





# **CHEMISTRY**

# **BOOKS - DISHA CHEMISTRY (HINGLISH)**

# **CHEMICAL KINETICS**

# Chemistry

**1.** The dcecomposition of a substance follows first order kinatics. Its concentraction is reduced to  $\frac{1}{8}$ th of its initial value in 24 minutes. The rate constant of the decomposition process is

A. 
$$\frac{1}{4}min^{-1}$$
  
B.  $\frac{0.692}{24}mol^{-1}$   
C.  $\frac{2.303}{24}\log\left(\frac{1}{8}\right)min^{-1}$   
D.  $\frac{2.303}{24}\log\left(\frac{8}{1}min^{-1}\right)$ 

#### Answer: D



2. Consider the reaction,  $\circ A + B \rightarrow$  products. When concentration of B alone was doubled the half life did not change. When the concentraction of A alone was doubled, the rate increased by two times. The unit of rate constant for this reaction is

A.  $s^{-1}$ 

B.  $L \cdot Mol^{-1} \cdot s^{-1}$ 

C. no units

D.  $mol \cdot L^- \cdot s^-$ 

**Answer: B** 



**3.** A reaction involving two different reactants can never be

A. bimolecular reaction

B. second order reaction

C. first order reaction

D. unimolecular reaction

Answer: D



4. Select the rate law that corresponding to the shown for the following reaction  $: A + B \rightarrow C$ 





A. 
$$Rate = k[B]^3$$

$$\mathsf{B.} Rate = k[B]^4$$

C. 
$$Rate = k[A][B]^3$$

D. 
$$Rate = k[A]^2[B]^2$$



5. Consider the reaction,

 $N_2(q) + 3H_2(q) \rightarrow 2NH_3(q)$ equility relationship The between  $rac{d[NH_3]}{dt} \, ext{ and } \, - rac{d[H_2]}{dt} \, ext{is}$  $\mathsf{A.} + rac{d[NH_3]}{dt} = \ - \ rac{2}{3} rac{d[H_2]}{dt}$  ${\sf B.} + rac{d[NH_3]}{dt} = \ - \ rac{3}{2} rac{d[H_2]}{dt}$  $\mathsf{C}.\, \frac{d[NH_3]}{dt} = \frac{d[H_2]}{dt}$  $\mathsf{D.} + rac{d[NH_3]}{dt} = -rac{1}{3}rac{d[H_2]}{dt}$ 

6. Diazonium salt decomposes as

 $C_6H_5N_2^+Cl^ightarrow C_6H_5Cl+N_2$ 

At  $0^{\circ}$  C, the evolution of  $N_2$  becomes two faster

when the initial concentration of the salt is doubled. Therefore, it is

A. a first order reaction

B. a second order reaction

C. independent of the initial concentration of

the salt

D. a zero order reaction



7. 
$$CHCl_3 + Cl_2 \rightarrow \mathbb{C}l_4 + HCl$$

Rate law for above reactionl will be

$$\mathsf{Rate} = K [CHCl_3] rac{\left[Cl_2
ight]^1}{2}$$

On the basis of information provided which of the

following option will be correct?

A. Rate law for any chemical reaction can be

predicted accurately by looking at balanced

chemical equation.

B. Rate law for a chemical reaction has to

determine experimentally.

C. Either determine experimentally or obtained

from balanced reaction rate will be same.

D. None of the above is correct

Answer: B

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8. Which of the following statements is incorrect?

A. Activation energy for the forward reaction equals to activation energy for the reverse reaction.

- B. For a reversible reaction, an increase in temperature increase the reaction rate for both the forward and the backward reaction.
  C. The larger the initial reactant concentration for a second order reaction, the shorter is its half-life.
- D. When  $\Delta t$  is infinitesimally small, the averga rate equals the intantaneous rate.

