GUJARAT TECHNOLOGICAL UNIVERSITY

M.Pharm PHARMACEUTICAL CHEMISTRY SEMESTER: I

Subject Name: REGULATORY AFFAIRS PRACTICAL - I

Subject Code: MRA105P

List of Practicals:

- 1. Case studies (4 Nos.) of each of Good Pharmaceutical Practices.
- 2. Documentation for in process and finished products Quality control tests for Solid, liquid, Semisolid and Sterile preparations.
- 3. Preparation of SOPs, Analytical reports (Stability and validation)
- 4. Protocol preparation for documentation of various types of records (BMR, MFR, DR)
- 5. Labeling comparison between brand & generics.
- 6. Preparation of clinical trial protocol for registering trial in India
- 7. Registration for conducting BA/BE studies in India
- 8. Import of drugs for research and developmental activities
- 9. Preparation of regulatory dossier as per Indian CTD format and submission in SUGAM
- 10. Registering for different Intellectual Property Rights in India
- 11. GMP Audit Requirements as per CDSCO
- 12. Preparation and documentation for Indian Patent application.
- 13. Preparation of checklist for registration of IND as per ICH CTD format.
- 14. Preparation of checklist for registration of NDA as per ICH CTD format.
- 15. Preparation of checklist for registration of ANDA as per ICH CTD format.
- 16. Case studies on response with scientific rationale to USFDA Warning Letter
- 17. Preparation of submission checklist of IMPD for EU submission.
- 18. Comparison study of marketing authorization procedures in EU.
- 19. Comparative study of DMF system in US, EU and Japan
- 20. Preparation of regulatory submission using eCTD software
- 21. Preparation of Clinical Trial Application (CTA) for US submission
- 22. Preparation of Clinical Trial Application (CTA) for EU submission
- 23. Comparison of Clinical Trial Application requirements of US, EU and Japan of a dosage form.
- 24. Regulatory requirements checklist for conducting clinical trials in India.
- 25. Regulatory requirements checklist for conducting clinical trials in Europe.
- 26. Regulatory requirements checklist for conducting clinical trials in USA