For a particular function X (where X is either U, H, F, or G), the following relationship holds:

What is the thermodynamic potential, X?

- A. Internal Energy, U
- B. Enthalpy, H
- C. Helmholtz Free Energy, F'
- D. Gibbs Free Energy, G



 $\left(\frac{\partial X}{\partial V}\right)_{T.N} = -P$



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Derive the Maxwell relation corresponding to the 2nd mixed partials of G with respect to T and P. Make sure you include which variables are held constant.

By looking at the appropriate Maxwell relation, which of the following is equal to $\left(\frac{\partial S}{\partial P}\right)_{T,N}$?







most relevant for this question?

- A. Internal Energy, U
- B. Enthalpy, H
- C. Helmholtz Free Energy, F
- D. Gibbs Free Energy, G



Suppose you are interested in the heat transfer at constant pressure of a particular chemical reaction. Which thermodynamic potential is



dA = dF (where A is the availability and F is the Helmholtz free energy) in the case of constant volume and temperature. Does this require that the constant temperature of the system T be equal to the constant temperature of the reservoir, T_0 ?

A. YesB. NoC. ???

Is carbonic acid a strong acid? That is, does H₂CO₃ spontaneously disassociate into HCO₃- and H+?

Substance (form)	$\Delta_f H$ (kJ)	$\Delta_f G$ (kJ)	S (J/K)	$C_P (\mathrm{J/K})$	$V (\mathrm{cm}^3)$
H_2CO_3 (aq)	-699.65	-623.08	187.4		
HCO_3^- (aq)	-691.99	-586.77	91.2		
H^+ (aq)	0	0	0	0	

A. Yes B. No C. ???



To induce more disassociation from H₂CO₃ into HCO₃- and H⁺, should we increase temperature or decrease temperature?

Substance (form)	$\Delta_f H$ (kJ)	$\Delta_f G$ (kJ)	S (J/K)	$C_P (\mathrm{J/K})$	$V (\mathrm{cm}^3)$
H_2CO_3 (aq)	-699.65	-623.08	187.4		
HCO_3^- (aq)	-691.99	-586.77	91.2		
H^+ (aq)	0	0	0	0	

A. Increase temperatureB. Decrease temperatureC. ???

What does the slope represent in this graph?

- A. Volume
- **B.** Pressure
- C. Entropy
- D. None of the above

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Suppose you are looking at a phase diagram (P vs. T). At the boundary between two phases,

- A. The entropy of the two phases is the same
- B. The Gibbs energy of the two phases is the same
- C. The Helmholtz energy of the two phases is the same
- D. The volume of the two phases is the same



Water boils at 100.0 degrees C at atmospheric pressure. What absolute pressure is required for it to boil at 101.0 degrees C?

- A. A pressure slightly higher than 1.0atm.
- B. A few atm (say, between 2.0atm and 9.0atm).
- C. Between 10-100 atm.
- D. Well over 100 atm.

2	Substance	∆H [°] f (25°C) kJ mol ^{−1}	<i>S</i> ° (25°C) J K ⁻¹ mol ⁻¹	ΔG_{f}° (25°C kJ mol $^{-1}$
	H₂O(ℓ)	-285.83	69.91	-237.18
	H ₂ O(<i>g</i>)	-241.82	188.72	-228.59

E. None of the above -it actually requires a lower pressure.



- A. The 2nd Law of Thermodynamics
- B. The Clausius-Clapeyron relation
- C. The fact that the Gibbs Free Energy is a state function
- D. None of the above the choice is pretty arbitrary



In the Maxwell construction, which of the following best explains why the critical pressure line is drawn so that the "areas are equal"?