

Maxwell thermodynamic relation

Gibb's function $\rightarrow$ Gibb's function (G) is the energy liberated or absorbed in reversible process at constant pressure and constant temperature.

From the definition of enthalpy

$$H = U + pV$$

Change in enthalpy

$$dH = dU + d(pV)$$

$$dH = dU + p dV + V dp$$

$$dH = T ds + V dp$$

If the process is isothermal

$$T ds = d(Ts)$$

$$dH = d(Ts) + V dp$$

$$dH - d(Ts) = V dp$$

$$d(H - Ts) = V dp$$

Here $(H - Ts) = G$ is known as Gibb's function.

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sem IV Unit IV

Maxwell's thermodynamic relation

G remains constant if a thermodynamic process remains isothermal as well as isobaric

$$dG = 0 \text{ when } dT = 0 \text{ and } dP = 0$$

$$G = H - TS$$

$$dG = dH - d(TS)$$

$$= dH - Tds - SdT$$

$$dG = Tds + VdP - Tds - SdT$$

$$dG = VdP - SdT$$

Partial differentiation with respect to variables

$$P \& T \quad \left(\frac{\partial G}{\partial T} \right)_P = -S, \quad \left(\frac{\partial G}{\partial P} \right)_T = V$$

As dG is a perfect differential

$$\frac{\partial}{\partial P} \left(\frac{\partial G}{\partial T} \right)_P = \frac{\partial}{\partial T} \left(\frac{\partial G}{\partial P} \right)_T$$

$$\left(\frac{\partial V}{\partial T} \right)_P = - \left(\frac{\partial S}{\partial P} \right)_T$$

This equation is known as Maxwell's thermodynamic relation.