### Answer: A



**9.** In a reaction  $A \rightarrow$  products, when start is made from  $8.0 \times 10^{-2}$ M of A, half-life is found to be 120minute. For the initial concentration  $4.0 \times 10^{-2}$ M, the half-life of the reaction becomes 240 minutes. The order of the reaction is :

A. zero

B. one

C. two

#### D. 0.5

## Answer: C

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**10.** During decomposition of an activated comples.

(i).energy is always releasing

(ii).energy is always absorbed

(iii).energy does not change

(iv).reactants may be formed

A. (i), (ii) and (ii)

B. (i) and (iv)

C. (ii) and (iii)

D. (ii), (iii) and (iv)

#### **Answer: B**



# 11. Which of the following statements is incorrect?

A. Energy is always released when activated

complex decomposes to form products.

B. Peak of the energy distribution curve

corresponds to the most probable potential

energy.

C. Peak of the energy distribution curve corresponds to the most probable kinetic energy.

D. When the temperature is raised maximum of energy curve moves to higher energy value and broadens out.

Answer: B



12. A catalyst increase rate of reaction by

A. decrease enthalpy.

B. decreasing internal energy.

C. decreasing activation energy.

D. increasing activation energy.

Answer: C



**13.** Consider a reaction  $aG + bH \rightarrow \text{products.}$ When concentration of both the reactants G and H is doubled, the rate increase by eight times. However, when concentration of G is doubled keeping the concentration of H fixed, the rate is doubled. The overall order of the reaction is

**A.** 0

**B**. 1

 $\mathsf{C.}\,2$ 

D. 3

Answer: D



**14.** For a first order reaction  $(A) \rightarrow$  products the concentration of A changes from 0.1M to 0.025M in 40 minutes.

The rate of reaction of when the concentration of A is 0.01M is:

A. 
$$1.73 imes 10^{-5}M\cdot min^{-1}$$

B.  $3.47 imes 10^{-4}M\cdot min^{-1}$ 

C.  $3.47 imes 10^{-5} M \cdot min^{-1}$ 

D.  $1.73 imes 10^{-4} M \cdot min^{-1}$ 

#### Answer: B



# 15. The given reaction

 $FeCl_3 + SnCl_2 
ightarrow 2FeCl_2 + SnCl_4$  is an

### example of

A. first order concentration

B. second order reaction

C. third order reaction

D. None of these

# Answer: C

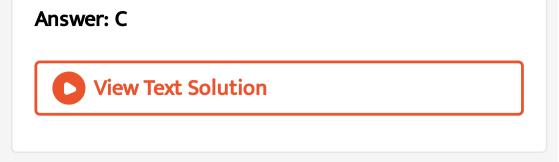


**16.** In a first order reaction,  $A+B o \;$  , If K is rate constant and initial concentration of the reactant A is 0.5M, then the half-life is

A. (log2)/(k)

B. (log2)/(ksqrt(0.5))

C. 
$$\frac{In2}{k}$$
  
D.  $\frac{0.693}{0.5k}$ 



- **17.** The integrated rate equations can be determined for
  - A. zero order reactions
  - B. first order reactions
  - C. second order reactions
  - D. Both (a) and (b)

Answer: D

18. The rate constant, the activation energy and the arrhenius paremeter of a chemical reaction at  $25^{\circ}$  are  $3.0 imes 10^{-4}s^-$ ,  $104.4kJ \cdot mol^{-1}$  and  $6.0 imes 10^{14}s^{-1}$  respectively. The value of the rate constant as  $T \to \infty$  is

A.  $2.0 imes10^{18}s^{-1}$ 

B.  $6.0 imes10^{14}s^{-1}$ 

C. Infinity

D.  $3.6 imes10^{30}s^{\,-1}$ 

### Answer: B



**19.** According to the adsorption theory of catalysis, the speed of the reaction increase because

A. in the process of adsorption, the activation

energy of the molecules becomes large

B. adsorption produces heat which increase the

speed of the reaction.

C. adsorption lowers the activation energy of

the reaction

D. the concentration of product molecules at

the active centres of the catalyst becomes

high due to adsorption.

