

CBSE Class 12 physics
Important Questions
Chapter 4
Chemical Kinetics

3 Marks Questions

1. For the reaction $A+B \rightarrow C+D$, the rate of reaction doubles when the concentration of A doubles, provided the concentration of B is constant. To what order does A enter into the rate expression?

Ans. $A+B \rightarrow C+D$

$$Rate[A]^x$$

Rate = 1 when $[A] = 1$ ----- 1)

Rate = 2 when $[A] = 2$ -----2)

Dividing equation 2) by 1)

$$\frac{2}{1} \propto \frac{(2)^x}{1^x}$$

$$2^1 \propto (2)^x$$

$$\therefore x = 1$$

The reaction is first order reaction.

2. A chemical reaction $2A \rightleftharpoons 4B+C$ in gas phase occurs in a closed vessel. The concentration of B is found to be increased by $5 \times 10^{-3} \text{ mole } L^{-1}$ in 10 second. Calculate (i) the rate of appearance of B (ii) the rate of disappearance of A?

Ans. $2A \rightarrow 4B + C$

$$-\frac{1}{2} \frac{d[A]}{dt} = \frac{1}{4} \frac{d[B]}{dt} = \frac{d[C]}{dt}$$

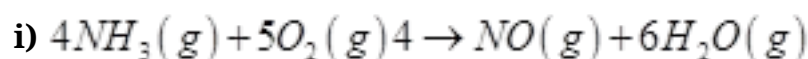
i) Rate of disappearance of B

$$= \frac{5 \times 10^{-3}}{10^5} \text{ mol L}^{-1} = 5 \times 10^{-4} \text{ mol L}^{-1} \text{ s}^{-1}$$

$$\text{ii) } \frac{-d[A]}{dt} = \frac{2}{4} \frac{d[B]}{dt} = \frac{1}{2} \frac{d[B]}{dt}$$

$$= \frac{1}{2} \times 5 \times 10^{-4} \text{ mol L}^{-1} \text{ s}^{-1} = 2.5 \times 10^{-4} \text{ mol L}^{-1} \text{ s}^{-1}$$

3. For the following reactions, write the rate of reaction expression in terms of reactants and products?



Ans.

In terms of reactant	In terms of products
i) $R_1 = -\frac{1}{4} \frac{\Delta[\text{NH}_3]}{\Delta t}$	$R_3 = \frac{1}{4} \frac{\Delta[\text{NO}]}{\Delta t}$
$R_2 = -\frac{1}{5} \frac{\Delta[\text{O}_2]}{\Delta t}$	$R_4 = \frac{1}{6} \frac{\Delta[\text{H}_2\text{O}]}{\Delta t}$

$$\frac{1}{4} R_1 = \frac{1}{5} R_2 = \frac{1}{4} R_3 = \frac{1}{6} R_4$$

$$-\frac{1}{4} \frac{\Delta[\text{NH}_3]}{\Delta t} = -\frac{1}{5} \frac{\Delta[\text{O}_2]}{\Delta t} = \frac{1}{4} \frac{\Delta[\text{NO}]}{\Delta t} = \frac{1}{6} \frac{\Delta[\text{H}_2\text{O}]}{\Delta t}$$

II) In terms of reactant	In terms of product

$R_1 = - \frac{\Delta[N_2O_5]}{\Delta t}$	$R_2 = \frac{\Delta[NO_2]}{\Delta t}$
	$R_3 = \frac{\Delta[O_2]}{\Delta t}$

$$\frac{1}{2}R_1 = \frac{1}{2}R_2 = R_3 = \frac{\Delta[N_2O_5]}{\Delta t} = \frac{1}{2} \frac{\Delta[NO_2]}{\Delta t} = \frac{\Delta[O_2]}{\Delta t}$$

4. The reaction $2N_2O_5(g) \rightarrow 2NO_2(g) + O_2(g)$ was studied and the following data were collected :

S.no (mol/L/min)	$[N_2O_5] \text{ mol L}^{-1}$	Rate of disappearance of $[N_2O_5]$ (mol/L/min)
1.	1.13×10^{-2}	34×10^{-5}
2.	0.84×10^{-2}	25×10^{-5}
3.	0.62×10^{-2}	18×10^{-5}

Determine

i) The order

ii) The rate law.

iii) Rate constant for the reaction.

