PHYSICAL / INORGANIC CHEMISTRY



DPP No. 47

Total Marks: 30

Max. Time: 30 min.

Topic: Chemical Kinetics

	of Questions choice Objective ('	–1' negative marking) Q	.1 to Q.10	(3 marks 3 min.)	M.M., Min. [30, 30]	
1.	2NO + 2H ₂ \longrightarrow N ₂ + 2H ₂ O. The experimental rate law for above reaction is, Rate = k [NO] ² [H ₂]. When time is in minutes and the concentration is in moles/L, the units for k are					
	(A) $\frac{\text{moles}^3}{\text{L}^3 - \text{min}}$	(B) $\frac{\text{moles}}{L - \text{min}}$	(C) $\frac{\text{moles}^2}{\text{L}^2 - \text{min}}$	(D) $\frac{L^2}{\text{moles}^2}$		
2.	The differential rate law equation for the elementary reaction A + 2B $\xrightarrow{\kappa}$ 3C, is :					
	$(A) - \frac{d[A]}{dt} = -\frac{d[B]}{dt}$	$= \frac{d[C]}{dt} = k[A][B]^2$	(B) $-\frac{d[A]}{dt}$	$=-\frac{1}{2}\frac{d[B]}{dt}=\frac{1}{3}\frac{d[C]}{dt}=k$	[A] ² [B]	
	$(C) - \frac{d[A]}{dt} = -\frac{1}{2} \frac{d[A]}{dt}$	$\frac{[B]}{dt} = \frac{1}{3} \frac{d[C]}{dt} = k[A][B]^2$	(D) None of	these		
3.	For the reaction 2A \longrightarrow B + 3C; if $-\frac{d[A]}{dt} = k_1[A]^2$; $\frac{d[B]}{dt} = k_2[A]^2$; $\frac{d[C]}{dt} = k_3[A]^2$, the correct relation					
	between \mathbf{k}_1 , \mathbf{k}_2 , and \mathbf{k}_3 is :					
	(A) $k_1 = k_2 = k_3$	(B) $2k_1 = k_2 = 3 k_3$	(C) $4k_1 = k_2 =$	$= 3 k_3$ (D) $\frac{k_1}{2} = k_2$	$=\frac{k_3}{3}$	
4.	(A) Unit of rate of d	ing statement is incorrect? isappearance is Ms ⁻¹ stant k is depend on orde	(B) Unit of ra	(B) Unit of rate of reaction is Ms ⁻¹ (D) Unit of k for first order reaction is Ms ⁻¹		
5.	The rate expression for reaction A (g) + B(g) \rightarrow C(g) is rate = k[A] ^{1/2} [B] ² . What change in rate if initi concentration of A and B increase by factor 4 and 2 respectively?					
	(A) 4	(B) 6	(C) 8	(D) None of	these	
6.	Reaction A \rightarrow B follows second order kinetics. Doubling the concentration of A will increase the rate of formation of B by a factor of :					
	(A) 1/4	(B) 1/2	(C) 2	(D) 4		
7.	For the reaction $2NO_2 \longrightarrow N_2O_2 + O_2$, rate expression is as follows					
	$-\frac{d[NO_2]}{dt}$ =K $[NO_2]^n$, where K = 3 × 10 ⁻³ mol ⁻¹ L sec ⁻¹ . If the rate of formation of oxygen is 1.5 × 10 ⁻⁴ mol L ⁻¹					
	sec ⁻¹ , then the molar concentration of NO ₂ in mole L ⁻¹ is					
	(A) 1.5×10^{-4}	(B) 0.0151	(C) 0.214	(D) 0.316		
8.	Sodium (Na = 23) crystallizes in bcc arrangement with the interfacial separation between the atoms at the edge 53.6 pm. The density of sodium crystal is:					
	(A) 2.07 g/cc	(B) 2.46 g/cc	(C) 1.19 g/cc			
9.	TIAI(SO ₄) ₂ . xH ₂ O is bcc·with 'a' = 1.22 nm. If the density of the solid is 2.32 g/cc, then the value of x is (Given: N _A = 6 × 10 ²³ ; at. wt. : TI = 204, AI = 27, S = 32).					
	(A) 2	(B) 4	(C) 47	(D) 70		

An atomic solid crystallizes in a body centre cubic lattice and the inner surface of the atoms at the adjacent corner are separated by 60.3 pm. If the atomic weight of A is 48, then density of the solid, is nearly:

(C) 3.5 g/cc

(B) 5.07 g/cc



(D) 1.75 g/cc

(A) 2.7 g/cc

10.