Answer: C



20. Consider a general chemical change2A+3B
ightarrow products. The rate with respect to A is

- $r_1$  and that with respect to B is  $r_2$ . The rates
- $r_1$  and  $r_2$  are related as

A. 
$$3r_1=2r_2$$

B. 
$$r_1 = r_2$$

C. 
$$2r_1=3r_2$$

D. 
$$r_1^2=2r_2^2$$



**21.** In a first order reaction, the concentration of the reactant, decrease from 0.8M to 0.4M in 15minutes. The time taken for the concentration to changes from 0.1M to 0.025M is

A. 7.5 minutes

B. 15 minutes

C. 30 minutes

D. 60 minutes

Answer: C



**22.** Activation energy  $(E_a)$  and rate constant  $(k_1 \text{ and } k_2)$  of chemical reaction at two different temperatures  $(T_1 \text{ and } T_2)$  are related by :

A. 
$$In \frac{k_2}{k_1} = \frac{E_a}{R} \left[ \frac{1}{T_1} - \frac{1}{T_2} \right]$$
  
B.  $In \frac{k_2}{k_1} = \frac{E_a}{R} \left[ \frac{1}{T_2} - \frac{1}{T_1} \right]$   
C.  $In \frac{k_2}{k_1} = \frac{E_a}{R} \left[ \frac{1}{T_2} + \frac{1}{T_1} \right]$   
D.  $In \frac{k_2}{k_1} = \frac{E_a}{R} \left[ \frac{1}{T_1} + \frac{1}{T_2} \right]$ 

#### Answer: A

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**23.**  $\frac{t_1}{4}$  can be taken as the time taken for the concentration of a reactant to drop to  $\frac{3}{4}$  of its initial value. If the rate constant for a first order reaction is k, the  $\frac{t_1}{4}$  can be writen as

A. 
$$\frac{075}{k}$$
  
B.  $\frac{0.69}{k}$   
C.  $\frac{0.29}{k}$   
D.  $\frac{0.10}{k}$ 

#### Answer: C



**24.** the decomposition of ammonia on tungsten surface at 500K follows zero order kinetics. The Half-life period of this reaction is 45 minutes when the initial pressure is 4 bar. The half-life period (minutes) of the reaction when the initial pressure is 16 bar at the same temperature is

A. 120

**B.** 60

C.240

D. 180

#### Answer: D





**25.** In a 1st order reaction, reactant concentration C varies with times t as:

A. 
$$\frac{1}{C}$$
 increase linearly with t  
B. logC decrease linearly with t  
C. C decrease with  $\frac{1}{t}$ 

D. logC decrease with  $\frac{1}{t}$ 

### Answer: B



**26.** For a reactions  $A \rightarrow \text{Products}$ , a plot of  $\log t_{\frac{1}{2}}$ vesus  $\log a_o$  is shown in the figure. If the initial concentration A is represented by  $a_0$ , the order of the reaction is



A. one

B. zero

C. two

D. three

**Answer: B** 



27. The rate of a chemical reaction doulets for every  $10^{\circ}C$  rise of temperature. If the temperature is raised by  $50^{\circ}$  C, the rate of the reaction by about,

A. 10 times

B. 24 times

C. 32 times

D. 64times

Answer: C



**28.** For a first order reaction  $t_{0.75}$  is 1368 seconds, therefore, the specific rate constant in sec<sup>-1</sup> is

A.  $10^{-3}$ 

B.  $10^{-2}$ 

 $C. 10^{-9}$ 

D.  $10^{-5}\,$ 



**29.** The integrated rate equations is  $R = \log C_0 - \log C_r$ 

The straight line graph is obtained by plotting

A. timesVs $\log C_t$ 

B. 
$$\frac{1}{\text{time}} VsC_t$$

C. timeVs $C_t$ 

D. 
$$\frac{1}{\text{time}} Vs \frac{1}{C_t}$$



30. The energies of activation for forward and reverse reactions for  $A_2 + B_2 \leftrightarrow 2AB$  are  $180kJmol^{-1}$  and  $200kJmol^{-1}$  respectively. The presence of catalyst lowers the activation energy of both (forward and reverse) reactions by  $100kJmol^{-1}$ . The enthalpy change of the reaction  $(A_2 + B_2 
ightarrow 2AB)$  in the presence of a catalyst will be (in  $kJmol^{-1}$ )

