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## David W. Husman, Ph.D., ASQ CPGP, RAC President and Principal Consultant, at David Husman Consulting, LLC

Dr. David Husman has more than 30 years of diverse international experience in quality assurance, quality control, and regulatory affairs within the pharmaceutical, biopharmaceutical, and biotech industries.

David is fully versed in all quality systems and has used his expertise to help numerous clients assess, develop, and improve their quality programs and compliance with US, EU, and worldwide regulatory agency requirements. He effectively interacts and communicates with all team members, including c-suite executives, scientists, managers, directors, and others.

David is certified in Good Manufacturing Practices (CPGP) and regulatory affairs (RAC). Having earned a doctorate in biochemistry, his expertise includes, but is not limited to:

- Compliance
- · Quality Systems
- Process Mapping
- Design and Implementation
- Auditing
- Validation
- Aseptic Processing
- Date Integrity
- Information Systems

David has worked with products that include both human and animal parenterals, medical devices, human tissues, blood products, biotech products, solid dosages, and vaccines.

David is experienced as a director, project manager, and auditor in vaccines, allergenic extracts, drug licensing, manufacturing, and quality. He has held several positions in quality assurance, quality control, and regulatory affairs in pharmaceutical, drug, and medical device industries. He has managed quality control laboratories, quality assurance and regulatory affairs departments, clinical trials, as well as numerous audits and regulatory inspections.

David is established as an effective trainer and skilled technical reviewer/editor. He has published articles on Data Integrity and Deviation Management, and has been a featured speaker at several conferences, including SQA, RAPS, and IVT. He earned his bachelor's degree and doctorate in biochemistry from The Ohio State University in Columbus, Ohio.