

**Q1. 21 January Shift 1**

Pre-exponential factors of two different reactions of same order are identical. Let activation energy of first reaction exceeds the activation energy of second reaction by  $20 \text{ kJ mol}^{-1}$ . If  $k_1$  and  $k_2$  are the rate constants of first and second reaction respectively at  $300 \text{ K}$ , then  $\ln \frac{k_2}{k_1}$  will be \_\_\_\_\_. (nearest integer) [ $R = 8.3 \text{ J K}^{-1} \text{ mol}^{-1}$ ]

**Q2. 21 January Shift 2**

Decomposition of A is a first order reaction at  $T(\text{K})$  and is given by  $\text{A}(\text{g}) \longrightarrow \text{B}(\text{g}) + \text{C}(\text{g})$ . In a closed  $1 \text{ L}$  vessel,  $1 \text{ bar A}(\text{g})$  is allowed to decompose at  $T(\text{K})$ . After  $100$  minutes, the total pressure was  $1.5 \text{ bar}$ . What is the rate constant ( $\text{in min}^{-1}$ ) of the reaction? ( $\log 2 = 0.3$ )

- (1)  $6.9 \times 10^{-3}$  (2)  $6.9 \times 10^{-4}$  (3)  $6.9 \times 10^{-1}$  (4)  $6.9 \times 10^{-2}$

**Q3. 22 January Shift 1**

$\text{A} \rightarrow \text{product}$  (First order reaction). Three sets of experiment were performed for a reaction under similar experimental conditions:

Run 1  $\Rightarrow$   $100 \text{ mL}$  of  $10 \text{ M}$  solution of reactant A

Run 2  $\Rightarrow$   $200 \text{ mL}$  of  $10 \text{ M}$  solution of reactant A

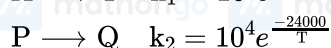
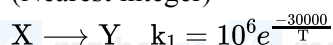
Run 3  $\Rightarrow$   $100 \text{ mL}$  of  $10 \text{ M}$  solution of reactant A +  $100 \text{ mL}$  of  $\text{H}_2\text{O}$  added.

The correct variation of rate of reaction is

- (1) Run 1 < Run 2 < Run 3 (2) Run 3 < Run 1 < Run 2  
(3) Run 3 < Run 1 = Run 2 (4) Run 1 = Run 2 = Run 3

**Q4. 22 January Shift 1**

The temperature at which the rate constants of the given below two gaseous reactions become equal is \_\_\_\_ K. (Nearest integer)



Given:  $\ln 10 = 2.303$

**Q5. 22 January Shift 2**

Correct statements regarding Arrhenius equation among the following are :

- A. Factor  $e^{-E_a/RT}$  corresponds to fraction of molecules having kinetic energy less than  $E_a$ .  
B. At a given temperature, lower the  $E_a$ , faster is the reaction.  
C. Increase in temperature by about  $10^\circ\text{C}$  doubles the rate of reaction.  
D. Plot of  $\log k$  vs  $\frac{1}{T}$  gives a straight line with slope  $= -\frac{E_a}{R}$ .

Choose the correct answer from the options given below :

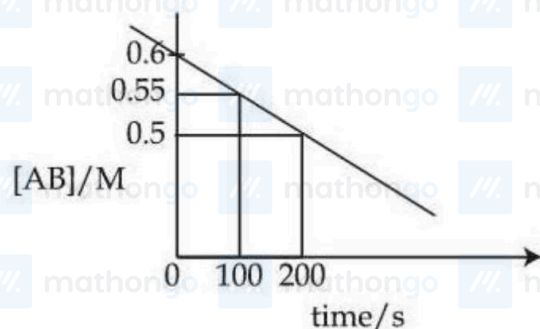
- (1) B and D Only (2) A and B Only (3) B and C Only (4) A and C Only

Q6. 22 January Shift 2

Consider  $A \xrightarrow{k_1} B$  and  $C \xrightarrow{k_2} D$  are two reactions. If the rate constant ( $k_1$ ) of the  $A \rightarrow B$  reaction can be expressed by the following equation  $\log_{10} k = 14.34 - \frac{1.5 \times 10^4}{T/K}$  and activation energy of  $C \rightarrow D$  reaction ( $E_{a2}$ ) is  $\frac{1}{5}$  th of the  $A \rightarrow B$  reaction ( $E_{a1}$ ), then the value of ( $E_{a2}$ ) is \_\_\_\_  $\text{kJmol}^{-1}$ . (Nearest Integer)

