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Requirements for the development,
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Specifies requirements for the
development, validation, and routine
control of a moist heat sterilization
process for medical
devices. ANSI/AAMI/ISO 17665-1:2006
(R2013) - Sterilization of ... This is a
revision of AAMI TIR13:1997, and with
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Technical Specification provides general
guidance on the development, validation
and routine control of moist heat
sterilization processes and is intended to
explain the requirements set forth in ISO
17665-1. The guidance given in this
Technical Specification is provided to
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heat sterilization processes and to assist
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active air removal systems;ISO - ISO 17665-1:2006 - Sterilization of health care ...ANSI/AAMI/ISO 17665-1:2006 -- Sterilization of health care products - Moist heat - Part 1 Requirements for the development, validation and routine control of a sterilization process for medical devices Paperback – January 1, 2006ANSI/AAMI/ISO 17665-1:2006 -- Sterilization of health care ...The adoption of ISO Technical Specification (TS) 17665-3, as an AAMI Technical Information Report was initiated by the AAMI Radiation Sterilization Working Group, which also functions as the U.S. Technical Advisory Group to the relevant work in the International Organization for Standardization (ISO). U.S. representatives from the AAMI ISO 17665-1:2006 Sterilization of health

care products - Moist heat - Part 1: Requirements for the development, validation and routine control of a sterilization process for medical devices. ISO 17665-1:2006 specifies requirements for the development, validation and routine control of a moist heat sterilization process for medical devices.

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The adoption of ISO Technical Specification (TS) 17665-3, as an AAMI Technical Information Report was initiated by the AAMI Radiation Sterilization Working Group, which also functions as the U.S. Technical Advisory Group to the relevant work in the International Organization for Standardization (ISO). U.S. representatives from the AAMI

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