



Array Validation Quality & Compliance, Inc.

We Deliver an Array of Dependable Compliance Solutions.

AVQC, Inc. is a Multinational Life Sciences Global Compliance Consulting Firm with 25 + years experience delivering Technical Writing & Validation Services in the Pharmaceutical, Medical Device, Biotechnology and Food Industries. We comply with FDA, MHRA, TGA, Health Canada, EMEA and ISO 90001, 14971 and 13485.

24/7 Customer
Service for Domestic
& International Clients:
+1.877.772.6375

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solutions@avqc.com

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4590 Deodar St.,
Silver Springs, NV 89429

We Ensure Your Compliance

We work closely with our clients to capture requirements, develop deliverable schedules, and deliver final high quality documentation on time and on budget.

Our Team Delivers Quality Service

Our team of professional consultants has decades of experience in a variety of areas in the Pharmaceutical, Medical Device, Biotechnology and Food Industries, serving both Domestic and International clients.

We Deliver a Full Range of Materials

We deliver a full range of Pharmaceutical and Medical Device Consulting Services in the areas of Compliance and Validation to help ensure your firm obtains and maintains GxP compliance.

Visit us at www.avqc.com to Schedule Your Free Consultation.



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We Deliver an Array of Regulatory, Validation, Quality and Compliance Solutions Designed to Exceed Your Expectations. **Contact Us to Design Your Solution.**

Regulatory, Quality & Compliance Services

- Strategic Planning for FDA Facility Approval
- Facility and Pre-approval Audits
- 483, Warning Letter & Consent Decree Remediation Projects
- On-site assistance during FDA inspections
- Quality System Development and Implementation
- CAPA Management
- Supplier Controls
- Customized Training Programs
- Batch Record Review
- Audit Prep, Vendor & 3rd Party audits
- Regulatory Submissions

Validation Services

- Equipment Validation
- Cleaning Validation
- Process Validation
- Test Method Validation
- Utilities Validation
- Computer System Validation
- Laboratory Instrument Validation
- Data Integrity Assessments
- Gap Analysis
- Technical & Procedural Remediation
- Creation of Functional; User Requirements; and Design Specifications
- Commissioning Documents, Site Acceptance Tests, Test Plans & Protocols, Protocol Execution and Validation Testing
- Site Validation Master Plans, Validation Master Plans, and Summary Reports

Technical Services

- Verification of Piping and Instrumentation Diagrams, Isometric Drawings, and Loop Sheets
- Document Research, Data Analysis, Feasibility Studies
- Creation, Modification & Revision of Technical Manuals, Training Manuals, Administrative Guides, White Papers and Standard Operating Procedures