

Recap Extensive	Intensive	
	Potential	Specific
S	$T = \partial U / \partial S$	$s = S/\alpha$
Ψ	$\Theta = \partial U / \partial \Psi$	$\psi = \Psi/\alpha$
$k+2$ state + extent	$k+1$ $\sum \Psi d\Theta = 0$	$k+1$ $\alpha(\Psi)$

variables of state

$$U(S, V, N) \rightarrow U(\Psi_1, \dots, \Psi_{k+2})$$



$$\alpha = n \rightarrow u(s, v)$$

$$\alpha = v \rightarrow u(s, n)$$

$$\alpha = s \rightarrow u(v, n)$$

$$\rightarrow u(\Psi_1, \dots, \Psi_{k+2})$$

$$\leftarrow \alpha = n$$

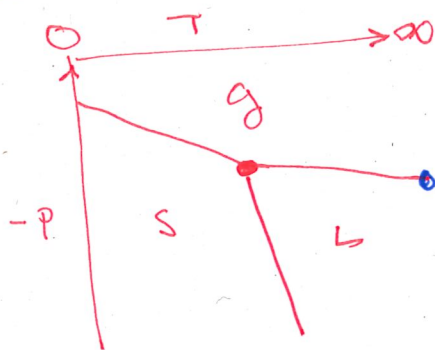
$$u = Ts + (-P)v + \mu n$$

$$du = Tds + (-P)dv$$

for now

Why specific variables $\Rightarrow$ Heterogeneous systems $\Rightarrow$ matter in different states $\Rightarrow$ potentials don't distinguish these states.
of course J/kg is also more convenient than J for 6.3 kg.

State of a heterogeneous system - is determined if the state and relative amounts of every phase of the system is known

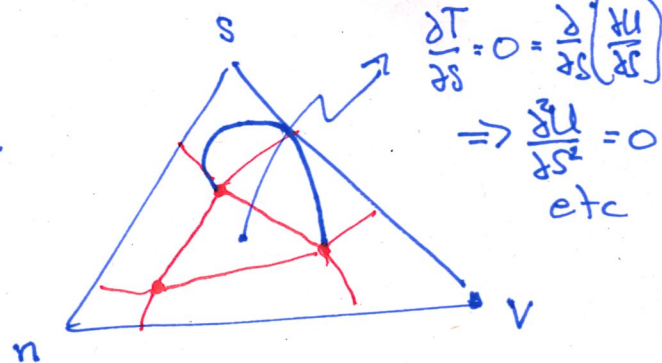


$$\alpha = S + V + N$$

$$\rightarrow s + v + n = 1$$

$$\downarrow u(s, v)$$

$$T = \frac{\partial u}{\partial s}$$



what if you don't have the phase diagram $\rightarrow$ the phase rule

Everything has to be true no matter how its written

2021 S.2

Heterogeneous system
of p -phases

Equality of potentials =
condition of internal eq.

rank $\underline{k+2}$



$$1 \leq p \leq k+2$$

if $p = k+1 \Rightarrow$

1 potential can be arbitrarily specified

if $p = k \Rightarrow$

2 ...

$$\Rightarrow p - k + 2$$

= the number of potentials that
can be changed independently

$F = p - k + 2 \Rightarrow$ is **NOT** the number of intensive
degrees of freedom, but it is
the number of potentials that
can be changed without causing
a change of phase.

Phase proportions (AKA "the lever rule")

