



# CHEM 101 2<sup>ND</sup> MID TERM EXAM 1STSEM 1434-1435

**First: Choose the correct answer:**

**(FOR CHEM 101  $10 \times 1 = 10$  marks) (FOR CHEM 103  $20 \times 1 = 20$  marks)**

1. The sign of  $\Delta H$  for the process  $\text{CO}_2(\text{s}) \rightleftharpoons \text{CO}_2(\text{g})$  is: (**The symbol "H" means enthalpy**)
 

A) positive and  $H_{\text{CO}_2(\text{s})} > H_{\text{CO}_2(\text{g})}$

B) positive and  $H_{\text{CO}_2(\text{g})} > H_{\text{CO}_2(\text{s})}$

C) negative and  $H_{\text{CO}_2(\text{s})} > H_{\text{CO}_2(\text{g})}$

D) negative and  $H_{\text{CO}_2(\text{g})} > H_{\text{CO}_2(\text{s})}$

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2. Which of  $\Delta H_{\text{rxn}}^\circ$  of the following equations represents  $\Delta H_{\text{f}, \text{K}_3\text{PO}_4(\text{s})}^\circ$ ?
 

A)  $3\text{K}(\text{s}) + \text{PO}_2(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{K}_3\text{PO}_4(\text{s})$

B)  $\text{K}_3(\text{s}) + \text{P}(\text{s}) + \text{O}_4(\text{g}) \rightarrow \text{K}_3\text{PO}_4(\text{s})$

C)  $\text{K}_3\text{P}(\text{s}) + 2\text{O}_2(\text{g}) \rightarrow \text{K}_3\text{PO}_4(\text{s})$

D)  $3\text{K}(\text{s}) + \text{P}(\text{s}) + 2\text{O}_2(\text{g}) \rightarrow \text{K}_3\text{PO}_4(\text{s})$

---

3. A balanced chemical equation with specified value of  $\Delta H$  and states of substances is called:
 

A) a thermochemical equation.

B) a combustion reaction.

C) the first law of thermodynamics.

D) Hess's law.

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4. Change in internal energy ( $\Delta E^\circ$ ), in kJ, of the following reaction is:
 

$2\text{NaHCO}_3(\text{s}) \rightarrow \text{Na}_2\text{CO}_3(\text{s}) + \text{H}_2\text{O}(\text{g}) + \text{CO}_2(\text{g})$

$\Delta H_{\text{rxn}}^\circ = 129 \text{ kJ}$

A) 121.04

B) 134.04

C) 124.04

D) 114.04

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5. If 10.0 g of a metal ( $C_s = 0.896 \text{ J/g K}$ ) at 298 K is supplied with 313.5 J of heat, its final temperature, in K, will be:
 

A) 353

B) 333

C) 323

D) 373

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6. From table below,  $\Delta H_{\text{rxn}}^\circ$  of the following reaction, in kJ, is:
 

$\text{PCl}_3(\text{g}) + 3\text{HCl}(\text{g}) \rightarrow 3\text{Cl}_2(\text{g}) + \text{PH}_3(\text{g})$

$\Delta H_{\text{rxn}}^\circ = ?$

Compound	$\text{PH}_3(\text{g})$	$\text{PCl}_3(\text{g})$	$\text{HCl}(\text{g})$
$\Delta H_{\text{f}}^\circ / \text{kJ mol}^{-1}$	+ 5.40	- 288.07	- 92.30

A) 570.37

B) 507.37

C) 705.37

D) 750.37

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7. Knowing that:
 

$\frac{1}{2}\text{H}_2(\text{g}) + \frac{1}{2}\text{Cl}_2(\text{g}) \rightarrow \text{HCl}(\text{g})$

$\Delta H_{\text{rxn}}^\circ = - 92.3 \text{ kJ}$

 the number of kilojoules (kJ) released if 100 g of  $\text{HCl}(\text{g})$  is produced, is:
 

A) 235.17

B) 325.17

C) 523.17

D) 253.17

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8. The process of surrounding solute particles by solvent particles is known as:
 

A) dilution

B) formation

C) solvation

D) osmosis

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9. The solubility of -----?----- in liquid is highly affected by changing pressure.
 

