

Standards Publications Available from AAMI

QUALITY MANAGEMENT, GENERAL SAFETY, AND DESIGN

ANSI/AAMI ES60601-1:2005 (IEC 60601-1:2005, MOD) and C1:2009	Medical electrical equipment, Part 1: General requirements for basic safety and essential performance, 1ed and 2009 Amendment
ANSI/AAMI HE74:2001(R)2009	Human factors design process for medical devices, 1ed
AAMI HE75:2009	Human factors engineering - Design of medical devices, 3ed
ANSI/AAMI/IEC 62304:2006	Medical device software - Software life cycle processes, 2ed
ANSI/AAMI/IEC 60601-1-2:2007	Medical electrical equipment, Part 1-2: General requirements for basic safety and essential performance—Collateral standard: Electromagnetic compatibility—Requirements and tests, 2ed
ANSI/AAMI/ISO 13485:2003(R)2009	Medical devices—Quality management systems—System requirements for regulatory purposes, 2ed
ANSI/AAMI/ISO TIR14969:2004	Medical devices—Quality management systems—Guidance on the application of ISO 13485, 2ed
ANSI/AAMI/ISO 14971:2007	Medical devices—Application of risk management to medical devices, 3ed
ANSI/AAMI/ISO 15223-1:2007 and A1:2008	Medical devices - Symbols to be used with medical device labels, labeling, and information to be supplied - Part 1: General requirements, 1ed and 2008 Amendment
ANSI/AAMI/ISO 15225:2000(R)2006 and A1:2004(R)2006	Nomenclature—Specification for a nomenclature system for medical devices for the purpose of regulatory data exchange, 1ed and 2004 Amendment
AAMI TIR18:1997	Guidance on electromagnetic compatibility of medical devices for clinical/biomedical engineers—Part 1: Radiated radio-frequency electromagnetic energy, 1ed
AAMI TIR24:1999	Acquisition and use of physiologic waveform databases for testing of medical devices, 1ed
AAMI TIR32:2004	Medical device software risk management, 1ed
AAMI TIR36:2007	Validation of software for regulated processes, 1ed
ANSI/AAMI/IEC TIR60878:2003	Graphical symbols for electrical equipment in medical practice, 1ed
ANSI/AAMI/IEC TIR62296:2003	Considerations of unaddressed safety aspects in the Second Edition of IEC 60601-1 and proposals for new requirements, 1ed
ANSI/AAMI/IEC TIR62348:2006	Mapping between the clauses of the third edition of IEC 60601-1 and the 1988 edition as amended, 1ed
ANSI/AAMI/IEC TIR80002-1:2009	Medical device software - Part 1: Guidance on the application of ISO 14971 to medical device software, 1ed
ANSI/AAMI/ISO TIR16142:2005	Medical devices—Guidance on the selection of standards in support of recognized essential principles of safety and performance of medical devices
ANSI/AAMI/ISO TIR19218:2005	Medical devices - Coding structure for adverse event type and cause, 1ed

BIOLOGICAL EVALUATION OF MEDICAL DEVICES

ANSI/AAMI/ISO 10993-1:2009	Part 1: Evaluation and testing within a risk management process, 4ed
ANSI/AAMI/ISO 10993-2:2006	Part 2: Animal welfare requirements, 2ed
ANSI/AAMI/ISO 10993-3:2003(R)2009	Part 3: Tests for genotoxicity, carcinogenicity, and reproductive toxicity, 2ed
ANSI/AAMI/ISO 10993-4:2002 and A1:2006	Part 4: Selection of tests for interactions with blood, 2ed and 2006 Amendment
ANSI/AAMI/ISO 10993-5:2009	Part 5: Tests for <i>in vitro</i> cytotoxicity, 3ed
ANSI/AAMI/ISO 10993-6:2007	Part 6: Tests for local effects after implantation, 2ed
ANSI/AAMI/ISO 10993-7:2008	Part 7: Ethylene oxide sterilization residuals, 3ed
ANSI/AAMI/ISO 10993-9:1999(R)2005	Part 9: Framework for identification and quantification of potential degradation products, 2ed
