

St. Xavier's College

CEMG

SYLLABUS

(RESTRUCTURED)

PHYSICAL CHEMISTRY

Module I:

(10 L)

Chemical Kinetics

- a. Definition of rate of a chemical reaction in terms of degree of advancement.
- b. Introduction of some methods to study a chemical reaction to find its rate.
- c. Definition of order of a reaction.
- d. Methods for determination of order of a reaction.
- e. Dimension of rate and rate constants for reactions with different orders.
- f. Integrated rate law for (i) zeroth, (ii) first, (iii) second and (iv) n'th order cases.
- g. Nature of variation in concentration with time for reactions with different orders.
- h. Use of the integrated rate laws in determining order of a chemical reaction.
- i. Half life period, its expression for reactions with different orders.
- j. Usage of half life period in determining order of a chemical reaction.
- k. Expression of rate constant for binary reaction involving gases.
- l. Variation of rate constant with temperature: Arrhenius equation.

Module II:

(10 L)

Electrical conductance

- a. Metallic and electrolytic conduction.
- b. Ions as charge carrier.
- c. Strong and weak electrolytes.
- d. Solvation of ions in solution: effect of charge to radius ratio.
- e. Interaction among ions in solution and its variation with concentration (qualitative idea of asymmetric and electrophoretic effect).
- f. Definition of specific and equivalent conductance.
- g. Variation in conductance, specific conductance and equivalent conductance with concentration.
- h. Equivalent conductance at infinite dilution: Kohlrausch law of independent migration.
- i. Interrelation among strength of current, specific conductance, ionic mobility and ion conductance.
- j. Definition of transport number and expression.

Module III:

(20 L)

Chemical Thermodynamics

- a. Validity of thermodynamic system.
- b. System, surrounding and boundary
- c. Classification of systems
- d. Mode of interaction between system and surrounding.
- e. Reversible and irreversible path of transformation of the system.
- f. Work as the path function.
- g. Calculation of work in hydrostatic system (IUPAC convention).
- h. Work as the area under the PV indicator diagram.
- i. State functions.
- j. Internal energy and its change.
- k. Mathematical statement of first law of thermodynamics.
- l. Calculation of heat transferred.
- m. Enthalpy (state function and calculation of its change)
- n. Heat capacities (at constant pressure, C_p and at constant volume, C_v).
- o. Difference between the heat capacities for ideal and van der Waals gas.
- p. Relation between the state parameters in case of reversible adiabatic processes involving ideal gases.
- q. Need for the second law.
- r. Definition of entropy.

- s. Entropy change of the universe to predict the direction of spontaneous change.
- t. Clausius inequality.
- u. Gibbs and Helmholtz free energies and their changes with interpretation.
- v. Open systems: partial molar quantities with special reference to chemical potential.
- w. Variation in chemical potential with pressure and temperature.
- x. Different form of chemical potential involving standard state in different scales.
- y. Four fundamental equations for change in (i) internal energy, (ii) enthalpy, (iii) Gibbs free energy and (iv) Helmholtz free energy.
- z. Definition of one mole of a reaction.
- aa. Expression of change in free energy per mole of the reaction.
- bb. van't Hoff isotherm
- cc. Equilibrium constant at constant pressure and at constant volume.
- dd. Variation in equilibrium constants with temperature and pressure.
- ee. Interrelation between the equilibrium constants.
- ff. Qualitative idea of Le Chatelier's principle.