Course Type	Course Code	Name of Course	L	Т	Р	Credit
DC	NCYC502	Formulation & Drug Delivery Technology	3	1	0	4

Course Objective				
• This course will introduce the students to the principles of drug formulation and delivery.				
Learning Outcomes				
The students should be able to:				
. Understanding here the despect forms can be tailand to the needs of the nations and discover to be				

- Understanding how the dosage form can be tailored to the needs of the patient and disease to be treated.
- Learn about the physicochemical and physiological principles that are forming the basis of a rational development of modern dosage forms.

Unit No.	Topics to be Covered	Lecture Hours	Learning Outcome
1	Preformulation studies: Preformulation studies of drug substances. Role of pre-formulation in drug discovery: Material properties in lead selection, high throughput pre-formulation studies. Preformulation as a support for formulation development, identification of challenges during formulation development.	7L+2T	At the end of the course students will be able to learn the Preformulation idea of drug substances, proteins and peptides.
2	Influence of Drug Properties and Routes of Drug Administration on the design of sustained and controlled release systems: Rationale for controlled drug delivery, physiochemical properties and biological factors influencing the design and performance of sustained/controlled release products. Preparation method of tablets, capsules, emulsions and suspensions. Overview of different carrier systems for drug delivery. Formulations using nano systems like Lipid nanoparticles; Polymeric nanoparticles; Polymeric micelles; Nanofibers; Dendrimers; Nanogels and biosilica; Quantum dots; Nanotubes; Magnetic and metallic nanoparticles.	10L+4T	Learn the criteria for selection of drugs and polymers for the development of Novel drug delivery systems.
3	Biopharmaceutic and pharmacokinetic aspects of PO CRDDS: Strategies and design, factor effecting controlled release drug delivery systems, Computation of desired release rate and dose for CRDDS, Pharmacokinetic design for DDS; in-vitro/in-vivo considerations, Intermittent zero order and first order release.	10L+4T	Explain the formulation and evaluation of Novel drug delivery systems.

	permeation, development and evaluation of transdermal devices, Case studies. Drug targeting: Different levels of targeting-first order, second order and third order targeting, active and passive targeting, prodrug based drug targeting, brain targeting, tumor targeting.	9L+2T	
	permeation, development and evaluation of transdermal devices, Case studies. Drug targeting: Different levels of targeting first order second order	9L+2T	
5	Principles of skin permeation, factors affecting percutaneous absorption of drugs, sorption promoters, absorption enhancement by energy input - iontophoresis, sonophoresis and electroporation, pharmacokinetics of skin		approaches for the development of targeted drug Delivery systems.
4	Peroral controlled-release delivery: Design and fabrication of oral systems, dissolution-controlled release, diffusion- controlled release, diffusion and dissolution-controlled release, ion- exchange resins, pH-independent formulations, osmotically controlled release, altered density formulations, Case studies.	6 L+2T	Explain the principles and technology used in the design of sustained release and controlled release drug delivery systems.

Text Books:

1) Drug Delivery Systems, KK Jain, Humana Press, 1st Ed. (2008).

Reference Books:

- Performulation in Solid Dosage Form Development, Edited by Moji ChristianahAdeyeye and Harry G. Brittain, CRC Press, 1st Ed. (2008).
- 2. Handbook of Performulation: Chemical, Biological and Botanical Drugs, Edited by Sarfaraz K. Niazi, CRC Press, 2nd Ed. (2019).
- 3. Handbook of Pharmaceutical Excipients, Rowe, R.C., P. J. Sheskey, Pharmaceutical Press, 6th Ed. (2009).
- 4. Pharmaceutical Excipients: Characterization by IR, Raman, NMR Spectroscopy, M. Dekker. Bugay, D.E. and W.P. Findlay, CRC Press, 6th Ed. (1999).