ANSI/AAMI/ISO BE78:2002(R)2008 and A1:2006(R)2008	Part 10: Tests for irritation and delayed type hypersensitivity, 2ed (<i>adoption of ISO 10993-10:2002 with national deviation</i>) and 2006 Amendment (identical to ISO amendment)
ANSI/AAMI/ISO 10993-11:2006	Part 11: Tests for systemic toxicity, 2ed
ANSI/AAMI/ISO 10993-12:2007	Part 12: Sample preparation and reference materials, 3ed
ANSI/AAMI/ISO 10993-13:1999(R)2004	Part 13: Identification and quantification of degradation products from polymeric medical devices, 1ed
ANSI/AAMI/ISO 10993-14:2001(R)2006	Part 14: Identification and quantification of degradation products from ceramics, 1ed
ANSI/AAMI/ISO 10993-15:2000(R)2006	Part 15: Identification and quantification of degradation products from metals and alloys, 1ed
ANSI/AAMI/ISO 10993-16:1997(R)2003	Part 16: Toxicokinetic study design for degradation products and leachables from medical devices, 1ed
ANSI/AAMI/ISO 10993-17:2002(R)2008	Part 17: Establishment of allowable limits for leachable substances, 1ed
ANSI/AAMI/ISO BE83:2006	Part 18: Chemical characterization of materials
ANSI/AAMI/ISO 14155-1:2003(R)2008	Clinical investigation of medical devices for human subjects—Part 1: General requirements, 1ed
ANSI/AAMI/ISO 14155-2:2003(R)2008	Clinical investigation of medical devices for human subjects—Part 2: Clinical investigation plans, 1ed
ANSI/AAMI/ISO 22442-1:2007	Medical devices utilizing animal tissues and their derivatives - Part 1: Application of risk management, 1ed
ANSI/AAMI/ISO 22442-2:2007	Medical devices utilizing animal tissues and their derivatives - Part 2: Controls on sourcing, collection and handling, 1ed
ANSI/AAMI/ISO 22442-3:2007	Medical devices utilizing animal tissues and their derivatives - Part 3: Validation of the elimination and/or inactivation of viruses and transmissible spongiform encephalopathy (TSE) agents, 1ed
AAMI TIR19:1998 and A1:1999	Guidance for ANSI/AAMI/ISO 10993-7:1995, Biological evaluation of medical devices—Part 7: Ethylene oxide sterilization residuals, 1ed and 1999 Amendment

ANSI/AAMI/ISO TIR10993-19:2006	Biological evaluation of medical devices - Part 19: Physio-chemical, morphological and topographical characterization of materials, 1ed
ANSI/AAMI/ISO TIR10993-20:2006	Biological evaluation of medical devices - Part 20: Principles and methods for immunotoxicology testing of medical devices, 1ed

BIOMEDICAL EQUIPMENT

ANSI/AAMI AT6:2005	Autologous transfusion devices, 3ed
ANSI/AAMI BF7:1989(R)2007	Blood transfusion micro-filters, 2ed
ANSI/AAMI BF64:2002(R)2007	Leukocyte reduction filters, 1ed
ANSI/AAMI BP22:1994(R)2006	Blood pressure transducers, 2ed
ANSI/AAMI DF80:2003	Medical electrical equipment, Part 2-4: Particular requirements for the safety of cardiac defibrillators [including automated external defibrillators], 4ed
ANSI/AAMI EC11:1991(R)2007	Diagnostic electrocardiographic devices, 2ed
ANSI/AAMI EC12:2000(R)2005	Disposable ECG electrodes, 3ed
ANSI/AAMI EC13:2002(R)2007	Cardiac monitors, heart rate meters, and alarms, 3ed
ANSI/AAMI EC38:2007	Medical electrical equipment - Part 2-47: Particular requirements for the safety, including essential performance, of ambulatory electrocardiographic systems, 3ed
ANSI/AAMI EC53:1995(R)2008 and A1:1998(R)2008	ECG cables and leadwires, 1ed and 1998 Amendment
ANSI/AAMI EC57:1998(R)2008	Testing and reporting performance results of cardiac rhythm and ST segment measurement algorithms, 2ed
ANSI/AAMI EC71:2001(R)2007	Standard communications protocol—Computer assisted electrocardiography, 1ed
