## Representing Enthalpy Changes

## Thermochemical Equations

* A balanced chemical equation that includes the heat transfer
$2 \mathrm{H}_{2(g)}+\mathrm{O}_{2(g)} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(1)+571.6 \mathrm{~kJ}$
$\mathrm{MgCO}_{3(\mathrm{sl}}+117.3 \mathrm{~kJ} \rightarrow \mathrm{MgO}_{(s)}+\mathrm{CO}_{2(g)}$
* To write a thermochemical equation, you must know:
* the balanced chemical equation
* heat of reaction for the substance


## Thermochemical Equations

* You can also $\triangle H$ values in thermochemical equations

$$
\begin{array}{ll}
2 \mathrm{H}_{2(g)}+\mathrm{O}_{2(g)} \rightarrow 2 \mathrm{H}_{2} \mathrm{OH}_{\mathrm{H}} & \Delta \mathrm{H}_{\text {rxn }}=-571.6 \mathrm{~kJ} \\
\mathrm{MgCO}_{3(\mathrm{~s})} \rightarrow \mathrm{MgO}_{(s)}+\mathrm{CO}_{2(g)} & \Delta \mathrm{H}_{\mathrm{rxn}}=117.3 \mathrm{~kJ}
\end{array}
$$

* Remember:
* exothermic reactions have $-\Delta H$
* endothermic reactions have $+\Delta H$


## Potential Energy Diagrams

* exothermic reactions: products are lower in energy than reactants lenergy lost to surroundings)
* endothermic reactions: products are higher in energy than reactants lenergy absorbed from surroundings)




## Homework

* Pg 295 \# 7-10

Molar Enthalpy

## First Law of Thermodynamics

* The first law of thermodynamics states that energy can be converted from one form to another but cannot be created or destroyed

$$
E_{\text {system }}=-E_{\text {surroundings }}
$$

## Enthalpy

* Enthalpy $(\triangle \boldsymbol{H})$ : is the measure of the total energy of a thermodynamic system plus pressure times volume

$$
\text { * } \triangle H=\triangle E+\triangle(P V)
$$

Most reactions we study we assume $\triangle P V$ is zero, so for our purposes $\triangle \boldsymbol{H}=\triangle \mathbf{E}=\mathbf{Q}$

# Second Law of Thermodynamics 

* The second law of thermodynamics states that when two objects are in close contact, heat is always transferred from the object of a higher temperature to an object of lower temperature until the two objects are the same temperature (thermal equilibrium)



## Molar Enthalpy $\left(\Delta H_{x}\right)$

* Molar Enthalpy: is the enthalpy change associated with a physical, chemical, or nuclear change involving one mole of a substance.


## Molar Enthalpy $\left(\Delta H_{x}\right)$

* A thermochemical equation represents the energy change that accompanies a chemical reaction.

$$
2 \mathrm{H}_{2(g)}+\mathrm{O}_{2(g)} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}_{(g)}+483.6 \mathrm{~kJ}
$$

## Molar Enthalpy $\left(\Delta H_{x}\right)$

* The equation representing the energy released when ONE mole of $\mathrm{H}_{2}$ combusts would be

$$
\mathrm{H}_{2(\mathrm{~g})}+\mathrm{y}_{2} \mathrm{O}_{2(\mathrm{~g})} \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{~g})+241.8 \mathrm{~kJ}
$$

# Examples of Molar Enthalpies of Reactions 

|  |  |
| :---: | :---: |
| Solution ( $\triangle \mathrm{H}_{\text {sol }}$ ) |  |
| Combustion ( $\triangle$ Hcomb) | $\mathrm{CH}_{4(g)}+2 \mathrm{O}_{2(\mathrm{~g})} \rightarrow \mathrm{CO}_{2(g)}+\mathrm{H}_{2} \mathrm{O}(1)$ |
| Vaporization ( $\triangle$ Hvap) | $\mathrm{CH}_{3} \mathrm{OH}_{(1)} \rightarrow \mathrm{CH}_{3} \mathrm{OH}_{(\mathrm{g})}$ |
| Freezing ( $\triangle H_{\text {fr }}$ ) | $\mathrm{H}_{2} \mathrm{O}(1) \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{s})$ |
| Neutralization ( $\triangle$ \#neut) | $\mathrm{NaOH}_{\text {(aq) }}+1 / 2 \mathrm{H}_{2} \mathrm{SO}_{4 \text { (aq) }} \rightarrow 1 / 2 \mathrm{Na}_{2} \mathrm{SO}_{4}($ aq) $)+\mathrm{H}_{2} \mathrm{O}(1)$ |
| Formation ( $\triangle H_{f}$ ) | $\mathrm{C}(\mathrm{s})+\mathrm{H}_{2(\mathrm{~g})}+1 / 2 \mathrm{O}_{2(\mathrm{~g})} \rightarrow \mathrm{CH}_{3} \mathrm{OH}_{(1)}$ |

