

19.3 Gibbs Free Energy

Gibbs Free Energy and Temperature

Free Energy and Temperature

Temperature 0 K

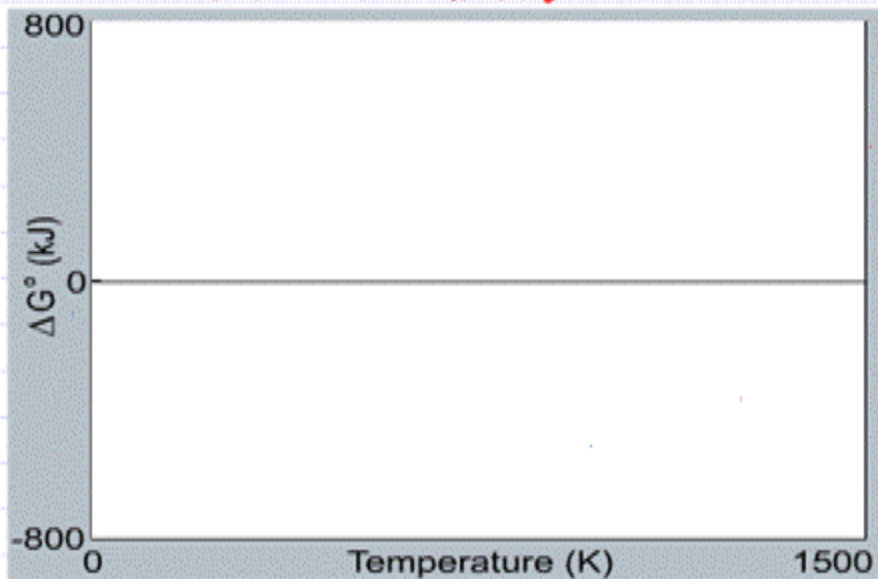
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Calculate

- $2 \text{Fe}_2\text{O}_3(\text{s}) + 3 \text{C}(\text{s}) \longrightarrow 3 \text{CO}_2(\text{g}) + 4 \text{Fe}(\text{s})$
- $\text{C}(\text{s}) + 2\text{H}_2(\text{g}) \longrightarrow \text{CH}_4(\text{g})$
- $\text{N}_2(\text{g}) + 2\text{O}_2(\text{g}) \longrightarrow 2 \text{NO}_2(\text{g})$
- $\text{CaCO}_3(\text{s}) \longrightarrow \text{CaO}(\text{s}) + \text{CO}_2(\text{g})$
- $2 \text{HCl}(\text{aq}) + \text{Na}_2\text{CO}_3(\text{s}) \longrightarrow 2 \text{NaCl}(\text{aq}) + \text{H}_2\text{O}(\ell) + \text{CO}_2(\text{g})$

$\Delta S^\circ = 561.00 \text{ J/K}\cdot\text{mol}$
 $\Delta H^\circ = 468.00 \text{ kJ/mol}$
 $\Delta G^\circ = 468.00 \text{ kJ/mol}$

See Cross Web Site:



i) $\Delta G^\circ_{\text{RXN}} = \sum \Delta G^\circ_f(\text{Products}) - \sum \Delta G^\circ_f(\text{Reactants})$

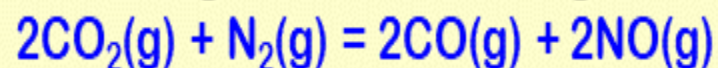
ii) $\Delta G^\circ_{\text{RXN}} = \Delta H^\circ_{\text{RXN}} - T\Delta S^\circ_{\text{RXN}}$ OR $\Delta G^\circ_{\text{RXN}} = \Delta H^\circ_{\text{RXN}} - T\Delta S^\circ_{\text{RXN}}$: This one often preferred as we can glean further insights into the reaction. ? What type of insights?

- $\Delta H^\circ > 0, \Delta S^\circ > 0$: Likely ΔG° starts out + due to ΔH° being in kJ but at what T does $-T\Delta S$ which is (-) make $\Delta G^\circ < 0$?
- $\Delta H^\circ < 0, \Delta S^\circ < 0$: Likely ΔG° starts out - due to ΔH° being in kJ but at what T does $-T\Delta S$ which is (+) make $\Delta G^\circ > 0$?
- $\Delta H^\circ > 0, \Delta S^\circ < 0$: $\Delta G^\circ > 0$ at all T, since ΔH° is + and $-T\Delta S$ is +. Nonspontaneous at all temperatures.
- $\Delta H^\circ < 0, \Delta S^\circ > 0$: $\Delta G^\circ < 0$ at all T, since ΔH° is - and $-T\Delta S$ is -. Spontaneous at all temperatures.