A) gases

B) liquids

C) solids

D) salts

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10. If 0.1 mol of solid glucose ( $\text{C}_6\text{H}_{12}\text{O}_6$ ) is dissolved in the same mass of each of the following solvents:
 

Solvent	Q	X	Y	Z
$K_b / ^\circ\text{C molal}^{-1}$	0.4	1.53	1.7	0.5

 the solvent which its boiling point is elevated more is:
 

A) Q

B) X

C) Y

D) Z

---

11. The magnitudes of the molal constant of boiling point elevation ( $K_b$ ) depend on:
 

A) temperature

B) pressure

C) nature of solute

D) nature of solvent

---

12. The aqueous solution with the highest boiling point is:
 

A) 0.1 M HI

B) 0.1 M  $(\text{NH}_4)_3\text{PO}_4$

C) 0.2 M  $\text{C}_2\text{H}_5\text{OH}$

D) 0.1 M  $\text{NH}_4\text{Cl}$



13. If 1 L carbonated water is bottled under pressure of 2.4 atm of  $\text{CO}_2(\text{g})$ , and Henry's law constant is  $3.36 \times 10^{-2} \text{ mol/L atm}$ , the number of grams of dissolved  $\text{CO}_2(\text{g})$  is:  
 A) 5.35                      B) 53.5                      C) 35.5                      D) 3.55

14. At  $30^\circ\text{C}$ , the osmotic pressure, in torr, of 0.108 M aqueous solution of a salt that is assumed to be totally ionized into three ions is:  
 A)  $3.16 \times 10^3$                       B)  $1.63 \times 10^3$                       C)  $6.13 \times 10^3$                       D)  $1.36 \times 10^3$

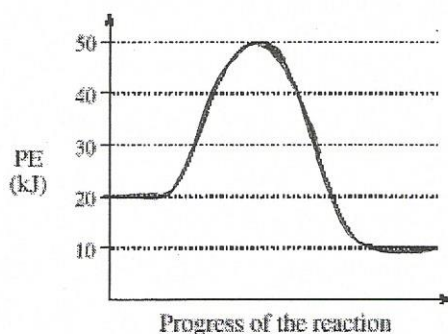
15. The minimum amount of energy required to overcome the energy barrier in a chemical reaction is the:  
 A) activation energy                      B) reaction's enthalpy  
 C) reactants' kinetic energy                      D) reactants' heat content

16. Increasing temperature increases reaction rate because it:  
 A) increases the activation energy                      B) decreases the activation energy  
 C) increases the number of collisions                      D) increases the reaction enthalpy

17. According to the following reaction:  

$$\text{N}_2\text{O}_5(\text{g}) \rightarrow \text{NO}_2(\text{g}) + \text{NO}_3(\text{g})$$
 if 0.8 mol of  $\text{N}_2\text{O}_5(\text{g})$  is initially put in 2 L-reaction vessel and is found to be 0.0125 mol after 2 min, the rate of disappearance of  $\text{N}_2\text{O}_5(\text{g})$ , in M/min, is:  
 A) 0.9169                      B) 0.1969                      C) 0.6919                      D) 0.9961

- 18 From the following reaction potential energy (PE) diagram:



which of the following is correct for the forward reaction:

	$\Delta H/\text{kJ}$	Activation energy, $E_a/\text{kJ}$	Type of reaction
A)	+ 10	40	exothermic
B)	+ 10	30	endothermic
C)	- 10	30	exothermic
D)	- 10	50	endothermic

- 19 In a first order reaction, if the concentration of the reactant changes from 0.1 M to 0.025 M in 40 minutes, the reaction rate, in M/min, when the initial concentration is 0.01 M is:  
 A)  $6.634 \times 10^{-4}$                       B)  $6.346 \times 10^{-4}$                       C)  $4.366 \times 10^{-4}$                       D)  $3.466 \times 10^{-4}$