Ans. Let the order of reaction be x

$$Rate = K[N_2O_5]^x$$

i) From the data -

$$34 \times 10^{-5} = (1.13 \times 10^{-2})^x \text{ -----1)}$$

$$25 \times 10^{-5} = (0.84 \times 10^{-2})^x \text{ -----2)}$$

$$18 \times 10^{-5} = (0.62 \times 10^{-2})^x \text{ -----3)}$$

Dividing 1) by 2)

$$\frac{34 \times 10^{-5}}{25 \times 10^{-5}} = \left(\frac{1.13 \times 10^{-2}}{0.84 \times 10^{-2}} \right)^x$$

$$(1.36) = (1.35)^x$$

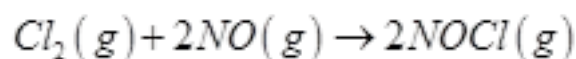
$$X=1$$

The order of reaction with respect with respect to N_2O_5 is 1

ii) Rate law $R = K [N_2O_5]$

iii) Rate constant , $K = \frac{\text{Rate}}{[N_2O_5]} = \frac{18 \times 10^{-5} \text{ mol/L/min}}{0.62 \times 10^{-2} \text{ mol/L}} = 0.29 \text{ min}^{-1}$

5. The following experimental data was collected for the reaction:



Trial	Intial conc. Of Cl_2 (mol / L)	[NO] mol/L	Initial Rate,(mol/L/s)
1	0.10	0.010	1.2×10^{-4}
2	0.10	0.030	10.8×10^{-4}
3	0.20	0.030	21.6×10^{-4}

Construct the rate equation for the reaction.

Ans. Order of NO is 2

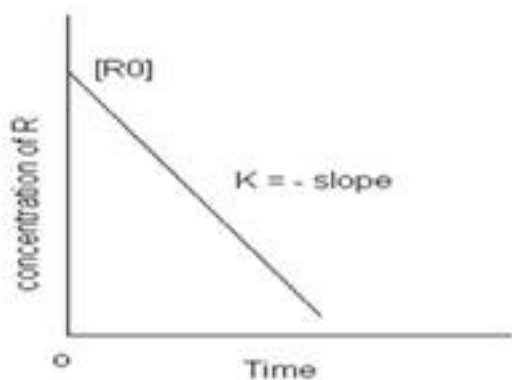
$$\text{Rate law} = K [Cl_2] [NO]^2$$

6. Draw a graph for

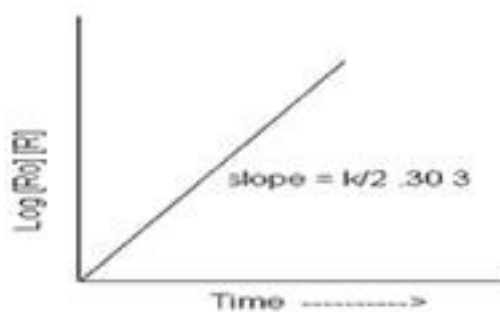
a) Concentration of reactant against time for a zero order reaction.

b) $\log [R_0]/[R]$ against time for a first order reaction.

Ans.



(a)



(b)

7. In general it is observed that the rate of a chemical reaction doubles with every 10° rise in temperature. If this generalization holds for a reaction in the temperature range 295K to 305K, what would be the activation energy for this reaction?

$$(R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1})$$

Ans. $T_1 = 295\text{K}$ $T_2 = 305\text{K}$

$$E_a = 2.303 R \left[\frac{T_2 T_1}{T_2 - T_1} \right] \left[\log \frac{k_2}{k_1} \right]$$

$$K_2 = 2k_1$$

$$E_a = 2.303 \times 8.314 \times \left[\frac{305 \times 295}{305 - 295} \right] \log \frac{2k_1}{k_1}$$

$$= 2.303 \times 8.314 \times 8997.5 \text{ K} \log 2$$

$$= 51855.2 \text{ J/mol} \quad (\log 2 = 0.3010)$$

8. The rate constant for a reaction is $1.5 \times 10^7 \text{ s}^{-1}$ at 50°C and $4.5 \times 10^7 \text{ s}^{-1}$ at 100°C .