ANSI/AAMI EQ56:1999(R)2008	Recommended practices for a medical equipment management program, 1ed
ANSI/AAMI ID26:2004(R)2009	Medical electrical equipment, Part 2: Particular requirements for the safety of infusion pumps and controllers, 3ed
ANSI/AAMI ID54:1996(R)2005	Enteral feeding set adapters and connectors, 1ed
AAMI NS4:1986(R)2009	Transcutaneous electrical nerve stimulators, 1ed
ANSI/AAMI NS28:1988(R)2006	Intracranial pressure monitoring devices, 1ed
ANSI/AAMI PB70:2003(R)2009	Liquid barrier performance and classification of protective apparel and drapes in health care facilities, 1ed
ANSI/AAMI PC69:2007	Active implantable medical devices—Electromagnetic compatibility—EMC test protocols for implantable cardiac pacemakers and implantable cardioverter defibrillators, 2ed
ANSI/AAMI/IEC 60601-2-2:2009	Medical electrical equipment, Part 2-2: Particular requirements for basic safety and essential performance of high frequency surgical equipment and high frequency surgical accessories, 5ed
ANSI/AAMI/IEC 60601-2-19:2009	Medical electrical equipment - Part 2-19: Particular requirements for basic safety and essential performance of baby incubators, 4ed
ANSI/AAMI/IEC 60601-2-20:2009	Medical electrical equipment - Part 2-20: Particular requirements for basic safety and essential performance of transport incubators, 3ed
ANSI/AAMI/IEC 60601-2-21:2009	Medical electrical equipment - Part 2-21: Particular requirements for basic safety and essential performance of infant radiant warmers, 2ed
ANSI/AAMI/IEC 60601-2-50:2009	Medical electrical equipment - Part 2-50: Particular requirements for basic safety and essential performance of infant phototherapy equipment, 2ed
ANSI/AAMI/IEC 80601-2-58:2008	Medical electrical equipment - Part 2-58: Particular requirements for basic safety and essential performance of lens removal and vitrectomy devices for ophthalmic surgery, 1ed
ANSI/AAMI/IEC 80601-2-30:2009	Medical electrical equipment - Part 2-30: Particular requirements for basic safety and essential performance of automated type non-invasive sphygmomanometers, 1ed
ANSI/AAMI/ISO 5840:2005	Cardiovascular implants—Cardiac valve prostheses, 4ed
ANSI/AAMI/ISO 7198:1998/2001(R)2004	Cardiovascular implants—Tubular vascular prostheses, 3ed
ANSI/AAMI/ISO 7199:2009	Cardiovascular implants and artificial organs—Blood-gas exchangers (oxygenators), 2ed
ANSI/AAMI/ISO 14708-3:2008	Implants for surgery - Active implantable medical devices - Part 3: Implantable neurostimulators, 1ed
ANSI/AAMI/ISO 14708-4:2008	Implants for surgery - Active implantable medical devices - Part 4: Implantable infusion pumps, 1ed
ANSI/AAMI/ISO 15674:2009	Cardiovascular implants and artificial organs—Hard-shell cardiomy/venous reservoir systems (with/without filter) and soft venous reservoir bags, 2ed
ANSI/AAMI/ISO 15675:2009	Cardiovascular implants and artificial organs—Cardiopulmonary bypass systems—Arterial line blood filters, 2ed
ANSI/AAMI/ISO 25539-1:2003(R)2009 and A1:2005(R)2009	Cardiovascular implants—Endovascular devices—Part 1: Endovascular prostheses, 1ed and 2005 Amendment
ANSI/AAMI/ISO 25539-2:2008	Cardiovascular implants - Endovascular devices - Part 2: Vascular stents, 1ed
ANSI/AAMI/ISO 81060-1:2007	Non invasive sphygmomanometers - Part 1: Requirements and test methods for non-automated measurement type, 1ed
ANSI/AAMI/ISO 81060-2:2009	Non-invasive sphygmomanometers - Part 2: Clinical validation of automated measurement type, 1ed.