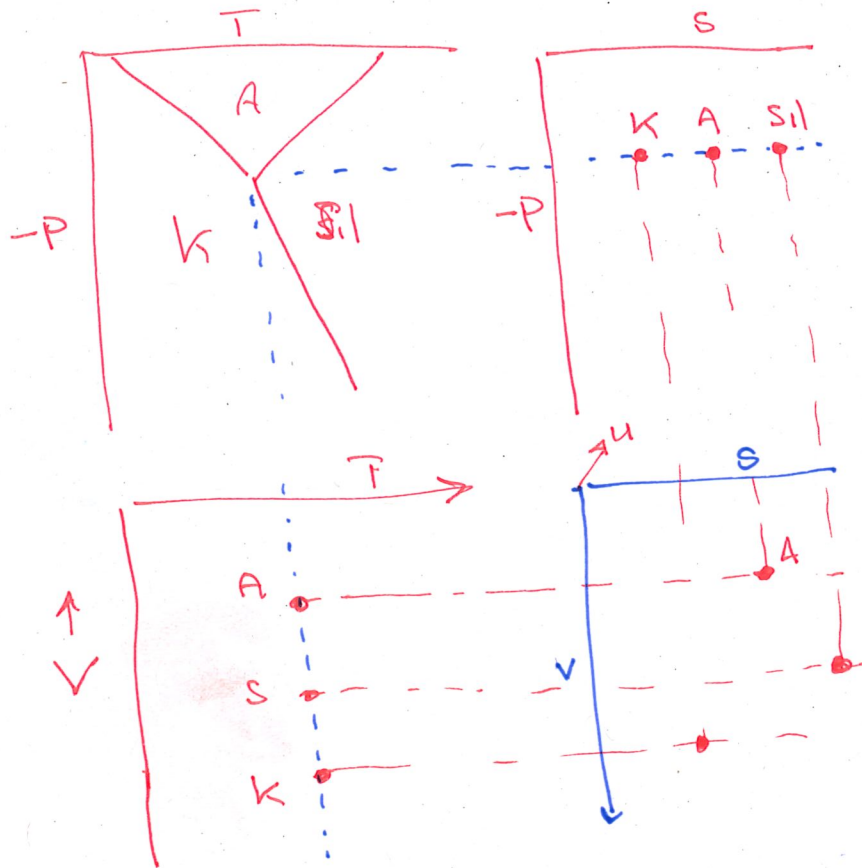
For any extensive property:

$$N^{\text{sys}} = \sum_j^p N^j = \sum_j^p \underbrace{x^j}_{\text{specific (intensive)}} \underbrace{n^j}_{\text{absolute amount}}$$

$$\frac{N^{\text{sys}}}{Q^{\text{sys}}} = \underbrace{\eta^{\text{sys}}}_{\text{intensive}} = \sum_j^p \underbrace{\frac{x^j}{Q^{\text{sys}}}}_{\text{relative amount}} n^j = \sum_j^p x^j \eta^j$$

each specific variable provides an eq in p -unknown
proportions $\rightarrow$ to determine the state of a system
containing p -phases the system must be described
by p specific variables.

Problem 2.1 $\frac{\partial \phi}{\partial \psi} > 0$ for reversible processes $\rightarrow \frac{\partial \psi}{\partial \phi} > 0$ 2021 S.3



$P, T \rightarrow P_{max} = 1$
 $V, T \rightarrow P_{max} = 2$
 $P, S \rightarrow P_{max} = 2$
 $S, V \rightarrow P_{max} = 3$

$$u^i = T s^i + (-P) v^i + \mu n^i$$

$$\begin{bmatrix} s^A & v^A & n^A \\ s^K & v^K & n^K \\ s^{Sil} & v^{Sil} & n^{Sil} \end{bmatrix} \begin{bmatrix} T \\ -P \\ \mu \end{bmatrix} = \begin{bmatrix} u^A \\ u^K \\ u^{Sil} \end{bmatrix}$$

$$\bar{A} \bar{x} = \bar{b}$$

$$\begin{bmatrix} s^A & s^K & s^{Sil} \\ v^A & v^K & v^{Sil} \\ n^A & n^K & n^{Sil} \end{bmatrix} \begin{bmatrix} x^A \\ x^K \\ x^{Sil} \end{bmatrix} = \begin{bmatrix} s^{sys} \\ v^{sys} \\ n^{sys} \end{bmatrix}$$

if $\alpha = N$ then the x^i 's are molar fractions

if $\alpha = V$?

1 $\leftarrow$ if $\alpha = N$