

Rate law and Rate constant

74. (b) It is an bimolecular and second order reaction.

75. (d) $\frac{dx}{dt} = (HI)^2$, then order of reaction = 2

76. (b) Rate = $K(\text{Sucrose})(H_2O)^0$

77. (b) Because in this reaction one molecule of N_2O_5 is used.

78. (c) Integrated velocity equation for first order reaction is: $k = \frac{2.303}{t} \log \frac{(A)_0}{(A)}$

79. **Definition of reaction order:**

- The order of a reaction is determined experimentally and depends on the **rate law** with respect to reactant concentrations.

Effect of surface area:

- Increasing the surface area of a solid reactant **increases the rate of reaction** because more particles are exposed.
- However, the **order of reaction** is a constant value for a given reaction and **does not change** just because the surface area changes.

Conclusion:

- Rate increases with surface area, but **order remains constant**.

Answer: (c) Remain constant

80. (c) $t_{\frac{1}{2}} = \frac{2.303 \log 2}{K} = \frac{0.693}{K}$



81. (b) Inversion of sugar is a pseudo-unimolecular reaction.

82. (b) For 1st order reaction

$$\log(a-x) = \log a - \frac{kt}{2.303}$$

$$y = c + mx$$

83. (c) $t_{\frac{1}{2}} = \frac{0.693}{k} = \frac{0.693}{3.46 \times 10^{-3}} = 200 \text{ min}$

84. (a) $\frac{1}{\text{Time}} \times (\text{conc.}) = \frac{\text{moles l}^{-1}}{\text{Time}} = \text{moles l}^{-1} \text{ time}^{-1}$ for zero order reaction.

85. (a) The order of reaction is $\frac{3}{2}$ and molecularity is 2.

87. (c) Molecularity can never be fractional.

88. (a) As doubling the initial conc. doubles the rate of reaction, order = 1.

89. (c) When B is in excess, it becomes a pseudo-unimolecular reaction.

90. (c) $k = \frac{0.693}{t_{\frac{1}{2}}} = \frac{0.693}{69.35} = 9.99 \times 10^{-3} = 0.01 \text{ s}^{-1}$

91. (b) $k = \frac{0.693}{24} \text{ hr}^{-1} = \frac{2.303}{96} \log \frac{10}{a-x}$

$$\text{or } \log \frac{10}{a-x} = 1.2036 \quad \text{or } 1 - \log(a-x) = 1.2036$$

$$\text{or } \log(a-x) = -0.2036 = 1.7964$$

$$\text{or } (a-x) = \text{antilog } 1.7964 = 0.6258 \text{ gm}$$

92. (b) 0.08 mol l^{-1} to 0.01 mol l^{-1} involves 3 half-life. So the t is 30 minutes





93. (c) $t_{\frac{1}{2}}$ of II order reaction is inversely proportional to initial concentration of reactants.
94. (b) As $r = k(H_2O_2)$, it is a reaction of 1st order.
95. The given reaction is:
 $RCI + H_2O \rightarrow ROH + HCl$

Molecularity:

Molecularity is defined as the number of molecules colliding in the *elementary step*.

Here, one molecule of $RCIRCI$ and one molecule of H_2OH_2O are involved.

\Rightarrow **Molecularity = 2**

□ **Order of reaction:**

In kinetics, when a reaction is carried out in the presence of a **large excess of water**, the concentration of water remains practically constant.

So, the rate law effectively depends only on $[RCI]$

$$\text{Rate} = k[RCI]$$

• \Rightarrow **Order of reaction = 1**

: (b) **Molecularity is 2, order of reaction is 1**

96. For first order reaction,

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

$$k = \frac{0.693}{120} = 0.005775 \text{ min}^{-1}$$

Time for 90% decomposition:

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



Here, $[a] = 0.1[a]_0 \Rightarrow \frac{[a]_0}{[a]} = 10$

$$t = \{2.303 / 0.005775\} \log 10$$

$$t = \frac{2.303}{0.005775} \times 1 \approx 399 \text{ minutes}$$

