

NATIONAL STANDARDS

Recently Published

NEW! ANSI/AAMI/ISO 10993-15:2019, *Biological evaluation of medical devices—Part 15: Identification and quantification of degradation products from metals and alloys*. [Purchase here](#).

NEW! AAMI TIR105:2020, *Risk management guidance for combination products*. [Purchase here](#).

ANSI/AAMI/IEC 80601-2-77:2020, *Medical electrical equipment—Part 2-77: Particular requirements for the basic safety and essential performance of robotically assisted surgical equipment*. [Purchase here](#).

ANSI/AAMI/IEC 80601-2-78:2020, *Medical electrical equipment—Part 2-78: Particular requirements for basic safety and essential performance of medical robots for rehabilitation, assessment, compensation or alleviation*. [Purchase here](#).

ANSI/AAMI/ISO 8637-3:2018, *Extracorporeal systems for blood purification – Part 3: Plasmafilters*. [Purchase here](#).

AAMI Call for Comments

If you would like to comment on one of the draft documents listed below, contact the individual indicated by e-mail to receive a PDF copy of the draft. These copies are free.

Published documents proposed for reaffirmation can be purchased from the [AAMI Store](#).

Comments due November 16, 2020

AAMI/IEC 60601-2-20, Medical electrical equipment - Part 2-20: Particular requirements for the basic safety and essential performance of infant transport incubators (proposed revised American National Standard). Applies to the basic safety and essential performance of transport incubators. This standard does not apply to heating devices intended for physiotherapy, baby incubators, radiant warmers.

Contact: jmoyer@aami.org

New Work

AAMI/CN, Small Bore Connectors Committee is working on the revision of AAMI/ISO 80369-7, *Small-bore connectors for liquids and gases in healthcare applications — Part 7: Connectors for intravascular or hypodermic applications*. Contact: [Colleen Elliott](#)

AAMI EQ, Medical Equipment Management Committee. The committee is working on the development of AAMI EQ110/Ed.1, *Guidance for health care technology management education programs*. This standard seeks to provide a recommended framework for new or established education programs in health care technology management (HTM). Contact: [Amanda Benedict](#).

AAMI QM-WG01, Application of Quality Systems to Medical Devices Working Group. The working group is working on the development of AAMI CR510, Consensus Report on Compliant Use of Cloud Computing for Quality Systems and Medical Devices. This Consensus Report will provide guidance to multiple stakeholders regarding the appropriate and compliant use of cloud computing both as a component of medical devices and in support of quality systems. Contact: [Joe Lewelling](#).

Consensus Body Members Needed

The committees listed below are seeking new members in the indicated stakeholder interest category to participate in the development of their documents. Definitions of the various stakeholder groups are:

User: *An individual or organizational representative, who purchases, utilizes or receives the materials, products, systems, or services covered in the scope of technical documents developed by AAMI in the delivery of healthcare; individuals in this interest category include clinicians, employees or representatives of Healthcare Delivery Organizations, clinical consultants, patients, etc.*

Industry: *An individual or organizational representative involved in the commercial production, promotion, sale, use or distribution of materials, products, systems, or services covered in the scope of technical documents developed by AAMI; this interest category includes manufacturers, those involved in supply chains, employees of test labs or commercial labs, industry consultants, etc.*

Regulatory: *An individual or organizational representative involved in the regulation of the materials, products, systems, or services covered in the scope of the technical documents developed by AAMI; this interest category includes those representing federal, state, local, foreign, or other government entities.*

General interest: *An individual or organizational representative with a general material interest in the materials, products, systems, or services covered in the scope of the technical documents developed by AAMI and who does not fit into any of the preceding categories; this interest category can include noncommercial academicians, noncommercial researchers, patient or*

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consumer advocates, representatives of accrediting organizations, representatives of other organizations, etc.

Please contact the staff person indicated for more information on how to join.

AAMI/CN, Small Bore Connectors Committee. The committee is seeking user, regulatory and general interest members to participate in the development of AAMI/ISO 80369-7, *Small-bore connectors for liquids and gases in healthcare applications — Part 7: Connectors for intravascular or hypodermic applications*. Contact: [Colleen Elliott](#)

AAMI EQ, Medical Equipment Management Committee. The committee is seeking industry, regulatory and general interest members to participate in the development of AAMI EQ110/Ed.1, *Guidance for health care technology management education programs*. Contact: [Amanda Benedict](#).

AAMI ST/WG 91, Resistometers. The working group is seeking user, industry, and regulatory/general interest stakeholders to participate in the US adoption of ISO 18472:2018, *Sterilization of health care equipment – Biological and chemical indicators – Test equipment*. Contact: [Cliff Bernier](#).

AAMI TIB, Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use. The committee and its affiliated working groups are seeking user, industry, and general interest/regulator members to participate in developing the US position towards documents under development in ISO/TC 76, *Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use, and other projects*.

AAMI/CV, Cardiac valves. The committee is seeking user, industry, and general interest/regulator members to participate in the US adoption of ISO 5840-1:202x, *Cardiovascular implants — Cardiac valve prostheses — Part 1: General requirements*; ISO 5840-2:202x, *Cardiovascular implants — Cardiac valve prostheses — Part 2: Surgically implanted heart valve substitutes*; ISO 5840-3:202x, *Cardiovascular implants — Cardiac valve prostheses — Part 3: Heart valve substitutes implanted by transcatheter techniques*; and the revision of ISO 5910:2018, *Cardiovascular implants and extracorporeal systems — Cardiac valve repair devices*. Contact: [Cliff Bernier](#).