A. 20

**B.** 300

**C**. 120

D. 280

## Answer: A



**31.** The half-life period of a order reaction is 15 minutes. The amount of substance left after one hour will be:

A. 
$$\frac{1}{4}$$
 of the original amount  
B.  $\frac{1}{8}$  of the original amount  
C.  $\frac{1}{16}$  of the original amount  
D.  $\frac{1}{32}$  of the original amount

# Answer: C



**32.** Reaction rate between two substance A and B is expressed as following,

 $\mathsf{rate} = k[A]^n[B]^m$ 

If the concentration of A is doubled and concentration of B is made half of initial concentration, the ratio of the new rate to the earlier rate will be,

A. m + n

B.n-m

$$\mathsf{C}.\,\frac{1}{2^{n+m}}$$

D. 
$$2^{n-m}$$

**Answer: D** 



**33.** The reaction of ozone with oxygen atoms in the presence of chlorine atoms can occur by two step process shown below,

$$egin{aligned} O_3(g)+Cl(g)& o O_2(g)+ClO(g)&...& ext{(i)}& ext{[}& ext{K}_i=5.2 imes10^9L\cdot mol^{-1}\cdot s^{-1} \end{aligned}$$

$$ClO(g) + O_2(g) \rightarrow O_2(g) + Cl(g)....$$
 (ii) [  
 $K_{ii} = 2.6 \times 10^{10} l \cdot mol^{-1} \cdot s^{-1}$ ]  
The closet rate constant for the overall reaction  
 $O_3(g) + O(g) \rightarrow 2O_2(g)$  is:  
A.  $1.4 \times 10^{20} l \cdot mol^{-1} \cdot s^{-1}$   
B.  $3.1 \times 10^{10} l \cdot mol^{-1} \cdot s^{-1}$   
C.  $5.2 \times 10^9 l \cdot mol^{-1} \cdot s^{-1}$   
D.  $2.6 \times 10^{10} l \cdot mol^{-1} \cdot s^{-1}$ 

### Answer: A



**34.** The temperature dependence of rate constant(k) of a chemical reaction is written in terms of Arrenius equation,  $k = A \cdot e^{-E_a}$  Activation energy  $(E_a)$  of the reaction can be calculated by plotting

A. 
$$kVs\frac{1}{lagT}$$
  
B.  $\log kVs\frac{1}{T}$   
C.  $\log kVs\frac{1}{\log T}$ 

D. 
$$kVsT$$

### Answer: B



35. For the reaction system,

 $2NO(g) + O_2(g) \rightarrow 2NO_2(g)$  volume is suddenly reduced to half its value by increasing the pressure on it. If the reaction is of first order with respect to  $O_2$  and second order with respect to No, the rate of reaction will

A. diminish to one-eighth of its initial value

B. increase to eighth times of its initial value

C. increase to fourth times of its initial value

D. diminish to one-fourth of its initial value

### Answer: B



**36.** In the reaction of formation of sulphur trioxide by contact process  $2SO_2 + O_2 \leftrightarrow 2SO_3$  the rate of reactionn was  $\frac{d[O_2]}{dt} = 2.5 \times 10^{-4} mol \cdot L^{-1} \cdot s^{-1}$ . The rate of reaction is terms of  $[SO_2]$  in  $mol \cdot L^{-1} \cdot s^{-1}$  will be