Calculate the value of activation energy for the reaction $R = 8.314 \text{ JK}^{-1} \text{ mol}^{-1}$?

Ans. $\log \frac{k_2}{k_1} = \frac{E_a}{2.303R} \left(\frac{T_2 - T_1}{T_1 T_2} \right)$

$$\log \frac{4.5 \times 10^7}{1.5 \times 10^7} = \frac{E_a}{2.303 \times 8.314} \left(\frac{373 - 323}{373 \times 323} \right)$$

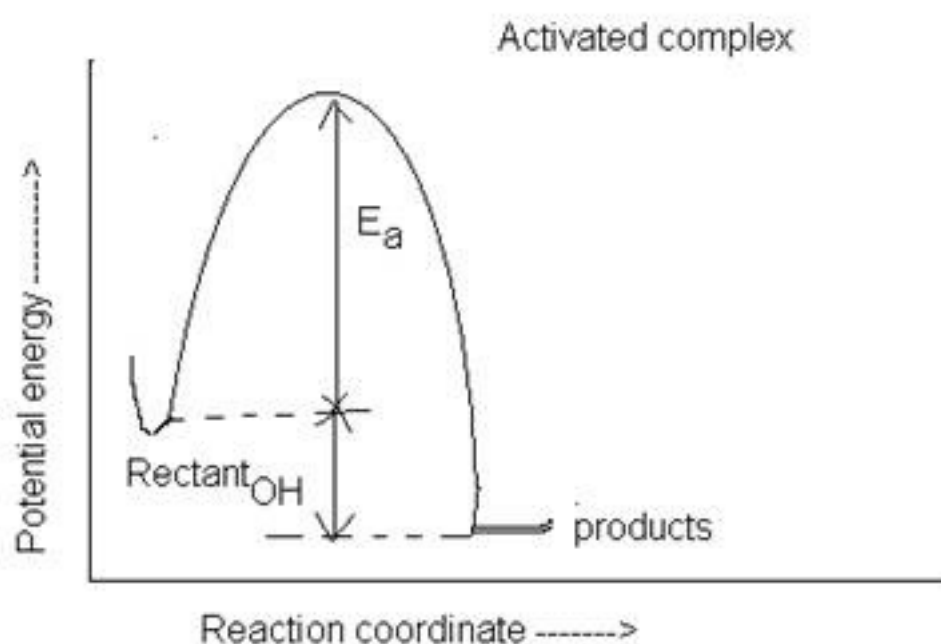
$$\log 1.5 = \frac{E_a}{2.303 \times 8.314} \left(\frac{50}{373 \times 323} \right)$$

$$E_a = \left(\frac{2.303 \times 8.314 \times 373 \times 323}{50} \right) \times \log 1.5$$

$$= 22 \text{ KJ / mol}$$

9. Plot a graph showing variation of potential energy with reaction. coordinate?

Ans.



10. The conversion of molecules X to Y follows second order kinetics. If concentration

of X is increased to three times how will it affect the rate of formation of Y?

Ans. The reaction $X \rightarrow Y$ follows second order kinetics.

Therefore, the rate equation for this reaction will be:

$$\text{Rate} = k[X]^2 \quad (1)$$

Let $[X] = a \text{ mol}^{-1}$, then equation (1) can be written as:

$$\text{Rate}_1 = k(a)_2$$

$$= ka_2$$

If the concentration of X is increased to three times, then $[X] = 3a \text{ mol L}^{-1}$

Now, the rate equation will be:

$$\text{Rate} = k(3a)_2$$

$$= 9(ka^2)$$

Hence, the rate of formation will increase by 9 times.

11. A first order reaction has a rate constant $1.15 \times 10^{-3} \text{ s}^{-1}$. How long will 5 g of this reactant take to reduce to 3 g?

Ans. From the question, we can write down the following information:

Initial amount = 5 g

Final concentration = 3 g

Rate constant = $1.15 \times 10^{-3} \text{ s}^{-1}$

We know that for a 1st order reaction,

$$t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$$

$$= \frac{2.303}{1.15 \times 10^{-3}} \log \frac{5}{3}$$

$$= \frac{2.303}{1.15 \times 10^{-3}} \times 0.2219$$

$$= 444.38 \text{ s}$$

$$= 444 \text{ s (approx)}$$

12. For the reaction $R \rightarrow P$, the concentration of a reactant changes from 0.03 M to 0.02 M in 25 minutes. Calculate the average rate of reaction using units of time both in minutes and Pseconds.

