PHARMACEUTICS SYLLABUS

Chapter	Topic
1.	 History of the profession of Pharmacy in India in relation to Pharmacy education, industry, pharmacy practice, and various professional associations.
	 Pharmacy as a career Pharmacopoeia: Introduction to IP, BP, USP, NF and Extra Pharmacopoeia. Salient
	features of Indian Pharmacopoeia.
2.	Packaging materials: Types, selection criteria, advantages and disadvantages of glass,
	plastic, metal, rubber as packaging materials.
3.	3.1 Pharmaceutical aids: Organoleptic (Colouring, flavouring and sweetening) agents. 3.2 Preservatives: Definition, types with examples and uses.
4.	 Unit operations: Definition, objectives/applications, principles, construction, and workings of: 4.1 Size reduction: hammer mill and ball mill.
	4.2 Size separation: Classification of powders according to IP, Cyclone separator, Sieves and standards of sieves.
	4.3 Mixing: Double cone blender, Turbine mixer, Triple roller mill and Silverson mixer homogenizer.
	 4.4 Filtration: Theory of filtration, membrane filter and sintered glass filter. 4.5 Drying: working of fluidized bed dryer and process of freeze drying. 4.6 Extraction: Definition, Classification, method, and applications.
5.	 5.1 Tablets: coated and uncoated, various modified tablets (sustained release, extended release, fast dissolving, multilayered, etc.) 5.2 Capsules: hard and soft gelatin capsules.
	5.3 Liquid oral preparations: solution, syrup, elixir, emulsion, suspension, dry powder for reconstitution.
	5.4 Topical preparations: ointments, creams, pastes, gels, liniments and lotions, suppositories, and pessaries, Nasal preparations, Ear preparations.
	5.5 Powders and granules: Insufflations, dusting powders, effervescent powders, and effervescent granules.
	5.6 Sterile formulations: Injectables, eye drops and eye ointments.5.7 Immunological products: Sera, vaccines, toxoids and their manufacturing methods.
6.	6.1 Basic structure, layout, sections, and activities of pharmaceutical manufacturing plants 6.2 Quality control and quality assurance: Definition and concepts of quality control
	and quality assurance, current good manufacturing practice (cGMP), Introduction to the concept of calibration and validation.
7.	Novel drug delivery systems: Introduction, Classification with examples, advantages, and challenges.