One problem ...

PATH. The amount of energy required for a process depends on how the process is carried out.

Example: Driving from Florence to Columbia. How much energy is required? (gas)

Jeep Cherokee vs Toyota Prius. The Jeep will use much more fuel than the Prius even though they start and end from exactly the same place. So the fuel usage is what we call a <u>PATH FUNCTION</u>, while the location is a STATE FUNCTION.

- so the heat of reaction depends on how the reaction is done.

- we need (for reporting) some kind of standard condition. At constant pressure, we can define a state function called ENTHALPY (H)

H = U + PV $\bigwedge H = Q$ constant pressure

 ΛH_{r}

... we record the "enthalpy change of reaction" in our data books.

SINCE the enthalpy change does NOT depend on path, this means that we can use standard values for enthalpy to predict the heat change in reactions that we have not tested in a calorimeter.

THERMOCHEMICAL EQUATIONS

- is like a regular chemical equation, except that phase labels are REQUIRED and the enthalpy for the reaction is given along with the equation.

$$CH_3(O(H_3(l) + 4O_2(g) \rightarrow 3(O_2(g) + 3H_2O(l); AH = -1800 kJ$$

- Why are phase labels required? Because phase changes either absorb or release energy.

 $\Delta H = -1600 \text{ kJ} \dots \text{ what does this mean}?$

 $1 \mod CH_{3}COCH_{3} = -1800 \text{ kJ}$ $4 \mod O_{2} = -1800 \text{ kJ}$ $3 \mod CO_{2} = -1800 \text{ kJ}$ $3 \mod H_{2}O = -1800 \text{ kJ}$

We treat the enthalpy change as if it's another product of the reaction!

USING A THERMOCHEMICAL EQUATION

 $CH_3(O(H_3(l) + 4O_2(g) \longrightarrow 3(O_2(g) + 3H_2O(l); AH = -1800 \text{ kJ})$ What would be the opthapy chapter when 25 a of water are produced by the reaction?

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

Convert 25 grams of water to moles. Use FORMULA WEIGHT.
 Convert moles of water to enthalpy change. Use THERMOCHEMICAL EQUATION.

$$\begin{array}{c} 1 \\ H_{20}; H = 27|.008 \\ 0 - \frac{1 \times 16.00}{1 \times 16.00} \\ 18.016gH_{20} = m_{0}| H_{20} \\ 18.016gH_{20} = m_{0}| H_{20} \\ 2S_{g}H_{20} \times \frac{m_{0}| H_{20}}{18.016gH_{20}} \times \frac{-1800 kJ}{3m_{0}| H_{20}} = -830 kJ \\ \hline 1 \end{array}$$

Notes:

A) The reaction is exothermic. We know that because the enthalpy change's sign is negative.

B) As long as this acetone is being burned at constant pressure, the enthapy change is equal to the heat, Q. We would expect 830 kJ of energy to be released as heat by this reaction.

FORMATION REACTIONS

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

$$(O_{2}(g): ((s, graphite) + O_{2}(g) \rightarrow (O_{2}(g)); AH = -\frac{393.5 \text{ kJ}}{(s, graphite)} + O_{2}(g) \rightarrow (O_{2}(g)); AH = -\frac{393.5 \text{ kJ}}{(s, graphite)}$$

$$(O_{2}(g): ((s, graphite) + O_{2}(g) \rightarrow (O_{2}(g)); AH = -\frac{190.5 \text{ kJ}}{(s, graphite)} + \frac{1}{2}O_{2}(g) \rightarrow (O_{2}(g)); AH = -\frac{110.5 \text{ kJ}}{(s, graphite)}$$

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.

 $AH_{f}^{\circ}, O_{2}(y) = O kJ/mol$

- What are formation reactions good for?

... finding enthalpies for more interesting reactions!

³² Let's say we would like to find the enthalpy of reaction for this equation: $2C_2H_2(g) + 5O_2(g) \longrightarrow 4(O_2(g) + 2H_2O(g); \Delta H = .$						
Hess' Law: If you add two reactions to get a new reaction, their enthalpies also add.						
		2(s) +			(2H2lg)	54=227.4
	C02 1	(1)	$O_2(y)$	\longrightarrow	$(O_2(g))$	14 = - 393.5
	H20 '	$H_2(g) + \frac{1}{2}$			$H_2O(g)$	DH= -241,8
	(2)	12 (g)	-)	2(s)	+ H2(y)	04 = -227.4
	C2	$H_2(g)$)	2 ((s)	+ 42 (g)	04 = -227.4
	Сc	s_{3} + $O_{2}(y)$	\rightarrow	(0 ₂ (g)	$\Delta H = -393.5$
	· · · · /	$(5) + O_2(y)$	\rightarrow	(02 (g)	$\Delta H = -393.5$
	(<u> </u>	$5) + O_2(y)$	\rightarrow	$lo_2($)	AH = -393.5
	Ĺ ι	$\mathfrak{S} + \mathcal{O}_2(\mathfrak{g})$	\rightarrow	$lo_2(g$)	$\Delta H = -393.5$
	42	$(q) + \frac{1}{2}O_2(q)$	\rightarrow		$H_2O(g)$	Aug - 744 6
	W2 ($(a) + \frac{1}{2} O_2(a)$	\rightarrow		$H_2O(a)$	ДН: -241,8 ДН: -241,8
	12	.]). 2. 6.97				14: 241,8
$2(2H_2(g) + SO_2(g) \longrightarrow 4CO_2(g) + 2H_2O(l)$						
$\Delta H = 2(-227.4) + 4(-393.5) + 2(-241.8) = -2512.4 \text{ kJ}$						

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* Enthalpy of formation data taken from Openstax Chemistry Appendix G

Hess' Law using enthalpy of formation:

2512.4kJ

 $\Delta H = \sum \Delta H_{F, products} - \sum \Delta H_{F, reacharts}$ $227.4 \qquad -393.5 \qquad -241.8$ $2(_2H_2(g) + 5O_2(g) \longrightarrow 4(O_2(g) + 2H_2O(g); \Delta H = ?$ $\Delta H = [4(-393.5) + 2(-241.8)] - [2(_{27}4) + 5(0)]$

* Remember:

See Appendix G in the Openstax textbook for enthalpy of formation data! - 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! ¹³⁴ Example problems:

$$2H_{2}(g) + O_{2}(g) \longrightarrow 2H_{2}O(g) ; \Delta H = -484 kJ$$

FORMULA WEIGHTS in g/mol

Calculate the enthalpy change for the combustion of 1000 g of hydrogen gas.

Convert 1000 g of hydrogen gas to moles. Use FORMULA WEIGHT.
 Convert moles hydrogen gas to enthalpy change. Use THERMOCHEMICAL EQUATION.

$$\frac{1}{2} 2 \log H_2 = \max H_2 \qquad (2) \sum \min H_2 = -484 kJ$$

$$\frac{1000 \cdot g H_2 \times \frac{mu}{2001 \log H_2} \times \frac{-484 kJ}{2mu} = -120000 kJ \text{ per kg H}_2$$