

Sterilization Validation and Routine Operation Handbook: Ethylene Oxide (Sterilization Validation and Routine Operation Handbook

Series)

Anne F. Booth



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Stringent regulations require you to validate sterilization processes and step-by-step guidelines are needed to develop and implement a suitable validation program. Sterilization Validation and Routine Operation Handbook: Ethylene Oxide is the best practical guide available for the validation of EtO process. The information provided complies with ANSI/AAMI/ISO 11135: 1994, Medical devices-Validation and routine control of ethylene oxide sterilization which is based on a standard developed by the European Standardization Committee (CEN) entitled EN 550, Sterilization of medical devices- Validation and routine control of ethylene oxide sterilization. The text defines methods to assist you in the interpretation and understanding of the requirements in the standard and offers logical procedures for the validation and routine monitoring of your specific ethylene oxide process.

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