AAMI TIR4:1989	Apnea monitoring by means of thoracic impedance pneumography, 1ed
AAMI TIR9:1992	Evaluation of clinical systems for invasive blood pressure monitoring, 1ed
AAMI TIR11:2005	Selection and use of protective apparel and surgical drapes in health care facilities, 2ed
AAMI TIR21:1998	Systems used to forecast remaining pacemaker battery service life, 1ed
AAMI TIR23:1999	Signal averaging, 1ed
AAMI TIR26:2000	Ventricular assist and heart replacement systems, 1ed

Standards Publications Available from AAMI (continued)

STERILIZATION

<p>ANSI/AAMI ST8:2008 Hospital steam sterilizers, 5ed</p> <p>ANSI/AAMI ST24:1999/(R)2009 Automatic, general-purpose ethylene oxide sterilizers and ethylene oxide sterilant sources intended for use in health care facilities, 3ed</p> <p>ANSI/AAMI ST40:2004 Table-top dry heat (heated air) sterilization and sterility assurance in health care facilities, 2ed</p> <p>ANSI/AAMI ST41:2008 Ethylene oxide sterilization in health care facilities: Safety and effectiveness, 4ed</p> <p>ANSI/AAMI ST50:2004 Dry heat (heated air) sterilizers, 2ed</p> <p>ANSI/AAMI ST55:2003/(R)2008 Table-top steam sterilizers, 2ed</p> <p>ANSI/AAMI ST58:2005 Chemical sterilization and high-level disinfection in health care facilities, 2ed</p> <p>ANSI/AAMI ST63:2002 Sterilization of health care products—Requirements for the development, validation and routine control of an industrial sterilization process for medical devices—Dry heat, 1ed</p> <p>ANSI/AAMI ST65:2008 Processing of reusable surgical textiles for use in health care facilities, 2ed</p> <p>ANSI/AAMI ST67:2003/(R)2008 Sterilization of health care products—Requirements for products labeled 'sterile', 1ed</p> <p>ANSI/AAMI ST72:2002 Bacterial endotoxins—Test methodologies, routine monitoring and alternatives to batch testing, 1ed</p> <p>ANSI/AAMI ST77:2006 Containment devices for reusable medical device sterilization, 1ed</p> <p>ANSI/AAMI ST79:2006, A1:2008 and A2:2009 Comprehensive guide to steam sterilization and sterility assurance in health care facilities, 1ed and Amendments 1 and 2</p> <p>ANSI/AAMI ST81:2004 Sterilization of medical devices - Information to be provided by the manufacturer for the processing of resterilizable medical devices, 1ed</p> <p>ANSI/AAMI/ISO 10993-7:2008 Biological evaluation of medical devices, Part 7: Ethylene oxide sterilization residuals, 3ed</p> <p>ANSI/AAMI/ISO 11135-1:2007 Sterilization of health care products - Ethylene oxide - Part 1: Requirements for the development, validation and routine control of a sterilization process for medical devices, 4ed</p> <p>TIR11135-2:2008 Sterilization of health care products - Ethylene oxide - Part 2: Guidance on the application of AAMI/ISO 11135-1, 1ed</p> <p>ANSI/AAMI/ISO 11137-1:2006 Sterilization of health care products - Radiation - Part 1: Requirements for the development, validation and routine control of a sterilization process for medical devices</p> <p>ANSI/AAMI/ISO 11137-2:2006 Sterilization of health care products - Radiation - Part 2: Establishing the sterilization dose</p> <p>ANSI/AAMI/ISO 11137-3:2006 Sterilization of health care products - Radiation - Part 3: Guidance on dosimetric aspects</p> <p>ANSI/AAMI/ISO 11138-1:2006 Sterilization of health care products - Biological indicators - Part 1: