CHINTAN RAVAL

Assistant Manager

Objective:

Using my knowledge and experience of pharmaceutical industry with my interest towards various aspects of analytical laboratory, contribute my best potential as a professional and as a human being to the organization to achieve a goal.

Work Experience:

- ***AMI LIFESCIENCE (P) LTD., Baroda, Gujarat.***

  Assistant Manager - ADL

  Jun - 2020 to till date.

  - Review of analyst qualification raw data, Qualification report.
  - Review of In-House Reference Standard / Working Standard protocol, report, raw data and COA.
  - Review of Intermediate material, API and Impurity qualification report and COA.
  - Preparation of URS and communicating to other laboratories for API testing.
  - Compilation and evaluation of analytical development data.
  - Ensure the good laboratory practices in laboratory.

- ***KALINTIES HEALTHCARE (P) LTD., Baroda, Gujarat.***

  Sr. Executive - ADL/QC Department

  Oct - 2019 to April - 2020.

  - Reviewed of analyst qualification raw data, Qualification report cleaning assessment raw data, investigational studies raw data and reports.
  - Reviewed of In-House Reference Standard / Working Standard protocol, report, raw data and COA.
  - Investigation of OOS, OOT, OOAL and Aberrant results.
  - Prepared and reviewed of change control.
  - Review of API and Intermediate stability studies.
  - Ensure the Instruments / Equipment is calibrated and maintain schedule for calibration.

CONTENT ME

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- chintanraval2111983@gmail.com
- O-18 / Komal Enclave,
  Opp. P.P.C.C. Ground,
  Near ShantivanChar Rasta,
  Paldi, Ahmedabad, Gujarat-380007

EDUCATIONS

- Master of Science-Chemistry
  Gujarat University/Second Class / 2006
- Bachelor of Science-Chemistry
  Gujarat University / First Class / 2003

- Resources and Team management.
- Compilation and evaluation of analytical development data.

➢ ***AMNEAL PHARMACEUTICALS PVT. LTD., Ahmedabad, Gujarat.***

Assistant Manager - ADL

April - 2014 to Sept – 2019

- Review of development and validation protocol, report and data.
- Troubleshooting of analytical problems occurs at various tests like Dissolution, BU, CU, Assay, RC, Saturated Solubility, BCS solubility, % Water content and Excipients compatibility study.
- Prepared partial validation report of dissolution test.
- Developed and Optimized of the dissolution parameters, discrimination media and bio-relevant media.
- Prepared Data compilation sheets and trend analysis sheet for stability samples.
- Reviewed of electronic records and audit trails in respective instruments.
- Review of analytical raw data, CMC studies raw data, characterization analytical raw data, protocol, reports, and Certificate of Analysis,
- Maintain TRFs, sample entry- destruction registers.

➢ ***CADILA PHARMACEUTICALS LTD., Dholka, Gujarat.***

Research Associate - ADL

Jan - 2012 to March – 2014

- Work plaining and distribution of TRF based development samples and Stability Samples for analysis.
- Reviewed of STP, ATR and Specification.
- Calibration and troubleshooting of instrument / equipment, like HPLC, Dissolution, K.F titrator, UV-spectrophotometer, FTIR, Polari meter, Balance, pH Meter etc....
- Developed of analytical method like assay, related substance, dissolution

➢ *TECHNICAL SKILLS*

- Analytical method development report preparation,
- Validation protocol/report preparation,
- Analytical method transfer,
- SOP preparation,
- Man power and resource management,
- Incident, Deviation, OOS, OOAT and Change control management,
- Reference Standard / Working Standard management,
- GDP/GLP review,
- Qualification, Calibration and Maintenance of Instrument / Equipment.

➢ *COMPUTER SKILLS*

- MS-Word,
- MS-Excel,
- MS-Power Point,
- Chromeleon,
- Labsolution
- Empower
- ERP Software

➢ *LANGUAGES KNOWN*

- English
- Gujarati
- Hindi

- ***DISHMAN PHARMACEUTICALS AND CHEMICALS LTD., Bavla, Gujarat.***

  Officer - QC Department

  Sept - 2009 to Dec – 2011

  - Analysis of API and intermediate.
  - To initiate and analyze the stability studies as per ICH guidelines
  - Calibration and troubleshooting of instrument / equipment
  - To perform working standard preparation and maintain in respective storage condition.

- ***MARCK BIOSCIENCE LTD, Kheda. Gujarat***

  Officer - QC Department

  May - 2007 to Aug – 2009

  - Analyzed the stability studies as per ICH guidelines.
  - To perform wet analysis for Pharmacopoeial and non-Pharmacopoeial products.
  - To perform the calibration of instruments / equipment as per laboratory schedule.
  - To perform working standard preparation and maintain in respective storage condition.

- ***NIRLIFE HEALTHCARE LTD (A Div. of Nirma Ltd.) Sachana, Gujarat***

  Chemist - QA/QC Department

  Sept - 2006 to April - 2007

  - Analyzed the stability studies as per ICH guidelines.
  - To perform wet analysis for Pharmacopoeial and non-Pharmacopoeial products.
  - To perform the calibration of instruments / equipment as per laboratory schedule.
  - To perform working standard preparation and maintain in respective storage condition.

➢ *PERSONAL DETAILS*

- Date of Birth : 21.01.1983
- Sex : Male
- Marital Status : Married
- Nationality : Indian