



**COMPLIANCE  
INSIGHT Inc.**  
**NANCY WATTS**

## **OBJECTIVE**

To advance and develop my career with my professional experience and educational background.

## **EXPERIENCE**

**5/2010 – 7/2011** **Forest Pharmaceuticals, Inc., Cincinnati, OH**  
3/2011- 7/2011 - *Data Review Supervisor, Quality Control*: Continue all previous duties, supervision of the data review group, train associates on cGMP laboratory documentation, advise Quality Assurance on laboratory audit processes.

5/2010-2/2011 - *Contract Quality Specialist, Quality Control*: Lead change control processes for the laboratory, IQ/OQ and Method Validation protocol/report writing, SOP development and update, analytical test method revisions, other CFR/ICH-GCP guideline improvements. Assist in Corrective Actions/Preventative Actions (CAPA) completion. Velquest/LabWare/Trackwise/Documentum

**11/2004 – 2/2010** **Patheon - PDS, Cincinnati, OH**  
*Chemist, Pharmaceutical Development Services (PDS)*: Drug assay via Dissolution, HPLC and other Wet Chemistry techniques, associated data analysis with PeakPro and EMPOWER data systems. Method verification, validation, and transfer including report and method writing. (part-time associate 9/2007 - 2/2010) Focus on documentation and analytical data review, training associates on review, including LIMS data storage entry and review. SOP revision and other cGMP, ICH-GCP guideline improvements.

**8/2002 – 10/2004** **Kendle International, Cincinnati, OH**  
9/2003-10/2004 - *Safety Specialist I*: Review and analyze adverse events occurring during clinical trials and other special projects for clients, report events per FDA guidelines using federally approved forms and client databases (also used for tracking and review purposes).  
8/2002 – 9/2003 - *Study Documentation Specialist*: Collection, tracking and organization of federally regulated documents associated with a clinical trial per GCP and ICH guidelines, site regulatory binders, site selection and management, quality assurance review of investigator files and site regulatory documents, informed consent and IRB submissions, main contact with IRB and sites (on a per study basis).

**7/2001 – 5/2002** **PerkinElmer Life Sciences, San Francisco, CA**  
*Genetics Research Analyst*: LC-MS/MS, Genetic Disease Research, Bloodspot analysis, Train colleagues on assay and instrumentation, assist in sample assay development, lab/instrumental implementation and validation, validation report compilation, data analysis and reporting, analytical method development and optimization. Sciex/Wallac/Gilson/Perkin-Elmer instrumentation/software.

**1/2001 – 6/2001** **Duramed Pharmaceuticals, Inc., Cincinnati, OH**  
*Contract Chemist*: Use of HPLC, UV-Vis, FTIR, TLC, Dissolution, and Karl Fischer Titration for the assay of finished product pharmaceuticals, stability and associated data as assigned by a team leader.

**9/1999 – 11/2000** **Alkermes, Inc., Cincinnati, OH**  
*Chemistry Associate*: GC-FID (HP 5890), GPC (RI, ELSD), DSC, HPLC, IR, Analyze biodegradable polymers for use in pharmaceutical delivery systems, train colleagues on assay and instrumentation.

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[www.Compliance-Insight.com](http://www.Compliance-Insight.com)



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## **EDUCATION**

**Bachelor of Science** in Chemistry, Forensic Emphasis  
Cum Laude, March 1999

**Ohio University**  
Athens, OH

Honors and Privileges: Arts and Sciences Dean's Scholarship, Alpha Lambda Delta Honor's Fraternity, Study Abroad at University of Wales, Swansea (Fall Qtr 1998).

Specialty Training:

\*Techniques of Capillary Gas Chromatography training course from HP, Penn State University, 2/2000

\*Waters GPC Theory/Instrumentation and Millennium software training course, Milford, MA, 6/2000

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