* The amount of energy involved in a change depends on the quantity of matter undergoing that change.
* To calculate an enthalpy change, $\Delta H(k J)$, for some amount other than a mole, use the formula

$$
\Delta H=n \Delta H_{X}
$$



$$
n=\# \text { of moles }
$$

## $\Delta H_{x}=$ molar enthalpy (kJ/mol)

## Example

$* \mathrm{H}_{2} \mathrm{O}(\mathrm{l})+40.8 \mathrm{~kJ} \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$

* Thus the molar enthalpy of vaporization for water is $\Delta H_{\text {vap }}=40.8 \mathrm{~kJ} / \mathrm{mol}$
* How much energy is needed to boil 90.10 g of water?
* Given:
* $\Delta H_{\text {vap }}=40.8 \mathrm{~kJ} / \mathrm{mol}$
* $m=90.10 \mathrm{~g}$
* $\mathrm{MH}_{\mathrm{H}_{2} \mathrm{O}}=18.02 \mathrm{~g} / \mathrm{mol}$


## * Given:

* $\Delta H_{\text {vap }}=40.8 \mathrm{~kJ} / \mathrm{mol}$
* $m=90.10 \mathrm{~g}$
* $\mathrm{M}_{\mathrm{H} 20}=18.02 \mathrm{~g} / \mathrm{mol}$


## * Required:

* $\Delta H$ ( and therefor $n$ )


## * Given:

* $\Delta H_{\text {vap }}=40.8 \mathrm{~kJ} / \mathrm{mol}$
* $m=90.10 \mathrm{~g}$
* $\mathrm{M}_{\mathrm{H}_{2} \mathrm{O}}=18.02 \mathrm{~g} / \mathrm{mol}$


## * Required:

* $\Delta H$ ( and therefor $n$ )


## * Equation:

* $n=m / M$ and $\Delta H=(n)(\Delta H x)$


## * $n=m / M$

* $n=(90.10 \mathrm{~g}) /(18.02 \mathrm{~g} / \mathrm{mol})$
* $n=5.00 \mathrm{~mol}$


## * $\Delta H=(n)(\Delta H x)$

* ( 5.00 mol ) ( $40.8 \mathrm{~kJ} / \mathrm{mol}$ )
* $=204 \mathrm{~kJ}$

Therefore 204 kJ of energy is required to boil 90.10 g of water.

## Example

* A common refrigerant (Freon-1 2, molar mass $120.91 \mathrm{~g} / \mathrm{mol}$ ) is alternately vaporized in tubes inside a refrigerator, absorbing heat, and condensed in tubes outside the refrigerator, releasing heat. This results in energy being transferred from the inside to the outside of the refrigerator. The molar enthalpy of vaporization for the refrigerant is $34.99 \mathrm{~kJ} / \mathrm{mol}$. If 500.0 g of the refrigerant is vaporized, what is the expected enthalpy change $\Delta H$ ?
* Given:
* $\Delta H_{\text {vap }}=34.99 \mathrm{~kJ} / \mathrm{mol}$
* $m=500.0 \mathrm{~g}$
* $M=120.91 \mathrm{~g} / \mathrm{mol}$


## * Given:

* $\Delta H_{\text {vap }}=34.99 \mathrm{~kJ} / \mathrm{mol}$
* $m=500.0 \mathrm{~g}$
* $M=120.91 \mathrm{~g} / \mathrm{mol}$


## * Required:

* $\Delta H$ ( and therefor $n$ )


## * Given:

* $\Delta H_{\text {vap }}=34.99 \mathrm{~kJ} / \mathrm{mol}$
* $m=500.0 \mathrm{~g}$
* $\mathrm{M}=120.91 \mathrm{~g} / \mathrm{mol}$


## * Required:

* $\Delta H$ ( and therefor $n$ )


## * Equation:

* $n=m / M$ and $\Delta H=n \Delta H x$
* $n=m / M$
* $n=(500.0 \mathrm{~g})$ $120.91 \mathrm{~g} / \mathrm{mol}$
* $n=4.135 \mathrm{~mol}$

$$
\begin{array}{ll}
* n=m / \mathrm{M} & * \Delta H=n \Delta H_{x} \\
* n=\frac{(500.0 \mathrm{~g})}{120.91 \mathrm{~g} / \mathrm{mol}} & * \Delta H=(4.135 \mathrm{~mol})(34.99 \mathrm{~kJ} / \mathrm{mol}) \\
* n=4.135 \mathrm{~mol} & * \Delta H=144.7 \mathrm{~kJ}
\end{array}
$$

## Therefore 144.7 kJ of energy is transferred.

## Homework

* pg 299 \#13, 15, 18


# Calculations Using a Calorimeter 

## Calorimeter

* Calorimeter: device used to measure the heat released or absorbed during a chemical or physical process.



## Calorimetry

* The analysis is based on the law of conservation of energy: the total energy change of the chemical system is equal to the total energy change of the surroundings


## $\Delta$ Iisystem $=-q_{\text {surroundings }}$

## Example

* In a calorimetry experiment, 5.56 g of sodium chloride is dissolved in 75.0 mL of water at an initial temperature of $24.9^{\circ} \mathrm{C}$. The final temperature of the solution is $23.7^{\circ} \mathrm{C}$. What is the molar enthalpy of solution of sodium chloride?