Ans. Average rate of reaction $-\frac{\Delta[R]}{\Delta t}$

$$= -\frac{[R]_2 - [R]_1}{t_2 - t_1}$$

$$= -\frac{0.02 - 0.03}{25} \text{ M min}^{-1}$$

$$= -\frac{-0.01}{25} \text{ M min}^{-1}$$

$$= 4 \times 10^{-4} \text{ M min}^{-1}$$

$$= \frac{4 \times 10^{-4}}{60} \text{ M s}^{-1}$$

$$= 6.67 \times 10^{-6} \text{ M s}^{-1}$$

13. For the reaction: $2A + B \rightarrow A_2B$ the rate = $k[A][B]_2$ with $k = 2.0 \times 10^{-6} \text{ mol}^{-2} \text{ L}^2 \text{ s}^{-1}$. Calculate the initial rate of the reaction when $[A] = 0.1 \text{ mol L}^{-1}$, $[B] = 0.2 \text{ mol L}^{-1}$. Calculate the rate of reaction after $[A]$ is reduced to 0.06 mol L^{-1} .

Ans. The initial rate of the reaction is

$$\text{Rate} = k[A][B]_2$$

$$= (2.0 \times 10^{-5} \text{ mol}^{-2} \text{ L}^2 \text{ s}^{-1}) (0.1 \text{ mol L}^{-1}) (0.2 \text{ mol L}^{-1})^2$$

$$= 8.0 \times 10^{-9} \text{ mol}^{-2} \text{ L}^2 \text{ s}^{-1}$$

When [A] is reduced from 0.1 mol L^{-1} to 0.06 mol L^{-1} , the concentration of A reacted = $(0.1 - 0.06) \text{ mol L}^{-1} = 0.04 \text{ mol L}^{-1}$

$$\text{Therefore, concentration of B reacted} = \frac{1}{2} \times 0.04 \text{ mol L}^{-1} = 0.02 \text{ mol L}^{-1}$$

Then, concentration of B available, $[B] = (0.2 - 0.02) \text{ mol L}^{-1}$

$$= 0.18 \text{ mol L}^{-1}$$

After [A] is reduced to 0.06 mol L^{-1} , the rate of the reaction is given by,

$$\text{Rate} = k[A][B]_2$$

$$= (2.0 \times 10^{-5} \text{ mol}^{-2} \text{ L}^2 \text{ s}^{-1}) (0.06 \text{ mol L}^{-1}) (0.18 \text{ mol L}^{-1})^2$$

$$= 3.89 \text{ mol L}^{-2} \text{ s}^{-1}$$

14. A reaction is second order with respect to a reactant. How is the rate of reaction affected if the concentration of the reactant is

(i) doubled (ii) reduced to half?

Ans. Let the concentration of the reactant be $[A] = a$

$$\text{Rate of reaction, } R = k[A]_2$$

$$= ka^2$$

(i) If the concentration of the reactant is doubled, i.e. $[A] = 2a$, then the rate of the reaction

would be

$$R' = k(2a)^2$$

$$= 4ka^2$$

$$= 4R$$

Therefore, the rate of the reaction would increase by 4 times.

(ii) If the concentration of the reactant is reduced to half, i.e. $[A] = \frac{1}{2}a$, then the rate of the

reaction would be $R' = k\left(\frac{1}{2}a\right)^2$

$$= \frac{1}{4}ka^2$$

$$= \frac{1}{4}R$$

Therefore, the rate of the reaction would be reduced to $\frac{1}{4}$ th

15. A reaction is first order in A and second order in B.

(i) Write the differential rate equation.

(ii) How is the rate affected on increasing the concentration of B three times?

(iii) How is the rate affected when the concentrations of both A and B are doubled?

Ans. (i) The differential rate equation will be

$$-\frac{d[R]}{dt} = k[A][B]^2$$

(ii) If the concentration of B is increased three times, then

$$-\frac{d[R]}{dt} = k[A][3B]^2$$
$$= 9.k[A][B]^2$$

Therefore, the rate of reaction will increase 9 times.