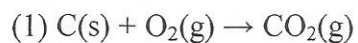
20. For the reaction:  

$$\text{N}_2\text{O}_5(\text{g}) \rightarrow 2\text{NO}_2(\text{g}) + \frac{1}{2}\text{O}_2(\text{g})$$
 if the value of the rate of disappearance of  $\text{N}_2\text{O}_5$  is  $6.25 \times 10^{-3} \text{ mol L}^{-1} \text{ s}^{-1}$ , the rate of appearance of  $\text{NO}_2$  is:  
 A)  $2.15 \times 10^{-2}$                       B)  $1.25 \times 10^{-2}$                       C)  $2.51 \times 10^{-2}$                       D)  $2.51 \times 10^{-2}$

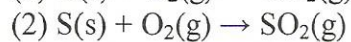
**Second: Solve, in details, the following problems:**

**(FOR CHEM 101 5 marks) (FOR CHEM 103 10 marks)**

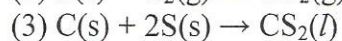
1. Given:



$$\Delta H_1 = -393.5 \text{ kJ}$$

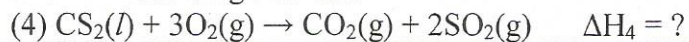


$$\Delta H_2 = -296.8 \text{ kJ}$$



$$\Delta H_3 = +87.9 \text{ kJ}$$

calculate  $\Delta H$  of the following reaction:



2. If dissolving 28.5 g of a sugar in 13.88 mol of water lowers the vapor pressure of water from 17.540 torr to 17.435 torr, calculate the molar mass of the sugar.

3. If a reaction rate constant is  $8.9 \times 10^{-3} \text{ M}^{-1} \text{ s}^{-1}$  at  $3^\circ \text{C}$  and  $7.1 \times 10^{-2} \text{ M}^{-1} \text{ s}^{-1}$  at  $35^\circ \text{C}$ , calculate its activation energy in kJ/mol?



الاسم :- عبدالله غازي عبدالله بن سعدون

الرقم الجامعي :-

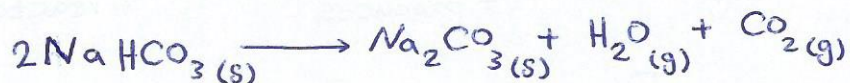
First : Choose the correct answer :-

Q1 :- (B) :- positive and  $H_{CO_2(g)} > H_{CO_2(s)}$

Q2 :- (D) :-  $3K_{(s)} + P_{(s)} + 2O_{2(g)} \longrightarrow K_3PO_{4(s)}$

Q3 :- (A) :- a thermochemical equation .

Q4 :- الكل :-  $R = 8,314 \text{ J mol}^{-1} \text{ K}^{-1}$   
 $\Delta H_{rxn}^{\circ} = 129 \text{ KJ}$  ,  $\Delta E^{\circ} = ?$  ,  $T = 25^{\circ} \text{C} = 298 \text{ K}$



$$\Delta E^{\circ} = \Delta H_{rxn} - \Delta nRT$$

$$\Delta nRT = (n_f - n_i)RT = (2 - 0) \times 8,314 \times 298 = 4955,144 \text{ J}$$

الفرق في عدد  
مولات الغازات

$$\therefore \Delta nRT = 4,955 \text{ KJ}$$

$$\therefore \Delta E^{\circ} = 129 - 4,955 = \boxed{124,04 \text{ KJ}}$$

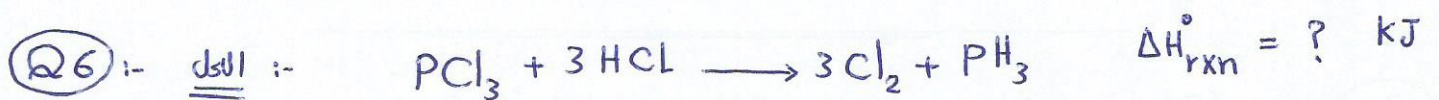
- Q5 :- الكل :-
- mass of metal (m) = 10 g
  - Specific heat ( $C_s$ ) = 0,896 J/g K
  - heat (q) = 313,5 J
  - $T_i = 298$  K
  - $T_f = ?$

$$\therefore q = m \times C_s \times (T_f - T_i)$$

$$\therefore T_f = \frac{q}{m \times C_s} + T_i = \frac{313,5}{10 \times 0,896} + 298 = 332,98 \approx \underline{\underline{333 \text{ K}}}$$

