

SCIENTIST  
Professional Summary

• Scientist with 15 years of analytical science experience within the pharmaceutical industry.

Education

Bachelor of Science , Life Sciences - Chemistry 1999 New York Institute of Technology ¼ City , State , USA

Experience

Scientist May 2011 to Current

Planet Pharma ¼ Clayton , NC

• Perform analytical method development and validation testing.

• Perform various routine analytical testing assignments.

• Set up and initiate stability studies.

• Author analytical procedures, method development reports, validation protocols/reports, technical reports, and laboratory equipment procedures.

• Demonstration/familiarization of analytical methods to fellow analysts for training purposes.

• Complete all assigned training to maintain qualification to perform tasks.

• Maintain and coordinate calibration and preventive maintenance schedule for all laboratory instrumentation.

• Participated in global SOP harmonization groups.

Associate Scientist Feb 2006 to May 2011

Planet Pharma ¼ Columbia , MD

• Performed routine testing using a variety of analytical techniques.

• Supported method development and validation for multiple projects and dosage forms.

• Perform feasibility experiments and compiled data for project review meeting.

• Prepared data tables and authored validation sections for regulatory filings.

• Initialized in-house stability studies.

• Participated in cross-functional analytical working group meetings.

Analytical Chemist Nov 2001 to Feb 2006

Entegris, Inc. ¼ Bloomington , MN

• Perform testing/methods validation etc. on all instrumentation (UV/Vis, AA, FTIR, Dissolution apparatus, HPLC, GC).

• In-Process Dissolution testing of products.

• Performed product specific and instrument training for analysts.

• Became a versatile chromatographer while performing release and stability testing on numerous products within a group (Tests included Content Uniformity, Assay among others).

• Wrote/Revised various SOP's, IOP's, CP's, and GTP's.

• Troubleshoot Analytical Instruments such as GC/HPLC.

• Assist in Analytical Methods development via Method Transfer to Quality Labs.

• Review analytical paperwork, maintaining accordance to current GMP guidelines.

• Performed instrument qualifications along with CSV Part 11 Remediation group.

• Member of Millennium Paperless Reporting group reducing the bulk of paperwork within batch records.

Organic Preparations Supervisor Mar 2000 to Apr 2001

Hampton Clarke/Veritech Laboratories ¼ City , STATE

• Supervised a group of five technicians in the preparation of samples for Gas Chromatography analysis by numerous EPA methods.

• Assisted in the preparations following set SOP's.

• Daily delegation of responsibilities from a generated back-log.

• Daily logging out of completed samples after preparation.

• Performed necessary troubleshooting and mechanical repairs within lab.

Organic Preparations Technician Sep 1999 to Mar 2000  
Severn-Trent Laboratories 1/4 City , STATE

• Responsible for routine preparations of samples for Gas Chromatography analysis as per EPA methods (8270, 640, etc.).

• Maintained proper documentation and used required lab instrumentation in accordance to set SOP's.

#### Qualifications

- Subject Matter Expert in multiple analytical techniques including HPLC, GC, Dissolution, UV/Vis Spectroscopy
- Excellent troubleshooting skills
- Detail oriented while ensuring project timelines are met.
- Proficient in the analysis of multiple dosage forms with emphasis on tablets and emulsions.

#### Certifications

Lean Six Sigma Yellow Belt (December 2011)

#### Presentations

##### Poster Presentations

• Bambrick, S.; "Development of an HPLC Method for the Determination of Lyso Degradant of Excipient Hydrogenated Phosphatidylcholine in reformulated Elocon Cream 0.1% using Charged Aerosol Detection (CAD)" 2013 Summit Analytical Community Day

• He, H.; Zhang, F.; Bambrick, S.; Haug, J.; Goldfarb, D.; Verduyck, T.;

Harris, D.; Liang, S.; and Waters, C.; "Investigation of Compound E Formation in Reformulated Elocon Cream as a function of Internal Phase pH and Raw Material (Aluminum Starch Octenylsuccinate) pH" 2012 Merck Manufacturing Division Science and Technology Days Conference

• Goldfarb, D.; Harris, D.; Verduyck, T.; Castermans, K.; Goor, S.; Vaes, T.; Tully, R.; Xiong, L.; Bambrick, S.; Henderson, A.; and Yee, O.; "Process Development in a Manufacturing Environment: Collaboration on Elocon Cream reformulation between PCT and Supply" 2010 Science and Technology Days Innovation, Collaboration, and Compliance

• Sigvardson, K.; Bambrick, S.; Christopher, D.; Dey, M.; Fett, J.; Nijhuis, M.; and Wynia, G.; "A General Experimental Approach for the Separation of Method and Product Variability in Content Uniformity Test Methods" 2009 Pharmaceutical Sciences and Drug Metabolism Science Symposium

• Drake, J.; Kachar, L.; Bambrick, S.; Xu, J.; and Fett J.; "NIR Method for Content Uniformity Determination of Commercial-Scale Batches Based on Laboratory-Scale Calibration Models" 2009 Pharmaceutical Sciences and Drug Metabolism Science Symposium

#### Computer Skills

- Waters Empower 2 and 3
- Electronic Laboratory Notebook
- Research LIMS
- Minitab (ver. 15, 16, and 17)
- JMP (ver. 9 and 10)
- Microsoft Office
- Trackwise
- ProCal
- UV/Vis Chemstation (version A.10.01)

## Organizations

• Boy Scouts of America (1989 – 1995)

## Awards and Recognition

• Merck Commercialization Award (2nd Half 2015)

• Merck Award of Excellence (2010 - 2016)

• Schering-Plough Shining Performance Appreciation Award (2004 - 2010)

• Schering-Plough Best of Everything Award (2002 and 2003)

• New York Institute of Technology - Deans List (1995 – 1999)

• Boy Scouts of America - Eagle Scout (1995)