## **CONTENT of COURSE**

Modules	Topics	Content	Learning Objective	Learning Outcome	Hrs. Assigned
Module 1	Awareness and Expertise on ISO 17025; Safety of Environment; GMP/GLP	Introduction to the principles of ISO/IEC 17025 Calibration, traceability and uncertainty of measurement Monitoring, witnessing and the human factor in assessments Safety of Environment; GMP/GLP	Management System, SOP guidelines, Preparation, & Change control	SOP,GMP and GLP	Lecture/ Seminar 14 hrs Practical/Project 20 hrs
Module 2	HPLC	Introduction to the principles of HPLC Instrumentation, working, applications & limitations Hands on practice on the instruments of HPLC	Sample processing Sample analysis and documentation	Quantitative and qualitative analysis of Drugs	Lecture/ Seminar 40 hrs Practical/Project 180 hrs
Module 3	Ultraviolet-Visible Spectroscopy; Polarimetry; HPTLC and Quantification of Natural Products	Introduction to the theory of Ultraviolet-Visible Spectroscopy & factors affecting the spectra. Theory and principle of HPTLC and Polarimetry Instrumentation, working, applications & limitations Hands on practice on Ultraviolet- Visible Spectrophotometer, HPTLC and Polarimeter	Sample preparation, Visualisation, Sample analysis, Data interpretation	Quantitative and qualitative analysis of Drugs, Optical rotation	Lecture/ Seminar 20 hrs Practical/Project 40 hrs
Module 4	Infrared Spectroscopy	Introduction to the theory of IR Spectroscopy & factors affecting the spectra.  Instrumentation, working, applications & limitations Introduction to Attenuated Total Reflectance Hands on practice on IR Spectrophotometer	Identification of Drugs	Determination of functional groups in molecules	Lecture/ Seminar 02 hrs Practical/Project 12 hrs
Module 5	Gas Chromatography	Introduction to the theory & principle of Gas Chromatography Instrumentation, working, applications & limitations Introduction to branches of Gas Chromatography Hands on practice on the instrument of Gas Chromatography	Sample processing Sample analysis and documentation	Quantitative and qualitative analysis of Drugs	Lecture/ Seminar 10 hrs Practical/Project 30 hrs

Module 6	Dissolution & Disintegration	Introduction to the theory and Principles of Dissolution & Disintegrator Instrumentation, working, applications & limitations Hands on practice on the Dissolution & Disintegrator apparatus	Developing and evaluating an IVIVC of solid dosage form	In vitro drug disintegration and drug release information	Lecture/ Seminar 04 hrs Practical/Project 50 hrs
Module 7	Karl- Fischer Titrator	Introduction to the theory & principle of KF Titration Instrumentation, working, applications & limitations Hands on practice on the Karl-Fischer Titrator	Understanding the importance of KF and sample handling techniques	Detection of water content	Lecture/ Seminar 02 hrs Practical/Project 12 hrs
Module 8	Titrimetric Analysis	Introduction to the theory & principle of various Titrations Instrumentation, working and applications Hands on practice on the related instruments	Types of solutions, pH, Buffers, Various methods of end- point determination	Qualitative Analysis Assay of Products	Lecture/ Seminar 08 hrs Practical/Project 30 hrs
Module 9	Chemical Analysis	Introduction to theory and principles of various methods of chemical analysis	understanding of various methods of chemical detection	Preparation of Solutions, Heavy metal Analysis	Lecture/ Seminar 08 hrs Practical/Project 48 hrs
Module 10	Microbiological Analysis of Drugs and Pharmaceuticals	Introduction to principles of various methods of Microbiological Analysis of Drugs and Pharmaceuticals	Microbiological Assay procedures	Analysis of Antibiotics, Microbiological limits, PET, BET & Sterility	Lecture/ Seminar 24 hrs Practical/Project 60 hrs
Module 11	Statistical Analysis	Introduction to various statistical methods for the evaluation of laboratory data	to design data, analyze data appropriately and interpret and draw conclusions	Accurate and consistent result	Lecture/ Seminar 12 hrs Practical/Project 10 hrs