

QUALITY PLAN

June 27, 2020

Part I **signoff**

Part II

revision history

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7.5.1 Control of production and service provision

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7.5.3 Installation Activities

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7.5.5 Particular requirements for sterile medical devices

7.5.6 Validation of processes for production and service provision

7.5.7 Particular requirements for validation of processes for sterilization and sterile barrier systems

7.5.8 Identification

7.5.9 Traceability

General

Particular requirements for implantable medical devices – N/A

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7.5.11 Preservation of product

7.6 Control of monitoring and measuring equipment

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