$$\therefore T_f = \boxed{333 \text{ K}}$$

final temperatur



$$\Delta H_f^\circ (\text{PH}_3) = +5,40 \text{ KJ mol}^{-1}$$

$$\Delta H_f^\circ (\text{PCl}_3) = -288,07 \text{ KJ mol}^{-1}$$

$$\Delta H_f^\circ (\text{HCl}) = -92,30 \text{ KJ mol}^{-1}$$

$$\Delta H_f^\circ (\text{Cl}_2) = 0 \text{ KJ mol}^{-1}$$

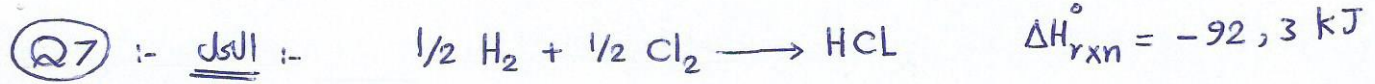
$$\Delta H_{\text{rxn}}^\circ = \sum n \Delta H_f^\circ \text{ products} - \sum m \Delta H_f^\circ \text{ reactants}$$

$$= [(3 \times 0) + (1 \times 5,40)] - [(1 \times -288,07) + (3 \times -92,30)]$$

$$= 5,40 + 564,97 = \underline{\underline{570,37 \text{ KJ}}}$$

$$\therefore \Delta H_{\text{rxn}}^\circ = \boxed{570,37 \text{ KJ}}$$





$\Delta H_{\text{rxn}}^\circ = -92,3 \text{ kJ} \longrightarrow$  هي حرارة تكوين مول واحد من (HCL)

$$\therefore 1 \text{ mol (HCL)} \longrightarrow 36,45 \text{ g/mol}$$

$$\therefore 1 \text{ mol (HCL)} \longrightarrow -92,3 \text{ kJ}$$

$$\therefore 36,45 \text{ g} \longrightarrow -92,3 \text{ kJ}$$

$$100 \text{ g} \longrightarrow -X$$

$$\therefore X = \frac{100 \times 92,3}{36,45} = \boxed{253,17 \text{ kJ}}$$

Q8 :- (C) Solvation .

Q9 :- (A) :- gases .

Q10 :- بما أن المولالية (m) متساوية في جميع المذيبات (X/Y/Z) :-  
لأن لهم نفس الكتلة .

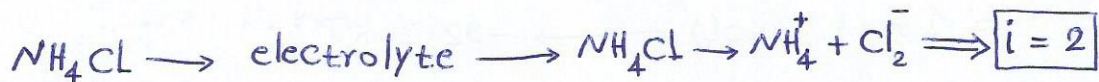
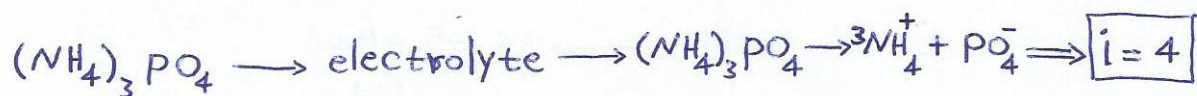
$K_b \longleftarrow$  ثابت للجميع

⊗ إذا ترفع درجة الغليان عندما يكون  $(\Delta T_b)$  هو الأكبر

$\therefore$  boiling point is elevated more is (Y) = 1,7

$\boxed{Y}$  :- الجواب هو : (C) :-

Q11 :- (D) :- nature of solvent .



\* ( ترفع درجة الغليان عندما يكون (i) هو الأكبر ) .

∴ الجواب الصحيح هو :- (B) ←  $0,1M (NH_4)_3PO_4$

Q13 :- الكل :- نطلع ذوبانية الغاز ( $S_g$ )

$$S_g = k P_g = 3,36 \times 10^{-2} \times 2,4 = \underline{\underline{0,08064 M \text{ or mol/L}}}$$

\* بما أن ذوبانية الغاز ( $S_g$ ) يعبر عنها بوحدة التركيز (M)

$$\therefore S_g = \text{Molarity (M)}$$

$$\therefore M = \frac{\text{Number of solute}}{\text{Volume of solution in (Litter)}}$$

$$\therefore n = M \times V = 0,08064 \times 1 = 0,08064 \text{ mole}$$

$$\therefore n = \frac{m}{M_{ut}} \Rightarrow m = n \times M_{ut} = 0,08064 \times 44 = \underline{\underline{3,55 g}}$$

