy by its stoichiometric coefficient from the reaction
- Enthalpy of formation of an element at its standard state is zero
- Watch phase labels. You will usually find SEVERAL enthalpies of formation for a given substance in different phases!
- For ionic substances in solution, remember that they exist as free ions, so look up the aqueous IONS!

$$2.016$$
 32.00 18.02 in purple $2.12(g) + O_2(g) \longrightarrow 2.12O(g)$; $\Delta H = -4.84 \text{ kJ}$

Calculate the enthalpy change for the combustion of 1.00 kg of hydrogen gas.

- 1 Convert 1.00 kg hydrogen gas to moles. Use formula weight.
- 2 Convert moles hydrogen gas to enthalpy change. Use thermochemical equation.

2 2mol H2 = -484 KJ

1.00 kg H₂ x
$$\frac{10^3g}{kg}$$
 x $\frac{nul H_2}{2.016g H_2}$ x $\frac{-484 kJ}{2 mel H_2} = \frac{-120000 KJ}{per kg H_2}$

What is the enthalpy change when 150. L of nitrogen monoxide are formed by this reaction at 25.0 C and 1.50 atm pressure?

- 1 Convert 150. L NO to moles. Use IDEAL GAS EQUATION.
- 2 Convert moles NO to enthalpy change. Use THERMOCHEMICAL EQUATION.