Focus Compliance & Validation Services

Pharmaceutical Packaging Capabilities

www.FocusCVS.com

DEMONSTRATED PROFESSIONAL EXPERIENCE

Within the pharmaceutical industry, FDA and other regulatory agencies exist to establish governance and guidance criteria establishing current good manufacturing practices (cGMP) including validation of machinery and systems, component specific validation, quality control activities, noncompliance and CAPA related activities, good documentation practices, etc. We offer our clients a unique blend of technical expertise, regulatory compliance insight, and practical hands-on experience in the pharmaceutical packaging environment.

PROFESSIONAL EXPERTISE IN PAHRMACEUTICAL PACKAGING

- SOP and Procedure Development
- Environmental Monitoring Procedure Development
- Validation Document Development
- AQL Sampling Plans
- Packaging Specification Development
- Internal Auditing Services
- Nonconformance & Complaint Management
- Validation & Qualification
- Part 11 (Electronic Records and Signatures)
- Mock Recalls
- Training
- Process Engineering
- Documentation Management Systems

FOCUS COMPLIANCE & VALIDATION SERVICES

Focus offers its clients a high level of experience. Focus personnel have extensive hands-on experience in the contract pharmaceutical, packaging industry. Our professional expertise and experience allows us to identify and implement the best solutions for our clients' specific applications.

FOCUS HISTORY

Established in November 1988, Focus has successfully completed projects for the pharmaceutical, medical device, biotechnology, chemical, explosives, petrochemical, metallurgical, textiles, and general manufacturing industries. These projects have been conducted throughout the United States, Puerto Rico, and more than 10 countries.

Why Choose Focus?

Immediate Impact

Based on our hands-on experience, Focus is able to interface quickly with you existing systems to determine where your company's and Focus resources are needed. Critical items will be given priority while time and effort ($costs) savings will be realized for low-impact issues.

Lasting Impact

Your organization's participants will benefit from the ability to gear up with additional resources as demanded by production or QA demands require.

Give Us A Call

Please give us a call or visit our Website – http://www.FocusCVS.com — we believe we can help you find the right solution.

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