A. 
$$-1.25 imes10^{-4}$$

$$\mathsf{B.}-2.50\times10^{-4}$$

$$\mathsf{C.}-3.75 imes10^{-4}$$

 $\mathsf{D.}-5.00 imes10^{-4}$ 

#### **Answer: D**



# **37.** A reactant (A) froms two products,

$$A \stackrel{k_1}{\longrightarrow} B$$
, Activation Energy  $E_{a_1}$ 

$$A \stackrel{k_2}{\longrightarrow} B$$
, Activation Energy  $E_{a_2}$ 

If  $E_{a_1} = 2E_{a_2}$ , then  $k_1 \,$  and  $\, k_2$  are related as:

A. 
$$k_2=k_1e^{rac{E_{a_1}}{RT}}$$

B. 
$$k_2=k_1e^{rac{E_{a_2}}{RT}}$$

C. 
$$k_1 = Ak_2 e^{rac{E_{a_1}}{RT}}$$

D. 
$$k_1=2k_2e^{rac{E_{a_1}}{RT}}$$

#### Answer: C

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**38.** Consider an endothermic reaction  $X \to Y$  with the activation energies  $E_b$  and  $E_i$ , for the backward and forward reactions, respectively. In general A. there is no definite relation between

 $E_b$  and  $E_f$ 

B.  $E_b = E_f$ 

C.  $E_b > E_f$ 

D.  $E_b < E_f$ 

Answer: D

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39. Which of the following influences the rate of a

chemical reactions performed in soution?

## A. Temperature

B. Activation energy

C. Presence of a catalyst

D. All of the above influence the rate

### Answer: D



**40.** The rate of a reaction quadruples when the tamperature changes from 300 to 310k. The activation energy of this reaction is : (Assume activation energy and pre-exponential factor are

independent of temperature , In2=0.693,

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

A.  $107.2kJ \cdot mol^{-1}$ 

B.  $53.6kJ \cdot mol^{-1}$ 

C.  $26.8kJ \cdot mol^{-1}$ 

D.  $214.4kJ \cdot mol^{-1}$ 

**Answer: A** 



41. Consider the reaction :

 $Cl_2(aq)+H_2S(aq) o S(s)+2H^+(aq)+2Cl^-(aq)$ the rate equation for this reaction is rate= $k[Cl_2][H_2S]$ 

Which of these mechanism is/ are consistent with this rate equation? A.  $Cl_2 + H_2S \rightarrow H^+ + Cl^- + HS^-$  (Slow)  $Cl^+ + HS^- \rightarrow H^+ + Cl^- + S$  (fast) B.  $H_2S \leftrightarrow H^+ + HS^-$  (fast equilibirium)  $Cl_2 + HS^- \rightarrow 2Cl^- + H^+ + S$  (Slow)

A. B only

B. Both A and B

C. neither A nor B

D. A only

Answer: D



42. The slope in Arrhenius plot, is equal to,

A. 
$$-rac{E_a}{2.303R}$$
  
B.  $rac{E_a}{R}$   
C.  $rac{R}{2.303E_2}$ 

D. None of these



**43.** A schematic plot of  $InK_{eq}$  versus inverse of temperature for a reaction is shown below

The reaction must be

A. highly spontaneous at ordinaty temperature

B. one with nigligible enthalpy change

C. endothermic

D. exothermic

## Answer: D



44. Plots showing the variation of the rate constant(k) with temperature (T) are given below. The plotthat follows Arrhenius equations is









### Answer: A



**45.** The reaction  $2NO_2(g) \rightarrow 4NO_2(g) + O_2(g)$ follows first order kinetics. The pressure of a vessel containing only  $N_2O_5$  was found to increase from 50mm Hg to 87.5mm Hg in 30min. The pressure exerted by the gases after 60min will be (assume temperature remains constant):

A. 106.2mm Hg

 $\mathsf{B}.\,150mm\;\mathsf{Hg}$ 

 $\mathrm{C.}\,125mm~\mathrm{Hg}$ 

 $\mathsf{D}.\,116.5mm~\mathsf{Hg}$ 

### Answer: A

