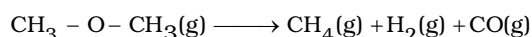


Date Planned : __ / __ / __	Daily Tutorial Sheet-2	Expected Duration : 90 Min
Actual Date of Attempt : __ / __ / __	JEE Advanced (Archive)	Exact Duration : _____

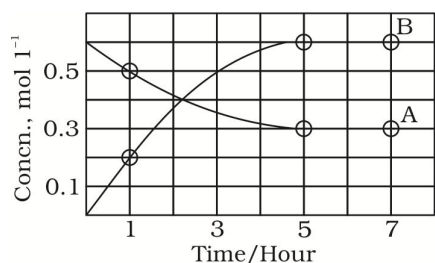
16. A first order reaction,  $A \rightarrow B$ , requires activation energy of  $70\text{kJ mol}^{-1}$ . When a 20% solution of A was kept at  $25^\circ\text{C}$  for 20 min, 25% decomposition took place. What will be percentage decomposition in the same time in a 30% solution maintained at  $40^\circ\text{C}$ ? Assume that activation energy remains constant in this range of temperature. ▶ (1993)

17. The gas phase decomposition of dimethyl ether follows first order kinetics : ▶ (1993)



The reaction is carried out in a constant volume container at  $500^\circ\text{C}$  and has a half-life of 14.5 min. Initially only dimethyl ether is present at a pressure of 0.40 atm. What is the total pressure of the system after 12 min? Assume ideal gas behaviour.

18. The progress of the reaction,  $A \rightleftharpoons nB$ , with time, is presented in figure given below. Determine. (1994)



- (i) The value of  $n$ ,  
(ii) The equilibrium constant  $K$ , and  
(iii) the initial rate of conversion of A.

19. For the reaction :  $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \longrightarrow 2\text{NH}_3(\text{g})$  (1994)

Under certain conditions of temperature and partial pressure of the reactants, the rate of formation of  $\text{NH}_3$  is  $0.001\text{ kg/h}^{-1}$ . The rate of conversion of  $\text{H}_2$  under the same condition is \_\_\_\_\_  $\text{kg/h}^{-1}$ .

20. From the following data for the reaction between A and B (1994)

[A], (mol/L)	[B], (mol/L)	Initial rate ( $\text{mol L}^{-1}\text{s}^{-1}$ ) at	
		300 K	320 K
$2.5 \times 10^{-4}$	$3.0 \times 10^{-5}$	$5.0 \times 10^{-4}$	$2.0 \times 10^{-3}$
$5.0 \times 10^{-4}$	$6.0 \times 10^{-5}$	$4.0 \times 10^{-3}$	--
$1.0 \times 10^{-3}$	$6.0 \times 10^{-5}$	$1.6 \times 10^{-2}$	--

Calculate:

- (i) the order of the reaction with respect to A and with respect to B  
(ii) the rate constant at 300 K (iii) the pre-exponential factor.
21. At  $380^\circ\text{C}$ , the half-life period for the first order decomposition of  $\text{H}_2\text{O}_2$  is 360 min. The energy of activation of the reaction is  $200\text{ kJ mol}^{-1}$ . Calculate the time required for 75% decomposition at  $450^\circ\text{C}$ . (1995)

22. In Arrhenius equation,  $k = A \exp^{(-E_a/RT)}$ . A may be termed as the rate constant at \_\_\_\_\_. (1997)
23. The rate constant for the first order decomposition of a certain reaction is described by the equation
- $$\log k(s^{-1}) = 14.34 - \frac{1.25 \times 10^4 K}{T} \quad (1997)$$
- (i) What is the energy of activation for the reaction?  
(ii) At what temperature will its half-life period be 256 min?
24. (i) The rate constant of a reaction is  $1.5 \times 10^7 s^{-1}$  at  $50^\circ C$  and  $4.5 \times 10^7 s^{-1}$  at  $100^\circ C$ . Evaluate the Arrhenius parameters A and  $E_a$ . (1998)  
(ii) For the reaction,  $N_2O_5(g) \longrightarrow 2NO_2(g) + \frac{1}{2}O_2(g)$   
Calculate the mole fraction  $N_2O_5(g)$  decomposed at a constant volume and temperature, if the initial pressure is 600 mm Hg and the pressure at any time is 960 mm Hg. Assume ideal gas behaviour.
- \*25. For the first order reaction, (1998)
- (A) the degree of dissociation is equal to  $(1 - e^{-kt})$   
(B) a plot of reciprocal concentration of the reactant vs time gives a straight line  
(C) the time taken for the completion of 75% reaction is thrice the  $\frac{1}{2}$  of the reaction  
(D) the pre-exponential factor in the Arrhenius equation has dimension of time,  $T^{-1}$  (▶)
- \*26. The following statement(s) is (are) correct: (1999)
- (A) A plot of  $\log k_p$  versus  $1/T$  is linear  
(B) A plot of  $\log [X]$  versus time is linear for a first-order reaction,  $X \rightarrow P$   
(C) A plot of P versus  $1/T$  is linear at constant volume  
(D) A plot of P versus  $1/V$  is linear at constant (▶)
27. The rate constant for an isomerisation reaction,  $A \longrightarrow B$  is  $3.42 \times 10^{-3} M / \text{min}$ . If the initial concentration of A is 1 M, calculate the rate of the reaction after 1 h. (1999)
28. A hydrogenation reaction is carried out at 500 K. If the same reaction is carried out in the presence of a catalyst at the same rate, the temperature required is 400 K. Calculate the activation energy of the reaction if the catalyst lowers the activation barrier by  $20 \text{ kJ mol}^{-1}$ . (2000)
29. The rate constant for the reaction,  $2N_2O_5 \longrightarrow 4NO_2 + O_2$  is  $1.3 \times 10^{-5} s^{-1}$ . If the rate is  $3.90 \times 10^{-5} \text{ mol L}^{-1} s^{-1}$ , then the concentration of  $N_2O_5$  (in  $\text{mol L}^{-1}$ ) is: (2000)
- (A) 1.4 (B) 1.2 (C) 0.04 (D) 3 (▶)
30. If 'I' is the intensity of absorbed light and 'C' is the concentration of AB for the photochemical process,  $AB + h\nu \rightarrow AB$ , the rate of formation of AB is directly proportional to: (2001)
- (A) C (B) I (C)  $I^2$  (D) C.I