Sherrie Liddell

451 IH-35 Georgetown, Texas 78628 sherrie.liddell@gmail.com 224-216-4033

Biographical Profile

Over 10 years of proven regulatory compliance adherence in chemical and pharmaceutical drug manufacture data and processes from raw material receipt, bulk product processing, finished product compounding and fill, product packaging, product market-release, to customer complaints.

— Key Qualifications —

- Accomplished quality professional with expertise in applying compliance regulations and updates, comprehension of analytical procedures including compendia approaches; ascertaining compliance within the laboratory environment and within manufacturing operations, including safety precautions; and manufacture risks assessment by auditing internal processes and its associated data.
- Extremely organized and detail-attentive in thorough review of quality control and assurance data, authorship of quality documents including management review summaries and corrective/preventive action documents.
- Efficient and accurate in management of quality data review and processes. Forthcoming in conveying compliance issues, risks assessments, and their resolve in written or oral formats.
- Balanced multi-tasker in fast paced manufacture environments. Assert compliance and quality expertise to manufacture companies. Welcome new responsibilities to incorporate compliance responsibilities in industry.

Professional Experience

First Priority, Inc. – Elgin, Illinois
COMPLIANCE MANAGEMENT, 1/2013 to 7/2015

Responsible for the daily compliance of quality data and processes. Also responsible for coaching personnel to maintain Good Laboratory Practices and Good Documentation Practices, conducting training of test procedures and compendia updates, observation and assurance of Good Manufacturing Practices.

Selected Contributions:

- 100% accuracy in individual compliance data upon FDA on-site audit data review.
- Authored a greater than 125 page corrective action/prevention action summary of Stability Program deficiencies and gaps.
- Boosted team good data and reduced repeat testing by laboratory personnel to a 90% accuracy rate through thorough data review, analyses mentoring, and coaching.

Kelly Scientific Contractual Placement at Alcon (A Novartis Company) – Des Plaines, Illinois
VALIDATION CONSULTANT, 11/2011 to 3/2012

Incorporated project directives for approaches of validating historic test methods with technological efficiency to newly derived test methods. Assessed instrument validation risk gaps. Authored validation summaries with analytical outcomes.

Continued…