

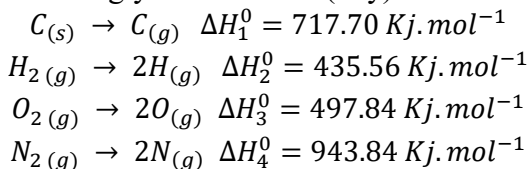
Series N ° : 4

Exercise N°1:

Calculate the bond energy of the **C=O** bond in the compound glycine (Gly) $\text{H}_2\text{N}-\text{CH}_2-\text{COOH}$, knowing that the standard molar enthalpy change for the formation of this compound from its elements in their standard state is $\Delta H^\circ_f(\text{Gly}) = -536.71 \text{ KJ/mol}$.

Given:

The standard enthalpy of sublimation of glycine $\Delta H^\circ_{\text{Sub}}(\text{Gly}) = 175.90 \text{ KJ/mol}$.



The bond	N-H	C-N	O-H	C-O	C-H	C-C
The bond energy $E_{(A-B)} (\text{ KJ/mol})$	390.41	304.30	462.31	357.39	412.57	345.27

Exercise N°2:

Consider 1 kg of air as an ideal gas undergoing a Carnot cycle. AB and CD are isothermal processes, and BC and DA are adiabatic processes, with temperatures at point A equal to $T_1 = 300\text{K}$ and pressures at points C, B, A respectively as $P_3 = 9 \text{ atm}$, $P_2 = 3 \text{ atm}$, $P_1 = 1 \text{ atm}$:

Calculate the efficiency of the cycle using two different methods:

- Using the heat balance (bilan).
- Using the temperatures at the two ends of the cycle.

Calculate the ΔS of air during the four transformations. Given $C_p = 103 \text{ J/Kg.K}$; $\gamma = 7/5$.

Exercise N°3:

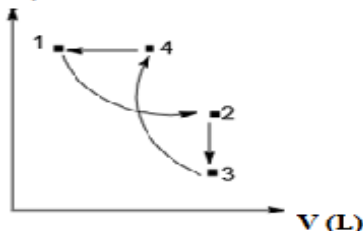
0.056 m^3 of an ideal gas undergoes heating at constant volume from the initial state ($T_1 = 38^\circ\text{C}$, $P_1 = 1.03 \text{ atm}$) to the second state ($P_2 = 1.72 \text{ atm}$), then undergoes heating at constant pressure to a volume of 0.126 m^3 .

For each transformation, calculate the final temperature, the amount of heat, work done, and the change in both internal energy and entropy. $C_v = 20.82 \text{ J/mol.K}$.

Exercise N°4:

Let the Clapeyron diagram represent a series of reversible transformations for one mole of an ideal gas with the given data in the table:

P (atm)



	P (atm)	V (L)	T (K)
1	10	1	600
2	2	5	600
3	1	5	300
4	10	1.25	750

Calculate ΔS for each transformation, then ΔS for the cycle.

Given: $C_p = 5.03 \text{ cal/K.mol}$; $R = 2 \text{ cal/mol.K}$; $C_v = 3.03 \text{ cal/K.mol}$.

Exercise N°5:

Ammonia gas at temperature 24 °C and pressure 10 atm is liquefied. We condense 1 mole of ammonia gas ($P_1 = 1$ atm, $T_1 = -20^\circ\text{C}$) to one mole of liquefied ammonia ($P_2 = 10$ atm, $T_2 = 24^\circ\text{C}$). Assume ideal behavior for ammonia gas and neglect the volume of the liquid state compared to the gas state.

Calculate ΔH , ΔU , and ΔS for the aforementioned transformation.

Given:

$C_p(\text{NH}_3)_g = 24.66 \text{ J/mol.K}$; $\Delta H_{\text{vap}}(\text{NH}_3) = 19.825 \text{ KJ/mol}$ at 24°C and 10 atm

Exercise N°6:

Calculate the change in entropy (ΔS) during the transformation of 1 mole of solid iodine at 25°C to gaseous iodine at 184°C under atmospheric pressure using the following data:

$C_p(\text{I}_2 \text{ solid}) = 54.6 \text{ J/K.mol}$; $C_p(\text{I}_2 \text{ liquid}) = 81.5 \text{ J/K.mol}$; $\Delta H_{\text{fus}} = 15633 \text{ J/mol}$.

$T_{\text{fusion}}(\text{I}_2 \text{ solid}) = 113.6^\circ\text{C}$; $\Delta H_{\text{vaporization}} = 25498 \text{ J/mol}$. $T_{\text{vaporization}}(\text{I}_2 \text{ liquid}) = 184^\circ\text{C}$.

Exercise N°7:

Consider the following chemical reaction, which takes place at a temperature of 25°C :



- 1- Calculate the change in standard enthalpy and the change in internal energy for this reaction.
- 2- Calculate the change in standard entropy for this reaction.
- 3- What is the nature of this reaction?
- 4- Calculate the change in standard enthalpy for this reaction at 800 K.

Given: Under standard conditions ($P = 1$ atm, $T = 298 \text{ K}$), we have:

	$\text{H}_2\text{S}_{(\text{g})}$	$\text{O}_{2(\text{g})}$	$\text{H}_2\text{O}_{(\text{g})}$	$\text{SO}_{2 (\text{g})}$
$\Delta H_f^\circ (\text{KJ} / \text{mole})$	-20.6	0	-241.8	-296.8
$S^\circ (\text{J} / \text{mole} \cdot ^\circ\text{K})$	205.8	205.1	188.8	248.2
$C_p (\text{J} / \text{mole} \cdot ^\circ\text{K})$	34.2	29.4	33.6	39.9