∴ the number of grams (m) of dissolved  $CO_2$  is =  $\boxed{3,55g}$



Q14 :- الحل :-  $T = 30^\circ\text{C} = 303\text{ K}$  ,  $M = 0,108\text{ M}$   
 $i = 3\text{ ions}$  ,  $R = 0,0821\text{ atm L mol}^{-1}\text{ K}^{-1}$   
 $R = 0,0821 \times 760 = 62,396$   
 $\Pi = MRTi = 0,108 \times 62,396 \times 303 \times 3$   
 $= 6,13 \times 10^3\text{ torr}$

$\therefore$  the osmotic pressure ( $\Pi$ ) =  $6,13 \times 10^3\text{ torr}$

Q15 :- (A) :- activation energy.

Q16 :- (C) :- increases the number of collisions .

Q17 :- الحل :-  $n_i = 0,8\text{ mol}$  ,  $n_f = 0,0125\text{ mol}$   
 $V = 2\text{ L}$  ,  $\Delta t = 2\text{ min}$

$\therefore \text{Rate} = \frac{\Delta[M]}{\Delta t}$    
 مقدار التغير في التركيز  $\rightarrow$    
 مقدار التغير في الزمن  $\rightarrow$

$\therefore \Delta[M]$  نوجد

$M_i = \frac{n_i}{V} = \frac{0,8}{2} = 0,4\text{ M}$

$M_f = \frac{n_f}{V} = \frac{0,0125}{2} = 0,00625\text{ M}$

$\therefore \Delta[M] = 0,4 - 0,00625 = 0,39375\text{ M}$

$\therefore \text{Rate} = \frac{\Delta[M]}{\Delta t} = \frac{0,39375}{2} = 0,1969\text{ M/min}$

- Q 18 :- إذا كان التفاعل في الاتجاه الأمامي (forward) فإن قيمة  $(\Delta H)$  تكون (-) لأن التفاعل طارد للحرارة (exothermic).

الاجواب الصحيح هو :- (C)

	$\Delta H / \text{kJ}$	Activation energy , $E_a / \text{kJ}$	Type of reaction
C)	-10	30	exothermic

- Q 19 :- الجد :-  $[A]^0 = 0,1 \text{ M}$  ,  $[A]_t = 0,025 \text{ M}$   
 $t = 40 \text{ minutes}$

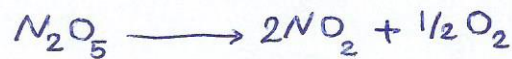
نطلع (k) ثابت سرعة التفاعل :-

$$\ln \frac{[A]_t}{[A]^0} = -kt \Rightarrow \ln \left( \frac{0,025}{0,1} \right) = -k \times 40$$

$$\therefore k = 0,03466 \text{ min}^{-1}$$

$$\therefore \text{Rate} = k [A] = 0,03466 \times 0,01 = 3,466 \times 10^{-4} \text{ M/min}$$

- Q 20 :- الجد :- Rate of  $\text{N}_2\text{O}_5 = 6,25 \times 10^{-3} \text{ mol L}^{-1} \text{ s}^{-1}$



$$\text{Rate} = -\frac{\Delta[\text{N}_2\text{O}_5]}{\Delta t} = \frac{1}{2} \frac{\Delta[\text{NO}_2]}{\Delta t}$$

$$\therefore \frac{\Delta[\text{N}_2\text{O}_5]}{\Delta t} = 6,25 \times 10^{-3}$$