Answer Kev

DPP No. # 47

1. (D) 2. (C)

(D)

(D)

(C)

(D)

6. (D) 7.

(D)

(C)

(C)

nts & Solutions

PHYSICAL / INORGANIC CHEMISTRY

DPP No. #47

$$\frac{\text{conc.}}{x} = k \text{ (conc.)}^3$$

$$\frac{\text{conc.}}{\text{time}} = k \text{ (conc.)}^3 \implies k = \frac{1}{\text{time(conc.)}^2}$$

$$k = \frac{1}{\frac{\text{moles}^2}{L^2}(\text{min})} = \frac{L^2}{\text{moles}^2 - \text{min}}$$

2. A + 2B
$$\xrightarrow{\kappa}$$
 3C (elementary reaction)

Rate =
$$-\frac{d[A]}{dt} = -\frac{1}{2}\frac{d[B]}{dt} = \frac{1}{3}\frac{d[C]}{dt} = k[A][B]^2$$

3. Rate =
$$-\frac{1}{2} \frac{d[A]}{dt} = \frac{d[B]}{dt} = \frac{1}{3} \frac{d[C]}{dt}$$

$$\frac{k_1}{2}$$
 [A]² = K_2 [A]² = $\frac{k_3}{3}$ [A]²

$$\Rightarrow \frac{k_1}{2} = k_2 = \frac{k_3}{3}$$

5. Rate =
$$k[A]^{1/2}[B]^2$$

$$r_1 = k[a]^{1/2} [b]^2$$
;

$$r_2 = k[4a]^{1/2} [2b]^2$$

$$\Rightarrow \frac{r_1}{r_2} = \frac{1}{2 \times 4} = \frac{1}{8}$$

$$\Rightarrow r_2 = 8 \times r_1$$

$$\Rightarrow$$
 $r_2 = 8 \times r_1$



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

$$\frac{\left(\frac{d[B]}{dt}\right)_1}{\left(\frac{d[B]}{dt}\right)_2} = \frac{k(a)^2}{k(2a)^2} = \frac{1}{4}$$

7. From the unit of K, it is evident that it is a second order reaction.

$$-\frac{1}{2} \, \frac{d[NO_2]}{dt} = \frac{d[O_2]}{dt} \qquad \Rightarrow \qquad \therefore \qquad -\frac{d[NO_2]}{dt} = 2 \times \frac{d[O_2]}{dt} = 2 \times 1.5 \times 10^{-4} = 3 \times 10^{-4} \\ 3 \times 10^{-4} = K \, [NO_2]^2 = 3 \times 10^{-3} \, [NO_2]^2 \qquad \Rightarrow \qquad \therefore \qquad [NO_2] = 0.316.$$

a - 2r = 53.6 pm8.

also
$$4r = \sqrt{3}a$$
 \Rightarrow $a - \frac{\sqrt{3}}{2}a = 53.6$
 \Rightarrow $a = \frac{53.6 \times 2}{2 - \sqrt{3}} = 400 \text{ pm}$
Density $(\rho) = \frac{2 \times 23}{6.023 \times 10^{23} \times 4^3 \times 10^{-24}} = 1.19 \text{ g/cc}$

9.
$$2.32 = \frac{2 \times M}{6 \times 10^{23} (1.22)^3 \times 10^{-21}}$$
$$\Rightarrow M = 1264 \Rightarrow x \approx 47.$$

10. Given a – 2r = 60.3 and for bcc, $4r = \sqrt{3}a$

$$\Rightarrow \qquad a - \frac{\sqrt{3}}{2} \ a = 60.3 \Rightarrow a = 450 \text{ pm}$$

$$\text{Density } (\rho) = \frac{2 \times 48}{6.023 \times 10^{23} \times (4.5)^3 \times 10^{-24}} = 1.75 \text{ g/cc.}$$