General requirements, 2ed</p> <p>ANSI/AAMI/ISO 11138-2:2006 Sterilization of health care products - Biological indicators - Part 2: Biological indicators for ethylene oxide sterilization processes, 3ed</p> <p>ANSI/AAMI/ISO 11138-3:2006 Sterilization of health care products - Biological indicators - Part 3: Biological indicators for moist heat sterilization processes</p> <p>ANSI/AAMI/ISO 11138-4:2006 Sterilization of health care products - Biological indicators - Part 4: Biological indicators for dry heat sterilization processes</p> <p>ANSI/AAMI/ISO 11138-5:2006 Sterilization of health care products - Biological indicators - Part 5: Biological indicators for low-temperature steam and formaldehyde sterilization processes</p> <p>ANSI/AAMI/ISO 11140-1:2005 Sterilization of health care products - Chemical indicators - Part 1: General requirements, 2ed</p> <p>ANSI/AAMI/ISO 11140-3:2007 Sterilization of health care products - Chemical indicators - Part 3: Class 2 indicator systems for use in the Bowie and Dick steam penetration test</p> <p>ANSI/AAMI/ISO 11140-4:2007 Sterilization of health care products - Chemical indicators - Part 4: Class 2 indicators as an alternative to Bowie and Dick test for detection of steam penetration</p> <p>ANSI/AAMI/ISO 11140-5:2007 Sterilization of health care products - Chemical indicators - Part 5: Class 2 indicators for Bowie and Dick air removal test sheets and packs, 2ed</p> <p>ANSI/AAMI/ISO 11607-1:2006 Packaging for terminally sterilized medical devices - Part 1: Requirements for materials, sterile barrier systems and packaging, 3ed</p> <p>ANSI/AAMI/ISO 11607-2:2006 Packaging for terminally sterilized medical devices - Part 2: Validation requirements for forming, sealing and assembly processes, 1ed</p> <p>ANSI/AAMI/ISO 11737-1:2006 Sterilization of medical devices - Microbiological methods - Part 1: Determination of a population of microorganisms on products, 2ed</p> <p>ANSI/AAMI/ISO 11737-2:2009 Sterilization of medical devices—Microbiological methods—Part 2: Tests of sterility performed in the definition, validation and maintenance of a sterilization process, 2ed</p> <p>ANSI/AAMI/ISO 13408-1:2008 Aseptic processing of health care products - Part 1: General requirements, 1ed</p> <p>ANSI/AAMI/ISO 13408-2:2003 Aseptic processing of health care products - Part 2: Filtration, 1ed</p> <p>ANSI/AAMI/ISO 13408-3:2006 Aseptic processing of health care products - Part 3: Lyophilization, 1ed</p> <p>ANSI/AAMI/ISO 13408-4:2005 Aseptic processing of health care products - Part 4: Clean-in-place technologies, 1ed</p> <p>ANSI/AAMI/ISO 13408-5:2006 Aseptic processing of health care products - Part 5: Sterilization in place, 1ed</p> <p>ANSI/AAMI/ISO 13408-6:2005 Aseptic processing of health care products - Part 6: Isolator systems</p> <p>ANSI/AAMI/ISO 14160:1998/(R)2008 Sterilization of single-use medical devices incorporating materials of animal origin—Validation and routine control of sterilization by liquid chemical sterilants, 1ed</p>	<p>ANSI/AAMI/ISO 14161:2009 Sterilization of health care products—Biological indicators—Guidance for the selection, use, and interpretation of results, 3ed</p> <p>ANSI/AAMI/ISO 14937:2009 Sterilization of health care products—General requirements for characterization of a sterilizing agent and