Mrs. Anagha Maharao

Qualification – M.Sc. Organic Chemistry, University of Mumbai

Professional experience - 30 years in QC, QA and RA of API and formulation companies

Present Profile

- Managing Director & Principal trainer at Institute of Pharmaceutical Management for QC, QA and RA courses since 2007
- GMP and RA Consultant to renowned pharmaceutical organizations for 10 years

Accomplishments

- Audited US, Indian and Chinese organizations to evaluate Quality System as per GMP guidelines
- Audited API, intermediate, starting material vendors and warehouse facilities
- Assisted pharma companies for facing regulatory inspections (EDQM, USFDA, WHO Geneva)
- Given guidance on DMF submissions to USFDA, EDQM, Health Canada, WHO etc
- Trained over 4000 pharma professionals on regulatory and GMP aspects
- Conducted seminars on 'Current regulatory requirements of DMF'
- Conducted 10 days regulatory affairs course at Astra-Zeneca, Bangalore
- Conducted training sessions on GMP, impurities, genotoxicity, PQR, change control, stability studies, good documentation, ICH Q11 etc in various organizations
- Felicitated on the eve of National conference on 'Women Entrepreneur of 21st Century: Trends and developments organized by University of Mumbai on 20th-21st November'09
- Published Indian patent (2006-06-16) on "Improved process for Zopiclone synthesis".
- Presented paper on "Studies on the application of hydrotropes as solvent media for some organic reactions" at ICCE 2005, Indore.
- Attended conventions like CPhI at Frankfurt, Paris & Spain and Informex at USA

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