AAMI/VP, Vascular Prostheses. The committee is seeking user, industry, and general interest/regulator members to participate in the US adoption of ISO 25539-2:2020, *Cardiovascular implants — Endovascular devices — Part 2: Vascular stents*; the revision of ISO 25539-3, *Cardiovascular implants — Endovascular devices — Part 3: Vena cava filters*; and the development of ISO 25539-4, *Cardiovascular implants — Endovascular devices — Part 4: Application of ISO 17327-1 for coated endovascular devices*. Contact: [Cliff Bernier](#)

AAMI/BG, Blood/Gas Exchange Device Committee. The committee is seeking user, industry, and general interest/regulator members to participate in the development of the following Cardiovascular implants and artificial organs documents: ISO 18193, *Cannulae for extracorporeal circulation*; Amendment 1 to ISO 18242:2016 *Centrifugal blood pumps for pulsatile pumps*; and revision of ISO 7199, *Blood-gas exchangers*. Contact: [Cliff Bernier](#)

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AAMI/VP-WG 01, Vascular Device-Drug Combination Products. The committee is seeking user, industry, and general interest/regulator members to participate in the revision of ISO 12417-1:2015, *Cardiovascular implants and extracorporeal systems — Vascular device-drug combination products — Part 1: General requirements* and the revision of ISO/TR 12417-2:2017, *Cardiovascular implants and extracorporeal systems — Vascular device-drug combination products — Part 2: Local regulatory information*. Contact: [Cliff Bernier](#)

AAMI/VI, Cardiovascular absorbable implants. The committee is seeking user, industry, and general interest/regulator members to participate in the revision of ISO/TS 17137:2019, *Cardiovascular implants and extracorporeal systems - Cardiovascular absorbable implants*. Contact: [Cliff Bernier](#)

AAMI/CO, Cardiac Occluders. The committee is seeking user, industry, and general interest/regulator members to participate in the development of ISO 22679, *Cardiovascular implants — Transcatheter cardiac occluders*. Contact: [Cliff Bernier](#)

AAMI/RD, Renal Disease and Detoxification Committee. The committee is seeking industry, user, and general interest/regulator members to participate in the revision of the ISO 23500:2019, Preparation and quality management of fluids for haemodialysis and related therapies series standards: Part 1: General requirements; Part 2: Water treatment equipment for haemodialysis applications and related therapies; Part 3, Water for haemodialysis and related therapies; Part 4: Concentrates for haemodialysis and related therapies; Part 5, Quality of dialysis fluids for haemodialysis and related therapies; and the revision of the ISO 8637, Extracorporeal systems for blood purification series standards: Part 1, Haemodialysers, haemodiafilters, haemofilters, and haemoconcentrators,;Part 2: Extracorporeal blood circuit for haemodialysers, haemodiafilters, and haemofilters; and Part 3, Plasmafilters. Contact: [Cliff Bernier](#)

AAMI/HIT, Health IT Committee. The working group is seeking industry, general interest, and regulator members to participate in the development of AAMI HIT1000-1/Ed., *Health IT Software and Systems – Part 1: Fundamental concepts and principles*. Contact: [Emily Hoefler](#)

AAMI/HIT-WG02, Health IT Quality Systems Working Group. The working group is seeking industry, general interest, and regulator members to participate in the development of AAMI HIT1000-2/Ed., *Health IT software and systems — Part 2: Application of quality systems principles and practices* (Provisional Standard). Contact: [Emily Hoefler](#)

AAMI/II, Infant Incubators. The committee is seeking users, regulatory, and general interest members to participate in the development of AAMI/IEC 60601-2-20, *Medical electrical equipment - Part 2-20: Particular requirements for the basic safety and essential performance of infant transport incubators*. Contact: [Jennifer Moyer](#)

UPCOMING MEETINGS

AAMI Committees and U.S. TAGs

Open meetings of AAMI committees and U.S. Technical Advisory Groups (TAGs) are listed here and at the AAMI website (www.aami.org). Note: If you plan to attend a meeting, please send a brief note to the AAMI Standards Department (standards@aami.org) indicating the name and date of the meeting so that staff can contact you in the event of a last-minute cancellation.

December 2020

Biological Indicators Working Group (open web meeting). 9 December 2020, 12:00h to 15:00h ET. Virtual meeting. Contact: [Amanda Benedict](#).

Vaporized hydrogen peroxide sterilization Working Group (open web meeting). Date and time TBD. Virtual meeting. Contact: [Amanda Benedict](#).

INTERNATIONAL STANDARDS

Information on draft international standards under ballot can be found in [ANSI Standards Action](#).

International Committee and Working Group Meetings

Call or email the indicated staff person at AAMI for more information about upcoming international standards meetings.

December 2020

ISO/TC 198/WG 9, Aseptic processing (closed meeting), 3 December 2020, 06:00 h to 10:00 h ET, Zoom meeting. Contact: [Amanda Benedict](#)

ISO/TC 198/WG 8, Microbiological methods (closed meeting), 8 December 2020, 09:00 h to 10:00 h ET, Zoom meeting. Contact: [Amanda Benedict](#)

ISO/TC 198/WG 7, Packaging (closed meeting), 8-9 December 2020, 07:00 h to 11:00 h ET each day, Zoom meeting. Contact: [Amanda Benedict](#)

February 2021

ISO/TC 198/WG 1, Industrial EO sterilization (closed meeting), 23 February 2021, 08:00 h to 10:00 h ET, Zoom meeting. Contact: [Amanda Benedict](#)