the development, validation and routine control of a sterilization process for medical devices, 2ed</p> <p>ANSI/AAMI/ISO 15882:2008 Chemical indicators—Guidance for selection, use, and interpretation of results, 4ed</p> <p>ANSI/AAMI ST15883-1:2009 (ISO 15883-1:2006, MOD) Washer-disinfectors, Part 1: General requirements, terms and definitions and tests, 1ed</p> <p>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, 3ed</p> <p>ANSI/AAMI/ISO 18472:2006 Sterilization of health care products - Biological and chemical indicators - Test equipment, 1ed</p> <p>AAMI TIR12:2004 Designing, testing and labeling reusable medical devices for reprocessing in health care facilities: A guide for device manufacturers, 2ed</p> <p>AAMI TIR14:2009 Contract sterilization for ethylene oxide, 2ed</p> <p>AAMI TIR15:2009 Physical aspects of ethylene oxide sterilization, 2ed</p> <p>AAMI TIR16:2009 Microbiological aspects of ethylene oxide sterilization, 2ed</p> <p>AAMI TIR17:2008 Compatibility of materials subject to sterilization, 2ed</p> <p>AAMI TIR22:2007 and A1:2008 Guidance for ANSI/AAMI/ISO 11607, Packaging for terminally sterilized medical devices - Part 1 and Part 2:2006, 2ed and 2008 Amendment</p> <p>AAMI TIR28:2009 Product adoption and process equivalence for ethylene oxide sterilization, 2ed</p> <p>AAMI TIR29:2002 Guide for process control in radiation sterilization, 1ed</p> <p>AAMI TIR30:2003 A compendium of processes, materials, test methods, and acceptance criteria for cleaning reusable medical devices, 1ed</p> <p>AAMI TIR31:2008 Process challenge devices/test packs for use in healthcare facilities, 2ed</p> <p>AAMI TIR33:2005 Sterilization of health care products - Radiation - Substantiation of a selected sterilization dose - Method V_{Dmax}, 1ed</p> <p>AAMI TIR34:2007 Water for the reprocessing of medical devices, 1ed</p> <p>AAMI TIR35:2006 Sterilization of health care products - Radiation sterilization - Alternative sampling plans for verification dose experiments and sterilization dose audits, 1ed</p> <p>AAMI TIR37:2007 Sterilization of health care products—Radiation—Guidance on sterilization of human tissue-based products, 1ed</p> <p>AAMI TIR39:2009 Guidance on selecting a microbial challenge and inoculation sites for sterilization validation of medical devices, 1ed</p> <p>AAMI TIR40:2009 Sterilization of health care products - Radiation - Guidance on dose setting utilizing a Modified Method 2, 1ed</p> <p>ANSI/AAMI/ISO TIR11139:2006 Sterilization of health care products—Vocabulary, 2ed</p> <p>ANSI/AAMI/ISO TIR17665-2:2009 Sterilization of health care products - Moist heat - Part 2: Guidance on the application of ANSI/AAMI/ISO 17665-1, 1ed</p>
DIALYSIS	
<p>ANSI/AAMI RD5:2003/(R)2008 Hemodialysis systems, 3ed</p> <p>ANSI/AAMI RD16:2007 Cardiovascular implants and artificial organs - Hemodialyzers, hemodiafilters, hemofilters and hemoconcentrators, 3ed</p> <p>ANSI/AAMI RD17:2007 Cardiovascular implants and artificial organs - Extracorporeal blood circuit for hemodialyzers, hemodiafilters and hemofilters, 3ed</p> <p>ANSI/AAMI RD47:2008 Reprocessing of hemodialyzers, 4ed</p> <p>ANSI/AAMI RD52:2004 and Amendments 1 to 4 Dialysate for hemodialysis, 1ed (includes A1:2007, A2:2007, A3:2009 and A4:2009)</p> <p>ANSI/AAMI RD52:2004/A3:2009 and A4:2009 