Qualification

Sterile dis- | connectors

Benefits

- 100% traceable documentation right from the beginning
- All required certificates provided
- Comprehensive IQ | OQ documentation
- Compliance with the requirements of GLP | GMP standards
- Strict Sartorius test standards
- Specially trained service technicians will support you during your product qualification

Product Information

The qualification of equipment and manufacturing systems is mandatory in many sectors of the pharmaceutical and regulated industries, and is a prerequisite for later process validation. Sartorius will support you with specially trained experts in GLP- | GMP-compliant equipment qualification. We will provide reports specifically written for your equipment as documentation that you can easily integrated into your existing quality management systems.

Specifications

- Equipment qualification and documentation will be done based on the equipment-specific qualification documents
- Testing and documentation of the equipment supplied and review of the accompanying documents
- Function testing according to equipment specifications along with documentation of the results

The qualification is available for the following sterile dis- | connectors:

- BioWelder®
- BioWelder® TC | BioSealer®

Customer Prerequisites

- The equipment is already at the place of qualification
- Sufficient space for access to the equipment is available
- Power supply is available at the place of qualification
- The customer must designate the person responsible at the customer's site when placing an order

Interested? Just get in touch with us for a free service consultation. You will find all important information on our website at www.sartorius.com/service.

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