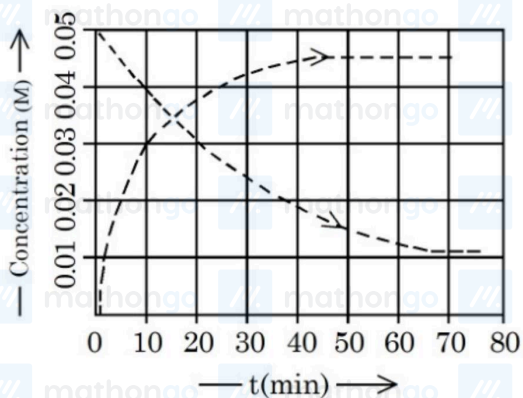
Q7. 23 January Shift 1

For the thermal decomposition of reactant  $AB(g)$ , the following plot is constructed.



The half life of the reaction is 'x' min.  $x =$  \_\_\_\_ min. (Nearest integer)

Q8. 23 January Shift 2



Given above is the concentration vs time plot for a dissociation reaction :  $A \rightarrow nB$ . Based on the data of the initial phase of the reaction (initial 10 min), the value of  $n$  is \_\_\_\_.

- (1) 3 (2) 2 (3) 4 (4) 5

**Q9. 23 January Shift 2**

Observe the following reactions at T(K).

I.  $A \rightarrow \text{products}$ .

II.  $5\text{Br}^-(\text{aq}) + \text{BrO}_3^-(\text{aq}) + 6\text{H}^+(\text{aq}) \rightarrow 3\text{Br}_2(\text{aq}) + 3\text{H}_2\text{O}(\text{l})$

Both the reactions are started at 10.00 am. The rates of these reactions at 10.10 am are same. The value of  $-\frac{\Delta[\text{Br}^-]}{\Delta t}$  at 10.10 am is  $2 \times 10^{-4} \text{ mol L}^{-1} \text{ min}^{-1}$ . The concentration of A at 10.10 am is  $10^{-2} \text{ mol L}^{-1}$ . What is the first order rate constant (in  $\text{min}^{-1}$ ) of reaction I ?

- (1)  $10^{-2}$  (2)  $10^{-3}$  (3)  $2 \times 10^{-3}$  (4)  $4 \times 10^{-3}$

**Q10. 24 January Shift 1**

At  $27^\circ\text{C}$  in presence of a catalyst, activation energy of a reaction is lowered by  $10 \text{ kJ mol}^{-1}$ . The logarithm of ratio of  $\frac{k(\text{catalysed})}{k(\text{uncatalysed})}$  is....

(Consider that the frequency factor for both the reactions is same)

- (1) 1.741 (2) 17.41 (3) 3.482 (4) 0.1741

**Q11. 24 January Shift 2**

The half-life of  $^{65}\text{Zn}$  is 245 days. After  $x$  days, 75% of original activity remained. The value of  $x$  in days is \_\_\_\_.

(Nearest integer)

(Given:  $\log 3 = 0.4771$  and  $\log 2 = 0.3010$ )

**Q12. 28 January Shift 1**

An organic compound undergoes first order decomposition. The time taken for decomposition to  $(\frac{1}{8})^{\text{th}}$  and  $(\frac{1}{10})^{\text{th}}$  of its initial concentration are  $t_{1/8}$  and  $t_{1/10}$  respectively. What is the value of  $\frac{t_{1/8}}{t_{1/10}} \times 10$  ? ( $\log 2 = 0.3$ )

- (1) 3 (2) 9 (3) 0.9 (4) 30

**Q13. 28 January Shift 2**

$A \rightarrow B$  (first reaction)

$C \rightarrow D$  (second reaction)

Consider the above two first-order reactions. The rate constant for first reaction at 500 K is double of the same at 300 K. At 500 K, 50% of the reaction becomes complete in 2 hour. The activation energy of the second reaction is half of that of first reaction. If the rate constant at 500 K of the second reaction becomes double of the rate constant of first reaction at the same temperature; then rate constant for the second reaction at 300 K is \_\_\_\_  $\times 10^{-1} \text{ hour}^{-1}$  (nearest integer).

**ANSWER KEYS**

1. 8 2. (1) 3. (3) 4. 1303 5. (3) 6. 57 7. 10 8. (1)

9. (4) 10. (1) 11. 102 12. (2) 13. 5