Dialysate for hemodialysis, Amendment s 3 and 4, 1ed</p> <p>ANSI/AAMI RD52:2004/A2:2007 Dialysate for hemodialysis, Amendment 2 - Annex D: Self-assessment of compliance with recommendations for dialysate preparation, 1ed</p> <p>ANSI/AAMI RD61:2006 Concentrates for hemodialysis, 2ed</p> <p>ANSI/AAMI RD62:2006 and A1:2009 Water treatment equipment for hemodialysis applications, 2ed and 2009 amendment</p>	<p>ANSI/AAMI RD5:2003/(R)2008 Hemodialysis systems, 3ed</p> <p>ANSI/AAMI RD16:2007 Cardiovascular implants and artificial organs - Hemodialyzers, hemodiafilters, hemofilters and hemoconcentrators, 3ed</p> <p>ANSI/AAMI RD17:2007 Cardiovascular implants and artificial organs - Extracorporeal blood circuit for hemodialyzers, hemodiafilters and hemofilters, 3ed</p> <p>ANSI/AAMI RD47:2008 Reprocessing of hemodialyzers, 4ed</p> <p>ANSI/AAMI RD52:2004 and Amendments 1 to 4 Dialysate for hemodialysis, 1ed (includes A1:2007, A2:2007, A3:2009 and A4:2009)</p> <p>ANSI/AAMI RD52:2004/A3:2009 and A4:2009 Dialysate for hemodialysis, Amendment s 3 and 4, 1ed</p> <p>ANSI/AAMI RD52:2004/A2:2007 Dialysate for hemodialysis, Amendment 2 - Annex D: Self-assessment of compliance with recommendations for dialysate preparation, 1ed</p> <p>ANSI/AAMI RD61:2006 Concentrates for hemodialysis, 2ed</p> <p>ANSI/AAMI RD62:2006 and A1:2009 Water treatment equipment for hemodialysis applications, 2ed and 2009 amendment</p>
AAMI STANDARDS COLLECTIONS	
<p>STBK09-1 Sterilization, Part 1: Sterilization in Health Care Facilities (2009) (book)</p> <p>STBK09-2 Sterilization, Part 2: Sterilization Equipment Design and Use (2009) (book)</p> <p>STBK09-3 Sterilization, Part 3: Industrial Process Control (2009) (book)</p> <p>STBK09-S Sterilization 3-Book Set (Current Editions of Parts 1-3) (books)</p> <p>STBKCD AAMI Standards on CD—Sterilization Edition (2009)</p> <p>DSBK08 Dialysis (2008) (book)</p> <p>DSBKCD AAMI Standards on CD—Dialysis Edition (2008)</p> <p>DSBK08-S Dialysis Set (Current edition of book and CD)</p> <p>DSBK08-PDF AAMI Standards on PDF—Dialysis Edition (2008; immediate download of complete collection)</p> <p>BIOTCD AAMI Standards on CD—Biological Evaluation of Medical Devices Series (2009)</p> <p>SYMBCD Medical Equipment Symbols and Safety Signs—Clip Art Collection (CD)</p> <p>STDSCD AAMI Standards on CD (subscription service)</p>	<p>STBK09-1 Sterilization, Part 1: Sterilization in Health Care Facilities (2009) (book)</p> <p>STBK09-2 Sterilization, Part 2: Sterilization Equipment Design and Use (2009) (book)</p> <p>STBK09-3 Sterilization, Part 3: Industrial Process Control (2009) (book)</p> <p>STBK09-S Sterilization 3-Book Set (Current Editions of Parts 1-3) (books)</p> <p>STBKCD AAMI Standards on CD—Sterilization Edition (2009)</p> <p>DSBK08 Dialysis (2008) (book)</p> <p>DSBKCD AAMI Standards on CD—Dialysis Edition (2008)</p> <p>DSBK08-S Dialysis Set (Current edition of book and CD)</p> <p>DSBK08-PDF AAMI Standards on PDF—Dialysis Edition (2008; immediate download of complete collection)</p> <p>BIOTCD AAMI Standards on CD—Biological Evaluation of Medical Devices Series (2009)</p> <p>SYMBCD Medical Equipment Symbols and Safety Signs—Clip Art Collection (CD)</p> <p>STDSCD AAMI Standards on CD (subscription service)</p>

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