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Without doing any calculations, match the following thermodynamic properties with their appropriate numerical sign for the following **endothermic reaction**.



a) ΔH_{rxn}

b) ΔS_{rxn}

c) ΔG_{rxn}

d) ΔS_{univ}



a. >0

b. <0

c. $=0$

d. >0 low T, <0 high T

e. <0 low T, >0 high T

$$\Delta G_{\text{rxn}} = \Delta H_{\text{rxn}} + (-T\Delta S_{\text{rxn}})$$

$\quad \quad \quad + \quad \quad \quad -$

a) a) ✓ : We are told that the reaction is endothermic.

b) a) ✓ : $\Delta S_{\text{rxn}} = S(\text{products}) - S(\text{reactants})$
4 gas molecules - 3 gas molecules

$$\therefore \Delta S_{\text{rxn}} > 0$$

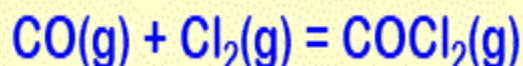
c) d) ✓ : With ΔH_{rxn} being >0 expect ΔG_{rxn} to be >0 ; however with $(-T\Delta S) < 0$ expect that at some high value of T that this will overcome ΔH and make $\Delta G < 0$.

d) e) ✓ : Tutor note :). It is easier to understand and predict ΔG_{rxn} . So focus on that and ΔS_{univ} is the reverse of ΔG_{rxn} . :)

19.3 Gibbs Free Energy

Gibbs Free Energy and Temperature

Without doing any calculations, match the following thermodynamic properties with their appropriate numerical sign for the following **exothermic reaction**.



a) ΔH_{rxn}

b) ΔS_{rxn}

c) ΔG_{rxn}

d) ΔS_{univ}



a. >0

b. <0

c. $=0$

d. >0 low T, <0 high T

e. <0 low T, >0 high T

$$\Delta G_{\text{rxn}} = \Delta H_{\text{rxn}} + (-T\Delta S_{\text{rxn}})$$

- +

a) b) ✓ : Reaction is exothermic.

b) b) ✓ : $\Delta S_{\text{rxn}} = S(\text{Products}) - S(\text{Reactants})$
1 gas molecule - 2 gas molecules

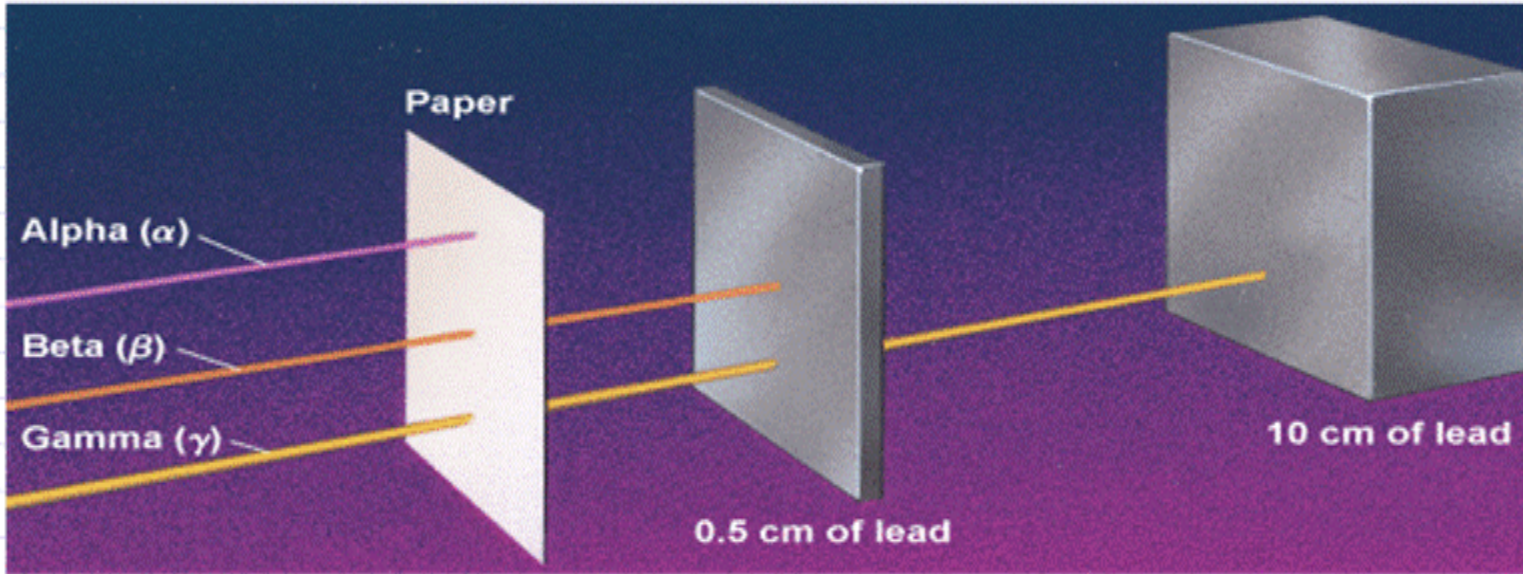
$$\therefore \Delta S_{\text{rxn}} < 0$$