## * Given:

* $m=75.0 \mathrm{~mL} \mathrm{x} 1 \mathrm{~g} / \mathrm{mL}=75.0 \mathrm{~g}$
* $c=4.18$
* $\Delta T=23.7-24.9=-1.2$
* Required:
* $\Delta H_{\text {sol }}$
* Equation:
* $n=m / M, Q=m c \Delta T, \Delta H=n \Delta H x$,


## First Find Q

$Q=m c \Delta T$
$Q=(75.0)(4.18)(-1.2)$
$Q=-376.2 \mathrm{~J}$

## First Find Q

## Then Find $n$

$Q=m c \Delta T$
$Q=(75.0)(4.18)(-1.2)$
$Q=-376.2 \mathrm{~J}$

## First Find Q

## Then Find $n$

$$
\begin{array}{ll}
Q=m c \Delta T & n=m / M \\
Q=(75.0)(4.18)(-1.2) & n=(5.56 \mathrm{~g}) /(58.44 \mathrm{~g} / \mathrm{mol}) \\
Q=-376.2 \mathrm{~J} & n=0.0951 \mathrm{~mol}
\end{array}
$$

*heat lost by water = heat absorbed by chemical system therefore $\Delta H=-q$

## Now Find $\Delta_{1 \text { isol }}$


$\Delta H_{s o l}=-q / n$
$\Delta H$ sol $=-(-376.2) /(0.0951)$
$\Delta H s o l=3.96 \times 103 \mathrm{~J} / \mathrm{mol}$ or $3.96 \mathrm{~kJ} / \mathrm{mol}$

# Therefore the molar enthalpy of solution is $3.96 \mathrm{~kJ} / \mathrm{mol}$ 

## Example

* A 50.00 mL volume of $0.300 \mathrm{~mol} / \mathrm{L}$ CuSO ${ }_{4}$ laq) is mixed with 50.00 mL of $\mathrm{NaOH}($ aq) . The initial temperature of both solutions is $21.40^{\circ} \mathrm{C}$. After mixing the two solutions in a calorimeter the peak temperature is $24.60^{\circ} \mathrm{C}$. Determine the enthalpy of change for the reaction.


## First Find Q

We know $c$ and $\triangle T$, we need $m$

$$
\begin{gathered}
V_{\text {Total }}=V_{\text {cusol }}+V_{\mathrm{NaOH}} \\
V_{\text {Total }}=50 \mathrm{~mL}+50 \mathrm{~mL} \\
m=V_{d} \\
m=(100 \mathrm{~mL})(1.00 \mathrm{~g} / \mathrm{mL}) \\
m=100 \mathrm{~g}
\end{gathered}
$$

$Q=m c \Delta T$
$Q=(100)(4.19)(24.60-21.40)$
$Q=1.341 \times 10^{3} \mathrm{~J}$ OR 1.34 kJ

## First Find Q

## Then Find $n$

We know $c$ and $\triangle I$, we need $m$

$$
\begin{aligned}
& V_{\text {Total }}=V_{\text {cusol }}+V_{\mathrm{NaOH}} \\
& V_{\text {Total }}=50 \mathrm{mLL}+50 \mathrm{~mL}
\end{aligned}
$$

$n=c V$
$n=(0.3 \mathrm{~mol} / \mathrm{L})(0.05 \mathrm{~L})$ $n=0.0150 \mathrm{~mol}$

$$
\begin{gathered}
m=(100 \mathrm{Vd} \\
m=100 \mathrm{~g} /(1.00 \mathrm{~g} / \mathrm{mL}) \\
m=1
\end{gathered}
$$

$Q=m c \Delta T$
$Q=(100)(4.19)(24.60-21.40)$
$Q=1.341 \times 10^{3} \mathrm{~J}$ OR 1.34 kJ

## First Find Q

## Then Find $n$

We know $c$ and $\triangle I$, we need $m$

$$
V_{\text {Total }}=V_{\text {cusO4 }}+V_{\mathrm{NaOH}}
$$

$$
n=c V
$$

$$
V_{\text {Total }}=50 \mathrm{~mL}+50 \mathrm{~mL}
$$

$$
n=(0.3 \mathrm{~mol} / \mathrm{L})(0.05 \mathrm{~L})
$$

$$
n=0.0150 \mathrm{~mol}
$$

$m=\mathrm{Vd}$ $m=(100 \mathrm{~mL})(1.00 \mathrm{~g} / \mathrm{mL})$ $m=100 \mathrm{~g}$

$$
\Delta H_{\text {sysstem }}=-q
$$

$Q=m c \Delta T$
$Q=(100)(4.19)(24.60-21.40) \quad \Delta H_{\text {system }}=-1.341 \mathrm{~kJ}$ $Q=1.341 \times 10^{3} \mathrm{~J}$ OR 1.341 kJ

Now Find $\Delta$ treaction
$\Delta H_{\text {reaction }}=(-1.341 \mathrm{~kJ}) /(0.0150 \mathrm{~mol})$ $=-89.4 \mathrm{~kJ} / \mathrm{mol}$

## Homework

* Page 305 \# 23, 24,25

