

ENERGY & MATTER SYNOPTIC

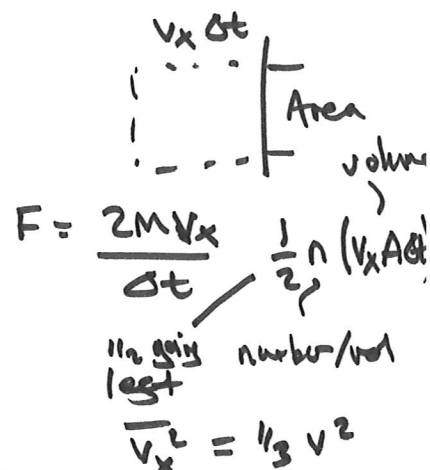
IDEAL GAS : molecules, N - uniform & random motion
 small relative to separation
 Newton's laws
 only elastic collisions with walls & each other

$$\rightarrow V, \rho = \frac{Nm}{V}, P = \frac{1}{3} n m \overline{v^2}$$

$$U = KE = \frac{1}{2} m \overline{v^2} N$$

$$\rightarrow PV = NRT_I$$

$$\frac{1}{2} m \overline{v^2} = \frac{3}{2} kT_I \quad \& \quad U = N \frac{3}{2} kT_I$$



EQUIPARTITION : $\frac{1}{2} kT$ energy / degree of freedom

Fails if $kT <$ quantum excitation

HEAT CAPACITY: $C_V = \left(\frac{\partial Q}{\partial T} \right)_V$ $C_P = \left(\frac{\partial Q}{\partial T} \right)_P$

monatomic: $U = \frac{3}{2} kT$ or $U = \frac{3}{2} n m \underbrace{N_A R}_R T$

BOLTZMANN FACTOR

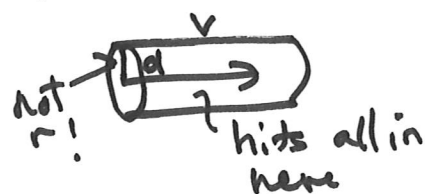
$P(E) \propto e^{-E/kT}$

MEAN FREE PATH

$\lambda = \text{speed} / \text{collision rate}$

$$\lambda \sim \frac{1}{\pi d^2 n}$$

$$\hookrightarrow \pi d^2 v n$$



RANDOM WALKS

In N steps travel $N\lambda$ but distance from start $\sqrt{N}\lambda$



DIFFUSION: $\frac{dn}{dt} = -DA \frac{dn}{dx}$ $\frac{1}{x} \rightarrow \propto$ density gradient

$D \sim \lambda \bar{v}$ "m²s⁻¹"

THERMAL CONDUCTION: $\frac{dQ}{dt} = -KA \frac{dT}{dx}$ \propto temperature gradient

$K \sim \lambda \bar{v} C$

THERMODYNAMICS

0th Law: A in thermal \equiv i/m with B & C \Rightarrow B & C in thermal \equiv i/m

1st Law: Energy conservation
"work generates heat \rightarrow energy"

$\Delta U = Q + W$

internal energy ? heat in ? work done on gas

Work by gas = $\int P dV$

\Rightarrow $PA \cdot dx$

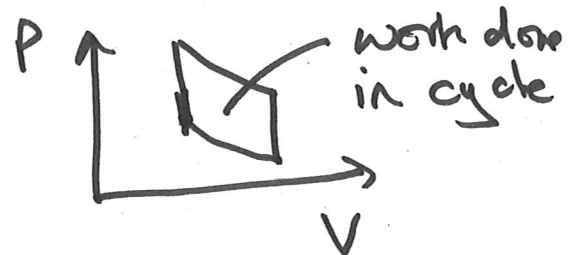
At constant P $PdV \xrightarrow[\text{mole}]{\text{mole}} RdT$

$C_p = C_v + R$

Isothermal: $T = \text{const}$ $W = \int P dV = \int_{V_1}^{V_2} \frac{n_m RT}{V} dV$

Adiabatic: no energy in $\rightarrow \Delta U = W \Rightarrow PV^\gamma = \text{constant}$

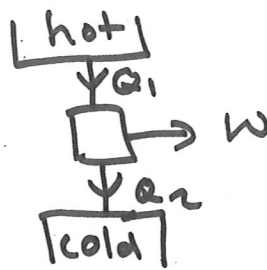
PV Indicator Diagrams



2nd Law: You can only extract work from heat flow from hot \rightarrow cold body

Carnot Engines

Reverse arrow
 \Rightarrow heat pump
 $\eta = \frac{Q_1}{W} > 1$



$$\eta = \frac{W}{Q_1}$$

at best

$$= \frac{Q_1 - Q_2}{Q_1}$$

Ideal Gas Realization



expand isothermally
 heat in



expand
 adiabatically



isothermal
 compress
 heat out



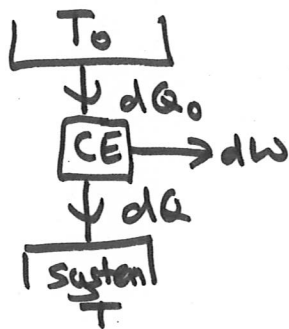
adiabatically
 expand



complete cycle
 reversible

Defining T : Use Q_1, Q_2 as Carnot to define
 $\propto T_1, T_2$ $T_{ideal} = T$

Entropy



$$\frac{dQ_0}{dQ} = \frac{T_0}{T}$$

Do process where CE
 does W & system
 somehow returned to
 initial state

$$Q_0 = \int dQ_0 = T_0 \int \frac{dQ}{T} \leq 0$$

you can't extract
 work just from
 hot thing

$dS = \frac{dQ}{T}$ is a function of the state of
 T_0 reservoir

Statistical interpretation is movement to most
 likely state $S = k \ln W$

2nd Law Refresh : Entropy always rises.

Enthalpy: energy changes involving gas expansion
 minimize $H = U + PV$

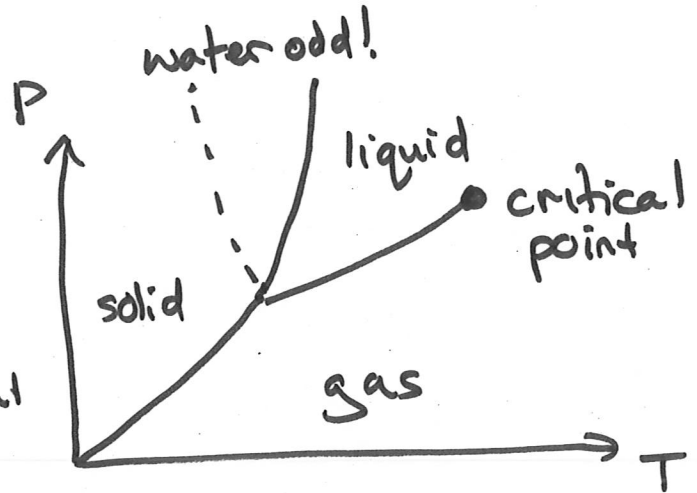
Gibb's Free Energy: is also allow to thermalize with environment then S as system + Universe must grow
 \Rightarrow minimize $G = U + PV - TS$

Phase Diagrams

Slopes given by Clausius - Clapyron eqn

$$\frac{dP}{dT} = \frac{L \in \text{Latent heat}}{T(v_2 - v_1)}$$

(volumes/mole)



Van der Waals Gases

$$\left(P + N^2 \frac{a}{V^2}\right) (V - Nb) = NkT$$

includes inter-atomic forces. size of force $\propto N/V$ & number of collisions $\propto N/V$

removes finite volume of atoms from V