c) e) ✓ : Since $\Delta H_{\text{rxn}} < 0$ expect ΔG_{rxn} to start < 0 . However since $(-T\Delta S_{\text{rxn}}) > 0$ expect that at some high value of T that this will surpass ΔH_{rxn} , thereby making $\Delta G_{\text{rxn}} > 0$.

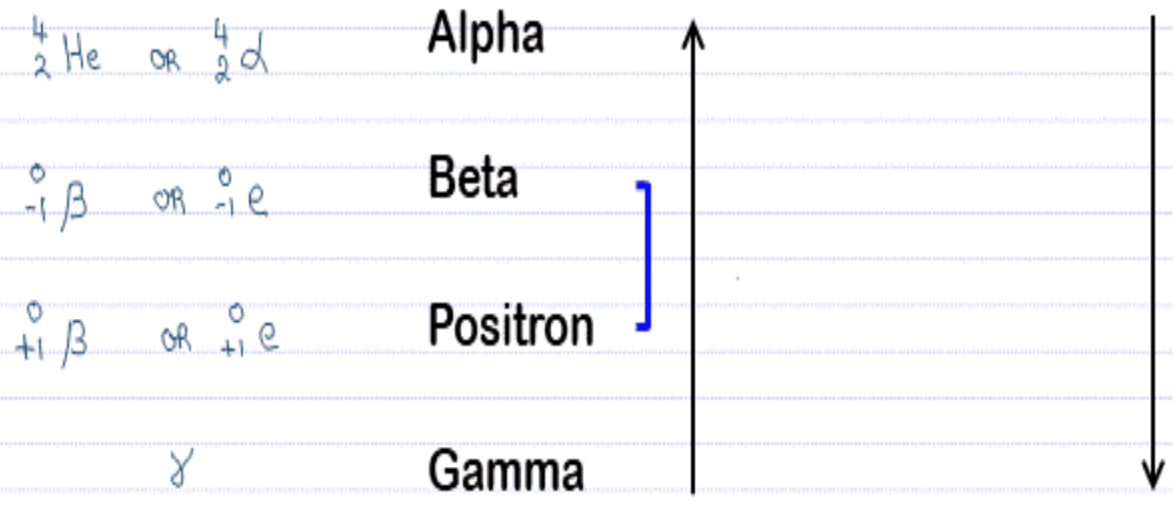
d) d) ✓ : See previous slide for simple trick to predict ΔS_{univ} .