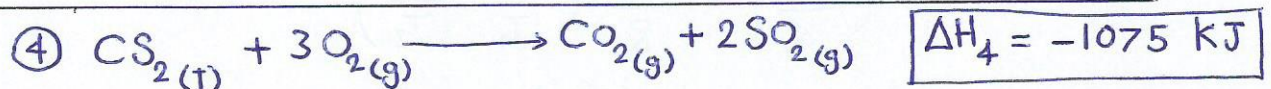
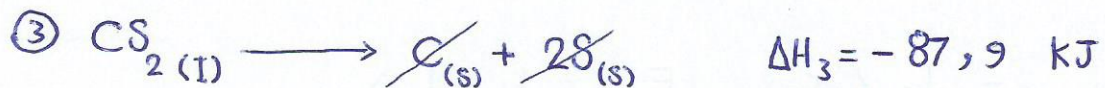
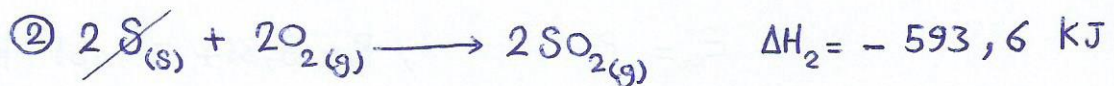
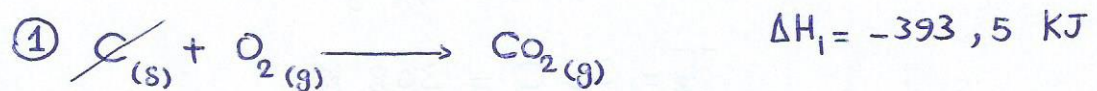
$$\therefore \frac{1}{2} \frac{\Delta[\text{NO}_2]}{\Delta t} = 6,25 \times 10^{-3}$$

$$\therefore \text{Rate of } \text{NO}_2 = 6,25 \times 10^{-3} \times 2 = 1,25 \times 10^{-2} \text{ mol L}^{-1} \text{ s}^{-1}$$



## Second :- Solve :-

Q1 :- الكل :- نقسب المعادلة (2) في (2) ونعكس المعادلة (3) ثم نأجمع المعادلات الثلاث :-



Q2 :- الكل :- - mass of sugar = 28,5 g

- No. mole of water = 13,88 mol

- Vapor pressure of water ( $P_w$ ) = 17,435 torr

- Vapor pressure of pure water ( $P_w^\circ$ ) = 17,540 torr

$$\therefore P_w = X_w P_w^\circ$$

$$\therefore P_w = \left( \frac{n_w}{n_w + n_s} \right) P_w^\circ$$

$\nwarrow$  عدد مولات الماء       $\swarrow$  عدد مولات السكر

$$17,435 = \left( \frac{13,88}{13,88 + n_s} \right) \times 17,540$$

$$\therefore n_s = \frac{13,88}{0,99401} - 13,88 = 0,0836 \text{ mol}$$

$$\therefore n_s = \frac{m}{M_{ut}} \Rightarrow M_{ut} = \frac{m}{n_s} = \frac{28,5}{0,0836} = \underline{\underline{340,91}} \approx \underline{\underline{341 \text{ g/mol}}}$$

$\therefore$  the molar mass of the sugar  $\approx \boxed{341 \text{ g/mol}}$

Q3 :- المطلوب :-

$$k_1 = 8,9 \times 10^{-3} \text{ M}^{-1} \text{ s}^{-1}$$

$$T_1 = 3 \text{ C}^\circ = 276 \text{ K}$$

$$k_2 = 7,1 \times 10^{-2} \text{ M}^{-1} \text{ s}^{-1}$$

$$T_2 = 35 \text{ C}^\circ = 308 \text{ K}$$

$$E_a = ? , R = 8,314 \text{ J mol}^{-1} \text{ K}^{-1}$$

$$\therefore \ln \left( \frac{k_2}{k_1} \right) = \frac{E_a}{R} \left( \frac{1}{T_1} - \frac{1}{T_2} \right)$$

$$\begin{aligned} \therefore E_a &= \frac{R \ln \left( \frac{k_2}{k_1} \right)}{\frac{1}{T_1} - \frac{1}{T_2}} \\ &= \frac{8,314 \ln \left( \frac{7,1 \times 10^{-2}}{8,9 \times 10^{-3}} \right)}{\frac{1}{276} - \frac{1}{308}} = \frac{17,26509}{3,7643 \times 10^{-4}} \end{aligned}$$

$$= 45865,34 \text{ J/mol}$$

$$\therefore E_a = \boxed{45,86 \text{ kJ/mol}}$$