



# Process, Product, Equipment & Software Validation

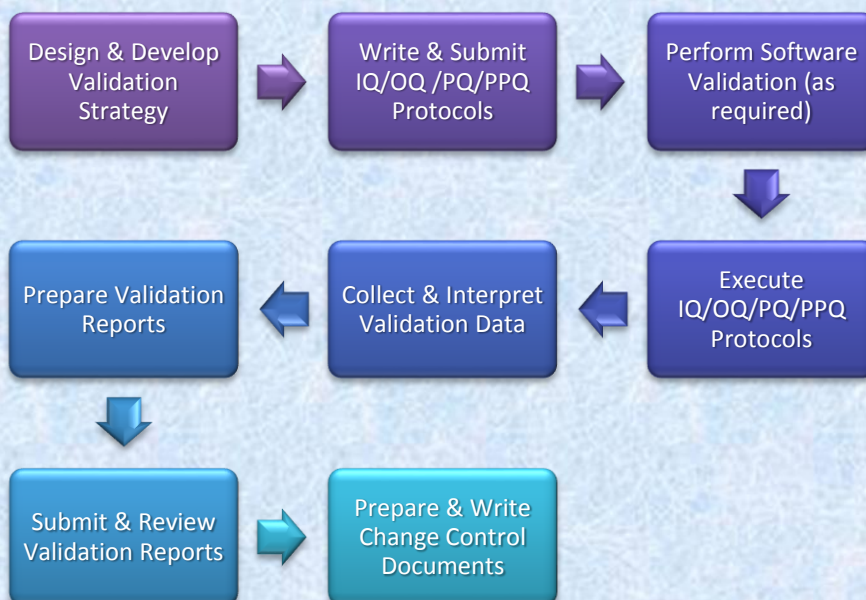
At BEPC, our mission is to provide our medical devices and biotech clients with innovative consulting services focused on continuous improvement through the implementation of Six-Sigma, Lean, and other related methodologies at an exceptional value.

The FDA defines validation as a process “establishing documented evidence which provides a high degree of assurance that a specific process will consistently produce a product meeting its pre-determined specifications and quality attributes.” BEPC utilizes the latest validation methods and industry knowledge to help our clients successfully comply with an evolving cGMP regulatory environment. We provide the following **Validation** services, including protocol generation and implementation:

- IQ (Installation Qualification)
- OQ (Operation Qualification)
- PQ (Product Qualification)
- PPQ (Process Performance Qualification)
- CSV (Computer Systems Validation)

Our Validation consultants are Subject Matter Experts (SMEs) in their respective fields and combines extensive training with years of on-site experience. With an international North American network of highly qualified consultants, BEPC is committed to fulfilling our clients' **Validation** needs.

## Process, Product, Equipment & Software Validation Methodology



### The BEPC Advantage

*Why should our clients choose BEPC?*

- Each client receives dedicated attention and a customized solution for their specific needs
- Flexibility to contract short or long term, fulltime or part time services
- Competitive negotiated rates with Volume Discounts
- Our network of consultants covers the US East, Central and West Coasts, as well as Mexico, minimizing our clients' costs
- High quality consultants ensures high quality results; our consultants are:
  - Highly skilled and qualified
  - Professional and committed
  - Experienced with short learning curves

### Contact Us:

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