24.1 Nuclear Reactions

The Penetrating Power of Radiation

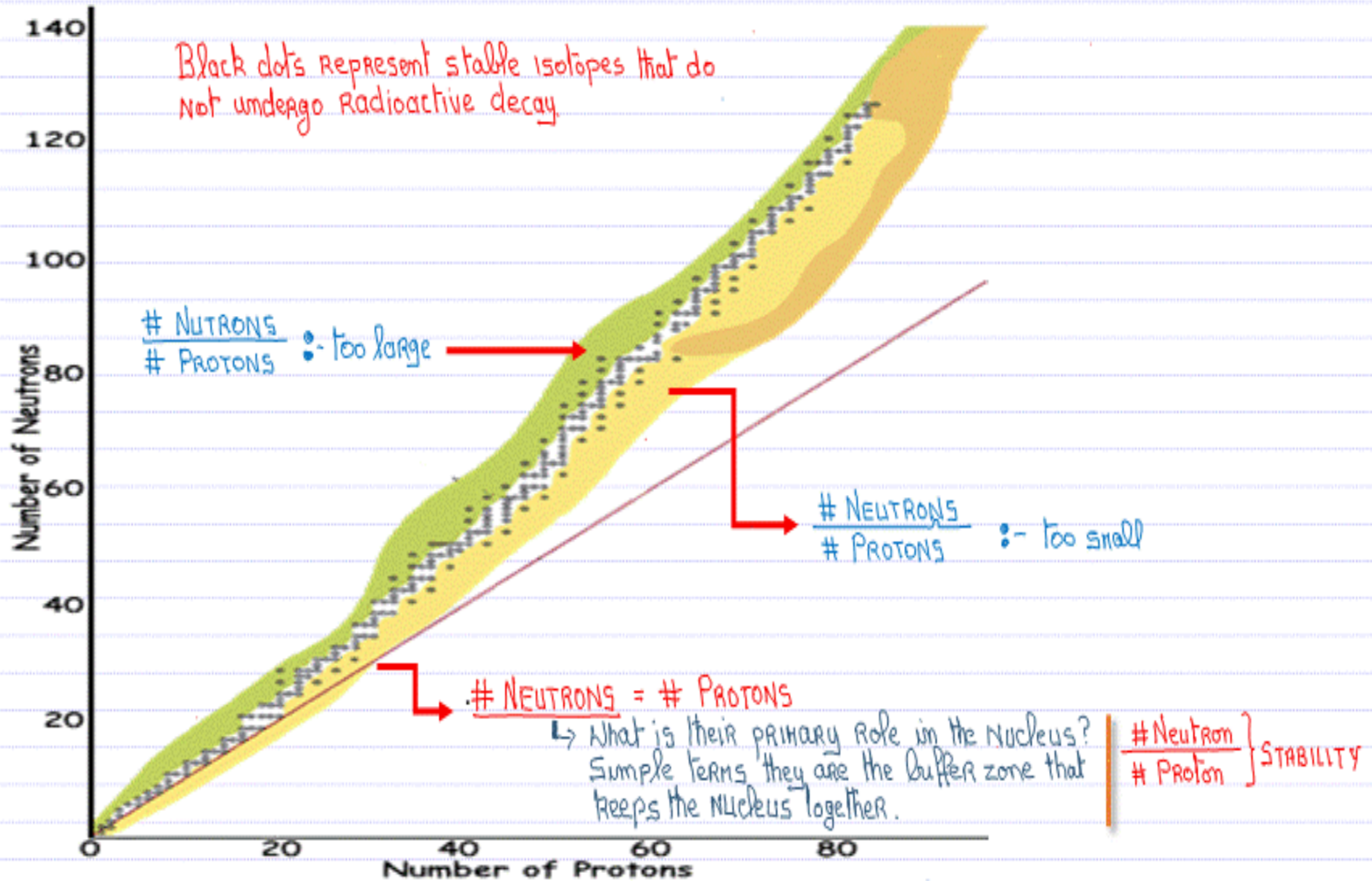


Particles: **Energy** **Vs** **Penetration Power**



24.2 Nuclear Stability

Band of Stability



24.2 Nuclear Stability

Natural Radioactive Decay

1. Alpha Emission: Nucleus kicks out an Alpha particle, ${}^4_2\text{He}$ or ${}^4_2\alpha$ ONL: ${}^4_2\text{He}$

2. Beta Emission: Nucleus kicks out a Beta particle, ${}^0_{-1}e$ or ${}^0_{-1}\beta$ ONL: $-1e^0$

3. Positron Emission: Nucleus kicks out a Positron, ${}^0_{+1}e$ or ${}^0_{+1}\beta$ ONL: $+1e^0$

4. Electron Capture: The nucleus captures one of its own electrons.

All of the above processes also result in emitting γ Rays.

Very Important Note: The most common mistake in balancing nuclear reaction is with 4.
In 1, 2 and 3 the particle is on the Product side.
In 4, the captured electron is on the Reactant side.

The driving force for all of these is to stabilize the Neutron to Proton ratio. Lets have a look at what each one does.