

Date Planned ://	Daily Tutorial Sheet-2	Expected Duration : 90 Min
Actual Date of Attempt : / /	JEE Advanced (Archive)	Exact Duration :

**16.** For a gaseous reaction 2B  $\longrightarrow$  A, the equilibrium constant  $K_P$  is ..... than  $K_C$ .

(1997)

- 17. The degree of dissociation is 0.4 at 400K and 1.0 atm for the gaseous reaction  $PCl_5 \rightleftharpoons PCl_3 + Cl_2$ . Assuming ideal behaviour of all the gases, calculate the density of equilibrium mixture at 400 K and 1.0 atm (relative atomic mass of P = 31.0 and Cl = 35.5). (1998)
- **18.** For the reversible reaction,  $N_2(g) + 3H_2(g) \Longrightarrow 2NH_3(g)$

(2000)

At 500°C, the value of  $K_p$  is  $1.44\times 10^{-5}$  when partial pressure is measured in atmosphere. The corresponding value of  $K_c$  with concentration in mol/L is:

(A)  $\frac{1.44 \times 10^{-5}}{(0.082 \times 500)^{-2}}$ 

**(B)**  $\frac{1.44 \times 10^{-5}}{(8.314 \times 773)^{-2}}$ 

(c)  $\frac{1.44 \times 10^{-5}}{(0.082 \times 773)^2}$ 

- (D)  $\frac{1.44 \times 10^{-5}}{(0.082 \times 773)^{-2}}$
- 19. At constant temperature, the equilibrium constant ( $K_p$ ) for the decomposition reaction,  $N_2O_4 \Longrightarrow 2NO_2$ , is expressed by  $K_p = \frac{4x^2p}{(1-x)^2}$ , where p = pressure, x = extent of decomposition. Which one of the following

statement is true?

(2001)

- (A)  $K_p$  increases with increases of p
- **(B)**  $K_n$  increases with increases of x
- (C)  $K_p$  increases with decreases of x
- (D)  $K_p$  remains constant with change in p and x
- **20.** Consider the following equilibrium in a closed container  $N_2O_4(g) \rightleftharpoons 2NO_2(g)$

At a fixed temperature, the volume of the reaction container is halved. For this change, which of the following statements hold true regarding the equilibrium constant  $(K_p)$  and degree of dissociation  $(\alpha)$ ?

- (A) Neither  $K_p$  nor  $\alpha$  changes
- **B)** Both  $K_p$  and  $\alpha$  changes
- (2002)

- (C)  $K_p$  changes but  $\alpha$  does not change
- (D)  $K_p$  does not change but  $\alpha$  changes
- 21. In the following equilibrium  $N_2O_4(g) \Longrightarrow 2NO_2(g)$  when 5 moles of each are taken, the temperature is kept at 298 K the total pressure was found to be 20 bar. Given that (2004)

$$\Delta G_f^{\circ}(N_2O_4) = 100 \text{kJ}, \ \Delta G_f^{\circ}(NO_2) = 50 \text{ kJ}$$

 $\odot$ 

- (i) Find  $\Delta G^{\circ}$  of the reaction.
- (ii) The direction of the reaction in which the equilibrium shifts.
- **22.**  $Ag^+ + NH_3 \rightleftharpoons [Ag(NH_3)]^+; K_1 = 3.5 \times 10^{-3}$

(2006)

 $[Ag(NH_3)]^+ + NH_3 \Longrightarrow [Ag(NH_3)_2]^+; K_2 = 1.7 \times 10^{-3}$ 



Then the formation constant of  $[Ag(NH_3)_2]^+$  is:

- (A)  $5.95 \times 10^{-6}$
- **(B)**  $5.95 \times 10^6$
- (C)  $5.95 \times 10^{-9}$
- **(D)** None of these



\*23. The thermal dissociation of equilibrium of CaCO<sub>3</sub>(s) is studied under different conditions. (2013)

$$CaCO_3(s) \Longrightarrow CaO(s) + CO_2(g)$$

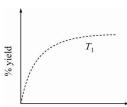
For this equilibrium, the correct statement(s) is:

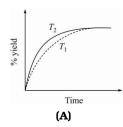
- (A)  $\Delta H$  is dependent on T
- **(B)** K is independent of the initial amount of CaCO3
- (C) K is dependent on the pressure of CO<sub>2</sub> at a given T
- **(D)**  $\Delta H$  is independent of the catalyst, if any
- 24. The %yield of ammonia as a function of time in the reaction

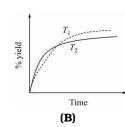
 $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g), \Delta H < 0 \text{ at } (P, T_1) \text{ is given below :}$ 

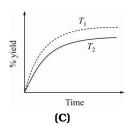
If this reaction is conducted at  $(P, T_2)$ , with  $T_2 > T_1$ , the %yield of (2015)

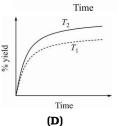
ammonia as a function of time represented by:











- \*25. The gas phase reaction  $2NO_2(g) \longrightarrow N_2O_4(g)$  is an exothermic reaction. The decomposition of  $N_2O_4$ , in equilibrium mixture of  $NO_2(g)$  and  $N_2O_4(g)$ , can be increased by : (2016)
  - (A) lowering the temperature
  - (B) increasing the pressure
  - (C) addition of an inert gas at constant volume
  - **(D)** addition of an inert gas at constant pressure
- $\Delta_f G^{\circ}$  at 500 K for substance 'S' in liquid state and gaseous state are +100.7 kcal mol<sup>-1</sup> and 26. +103 kcal mol<sup>-1</sup>, respectively. Vapour pressure of liquid 'S' at 500 K is approximately equal to:
  - (A) 0.1 atm
- (B) 1 atm
- (C) 10 atm
- (D) 100 atm
- 27. At a certain temperature in a 5 L vessel, 2 moles of carbon monoxide and 3 moles of chlorine were allowed to reach equilibrium according to the reaction,  $CO + Cl_2 \Longrightarrow COCl_2$ (2018)

At equilibrium, if one mole of CO is present then equilibrium constant (K<sub>o</sub>) for the reaction is:

 $(\blacktriangleright)$ 

- (A)
- (B) 2.5
- (C)
- For the following reaction, the equilibrium constant  $K_c$  at 298 K is  $1.6 \times 10^{17}$ 28.
- (2019)

$$Fe^{2+}(aq) + S^{2-}(aq) \Longrightarrow FeS(s)$$

When equal volumes of 0.06 M  $Fe^{2+}$ (aq) and 0.2 M  $S^{2-}$ (aq) solutions are mixed, the equilibrium concentration of  $\text{Fe}^{2+}$ (aq) is found to be  $Y \times 10^{-17} \text{M}$ . The value of Y is\_\_\_\_\_\_