DEPARTMENT OF QUALITY ASSURANCE

and

DEPARTMENT OF DRUG REGULATORY AFFAIRS

Department of Quality Assurance and Drug Regulatory Affairs are committed to incorporation of quality in every aspect of pharmaceutical development and manufacturing.

The faculty has extensive industrial experience in various aspects of drug development and manufacturing.

Regulatory Services

1. Regulatory submissions – dossier preparations
2. Regulatory feasibility evaluation for various pharmaceutical products for different countries

IPR Services

1. Preliminary Patentability searches
2. Preliminary Infringement analysis

QA Services

1. Preparation of procedural documentation like SOPs
2. Preparation of site master files
3. Support in various facility audits WHO, PICS, GMP
4. Establishment of quality management systems
5. Lean manufacturing
6. Pharmaceutical plant designing (detailed engineering not included)