

Thermodynamics!

at constant P

$$\Delta E = q_p - P\Delta V$$

$$\Delta E = \Delta H - P\Delta V$$

$$\Delta E = \Delta H - \Delta nRT$$

internal energy
 $\Delta E = q + w$
 work!

LAW
 Energy cannot be created or destroyed only changed in its form

$$H = E + PV \quad \text{definition of enthalpy}$$

CALORIMETRY

$$\begin{cases} \Delta E = q_v (\text{bomb calorimetry}) \\ \Delta H = q_p (\text{coffee-cup calorimetry}) \end{cases}$$

$$\begin{aligned} q &= n C_m \Delta T && \text{moles} \\ q &= m C_s \Delta T && \begin{aligned} \text{molar heat capacity in J/mol}\cdot\text{K} \\ \text{mass in grams} \end{aligned} \\ &&& \begin{aligned} \text{specific heat capacity in J/g}\cdot\text{K} \end{aligned} \end{aligned}$$

ISOTHERMAL expansion

$$\Delta T = 0, T \text{ is constant}$$



$$\Delta E = 0 \text{ and } \Delta H = 0$$

$$q = -w \quad \text{and! } P_1/V_1 = P_2/V_2 \quad (\text{Boyle's Law})$$

$$\int dw = \int -Pdv$$

$$\begin{aligned} * w &= -nRT \ln \frac{V_2}{V_1} \\ * q &= nRT \ln \frac{V_2}{V_1} \\ * \Delta S &= nR \ln \frac{V_2}{V_1} \end{aligned}$$

system heat is opposite in sign from the calorimeter!

If heating gases you need to know all the different Cv's & Cp's

Cv	Cp
monatomic $\frac{3}{2}R$	$\frac{5}{2}R$
linear $\frac{5}{2}R$	$\frac{7}{2}R$
non-linear $\frac{6}{2}R$	$\frac{9}{2}R$

$$S = k \ln W$$

↑ microstates

2nd LAW
 all spontaneous changes are accompanied by an increase in universal entropy

3rd LAW

The entropy of a perfectly crystalline substance at absolute zero is zero.

When $\Delta S_{univ} = 0$

for a process, you have reached EQUILIBRIUM

We PREFER

$$\Delta G = 0$$

HESS' LAW

$$\Delta H_{rxn} = \Delta H_1 + \Delta H_2 + \Delta H_3 + \dots$$

$$\Delta H_{rxn}^\circ = \sum \Delta H_f^\circ(\text{prod}) - \sum \Delta H_f^\circ(\text{react})$$

$$\Delta H_{rxn} = \sum \text{B.E.}(\text{react}) - \sum \text{B.E.}(\text{prod})$$

$$\Delta G_{rxn}^\circ = \sum \Delta G_f^\circ(\text{prod}) - \sum \Delta G_f^\circ(\text{react})$$

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FREE ENERGY

$$G = H - TS$$

at constant P + T

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

$$\Delta G = -T\Delta S_{\text{universe}}$$

governs 2nd Law

conditional!

$$\Delta G = 0 = \Delta H - T\Delta S$$

$$\begin{aligned} \Delta H &= T\Delta S \\ T &= \Delta H/\Delta S \end{aligned}$$

Phase Changes

$$\Delta S_{\text{trans}} = \frac{\Delta H_{\text{trans}}}{T_{\text{trans}}}$$

$$\Delta S_{rxn}^\circ = \sum S^\circ(\text{prod}) - \sum S^\circ(\text{react})$$