(iii) When the concentrations of both A and B are doubled,

$$-\frac{d[R]}{dt} = k[A][B]^2$$
$$= k[2A][2B]^2$$
$$= 8.k[A][B]^2$$

Therefore, the rate of reaction will increase 8 times.

16. Calculate the half-life of a first order reaction from their rate constants given below:

(i) 200 s^{-1} **(ii)** 2 min^{-1} **(iii)** 4 years⁻¹

Ans. (i) Half life, $t_{1/2} = \frac{0.693}{k}$

$$= \frac{0.693}{200\text{ min}^{-1}}$$
$$= 3.4 \times 10^{-3}\text{ s (approximately)}$$

(ii) Half life, $t_{1/2} = \frac{0.693}{k}$

$$= \frac{0.693}{2\text{ min}^{-1}}$$
$$= 0.35\text{ min (approximately)}$$

(iii) Half life, $t_{1/2} = \frac{0.693}{k}$

$$= \frac{0.693}{4 \text{ years}^{-1}}$$

$$= 0.173 \text{ years (approximately)}$$

17. The half-life for radioactive decay of ^{14}C is 5730 years. An archaeological artifact wood had only 80% of the ^{14}C found in a living tree. Estimate the age of the sample.

Ans. Here, $k = \frac{0.693}{t_{1/2}}$

$$= \frac{0.693}{5730} \text{ years}^{-1}$$

It is known that,

$$t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$$

$$= \frac{2.303}{\frac{0.693}{5730}} \log \frac{100}{80}$$

$$= 1845 \text{ years (approximately)}$$

Hence, the age of the sample is 1845 years.

18. The rate constant for a first order reaction is 60 s^{-1} . How much time will it take to reduce the initial concentration of the reactant to its 1/16th value?

Ans. It is known that,

$$t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$$

$$= \frac{2.303}{60\text{S}^{-1}} \log \frac{1}{\frac{1}{16}}$$

$$= \frac{2.303}{60\text{S}^{-1}} \log 16$$

$$= 4.6 \times 10^{-2} \text{ s (approximately)}$$

Hence, the required time is $4.6 \times 10^{-2} \text{ s}$.

19. A first order reaction takes 40 min for 30% decomposition. Calculate $t_{1/2}$.

Ans. For a first order reaction,

$$t_1 = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$$

$$k = \frac{2.303}{40\text{min}} \log \frac{100}{100 - 30}$$

$$= \frac{2.303}{40\text{min}} \log \frac{10}{7}$$

$$= 8.918 \times 10^{-3} \text{ min}^{-1}$$

Therefore, $t_{1/2}$ of the decomposition reaction is

$$t_{1/2} = \frac{0.693}{k}$$

$$= \frac{0.693}{8.918 \times 10^{-3}} \text{ min}$$

$$= 77.7 \text{ min (approximately)}$$

20. Consider a certain reaction $A \rightarrow \text{Products}$ with $k = 2.0 \times 10^{-2} \text{ s}^{-1}$. Calculate the concentration of A remaining after 100 s if the initial concentration of A is 1.0 mol L^{-1} .

Ans. $k = 2.0 \times 10^{-2} \text{ s}^{-1}$

$T = 100 \text{ s}$

$[A]_0 = 1.0 \text{ mol L}^{-1}$

Since the unit of k is s^{-1} , the given reaction is a first order reaction.

Therefore, $k = \frac{2.303}{t} \log \frac{[A]_0}{[A]}$

$$2.0 \times 10^{-2} \text{ s}^{-1} = \frac{2.303}{\log s} \log \frac{1.0}{[A]}$$

$$2.0 \times 10^{-2} \text{ s}^{-1} = \frac{2.303}{\log s} (-\log[A])$$

$$-\log[A] = \frac{2.0 \times 10^{-2} \times 100}{2.303}$$

$$[A] = \text{antilog} \left(-\frac{2.0 \times 10^{-2} \times 100}{2.303} \right)$$

$$= 0.135 \text{ mol L}^{-1} \text{ (approximately)}$$

Hence, the remaining concentration of A is 0.135 mol L^{-1} .