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CERTIFICATE OF ACCREDITATION

This is to attest that

CSA GROUP JAPAN

YOKOHAMA CONNECT SQUARE 12 FL., 3-3-3 MINATOMIRAI, NISHI-KU
YOKOHAMA-SHI, KANAGAWA, 220-0012, JAPAN

Testing Laboratory TL-685

has met the requirements of AC89, *IAS Accreditation Criteria for Testing Laboratories as well as the FDA ASCA Program specifications*, and has demonstrated compliance with ISO/IEC Standard 17025:2017, *General requirements for the competence of testing and calibration laboratories*. This organization is accredited to provide the services specified in the scope of accreditation.

Effective Date February 3, 2025



International Accreditation Service
Issued under the authority of IAS management

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SCOPE OF ACCREDITATION

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Contact Name Wayne Thomas

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Accredited to ISO/IEC 17025:2017

Effective Date February 3, 2025

FDA ASCA Program

FDA ASCA Program Scope

| Basic Safety and Essential Performance of Medical Electrical Equipment, Medical Electrical Systems and Laboratory Medical Equipment | |
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| ANSI AAMI ES60601-1:2005/(R)2012 & A1:2012, C1:2009/(R)2012 & A2:2010/(R)2012 (Cons. Text) [Incl. AMD2:2021] [19-46] | Medical electrical equipment - Part 1: General requirements for basic safety and essential performance (IEC 60601-1:2005, MOD) [Including Amendment 2 (2021)] |
| IEC 60601-1-3 Edition 2.2 2021-01 CONSOLIDATED VERSION [12-336] | Medical electrical equipment - Part 1-3: General requirements for basic safety and essential performance - Collateral Standard: Radiation protection in diagnostic X-ray equipment |
| IEC 60601-1-6 Edition 3.2 2020-07 CONSOLIDATED VERSION [5-132] | Medical electrical equipment – Part 1-6: General requirements for basic safety and essential performance – Collateral standard: Usability |
| IEC 60601-2-18: Edition 3.0 2009-08 [9-114] | Medical electrical equipment - part 2-18: particular requirements for the basic safety and essential performance of endoscopic equipment |
| IEC 60601-2-22 Edition 3.1 2012-10 [12-268] | Medical electrical equipment - part 2-22: particular requirements for the basic safety and essential performance of surgical cosmetic therapeutic and diagnostic laser equipment |
| IEC 60601-2-28 Edition 3.0 2017-06 [12-309] | Medical electrical equipment - part 2-28: particular requirements for the basic safety and essential performance of x-ray tube assemblies for medical diagnosis |
| IEC 60601-2-33 Ed. 3.2 b:2015 [12-295] | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| IEC 60601-2-37 Edition 2.1 2015 [12-293] | Medical electrical equipment - Part 2-37: particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment |
| IEC 60601-2-43 Edition 3.0 2022-12 [12-351] | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of X-ray equipment for interventional procedures |

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| IEC 60601-2-43 Edition 2.2 2019-10 CONSOLIDATED VERSION [12-329] | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of X-ray equipment for interventional procedures |
| IEC 60601-2-44 Edition 3.2: 2016 [12-302] | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| IEC 60601-2-45 Edition 3.1 2015 [12-294] | Medical electrical equipment - part 2-45: particular requirements for basic safety and essential performance of mammographic x- ray equipment and mammographic stereotactic devices |
| IEC 60601-2-54 Edition 2.0 2022-09 [12-348] | Medical electrical equipment - Part 2-54: particular requirements for the basic safety and essential performance of X-ray equipment for radiography and radioscopy |
| IEC 61010-1 Edition 3.1 2017-01 CONSOLIDATED VERSION [19-34] | Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements [Including: Corrigendum 1 (2019)] - Note: This standard is recognized with relevant US national differences applied, see reference #1 in Relevant FDA Guidance and/or Supportive Publication section |

Regular Scope

| Electrical and Electronics | |
|-----------------------------------|--|
| ANSI/AAMI ES60601-1 | Medical electrical equipment— part 1: general requirements for basic safety and essential performance |
| ANSI/AAMI ES60601-1+ AMD1 | Medical electrical equipment— part 1: general requirements for basic safety and essential performance |
| ANSI/AAMI ES60601-1+ AMD2 | Medical electrical equipment— part 1: general requirements for basic safety and essential performance |
| CAN/CSA C22.2 No. 14 | Industrial Control Equipment |
| CAN/CSA C22.2 No. 60065 | Audio, video and similar electronic apparatus - safety requirements |
| CAN/CSA C22.2 No. 60601-1 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| CAN/CSA C22.2 No. 60601-1+AMD1 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| CAN/CSA C22.2 No. 60601-1+AMD2 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| CAN/CSA C22.2 No. 60601-1-3 | Medical electrical equipment – part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| CAN/CSA C22.2 No. 60601-1-3+AMD1 | Medical electrical equipment – part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| CAN/CSA C22.2 No. 60601-1-3+AMD2 | Medical electrical equipment - Part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| CAN/CSA C22.2 No. 60601-1-6 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |

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| CAN/CSA C22.2 No. 60601-1-6+AMD1 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |
| CAN/CSA C22.2 No. 60601-1-6+AMD2 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |
| CAN/CSA C22.2 No. 60601-2-18 | Medical electrical equipment - part 2-18: particular requirements for the basic safety and essential performance of endoscopic equipment |
| CAN/CSA C22.2 No. 60601-2-18+AMD1 | Medical electrical equipment - part 2-18: particular requirements for the basic safety and essential performance of endoscopic equipment |
| CAN/CSA C22.2 No. 60601-2-28 | Medical electrical equipment - part 2-28: particular requirements for the basic safety and essential performance of x-ray tube assemblies for medical diagnosis |
| CAN/CSA C22.2 No. 60601-2-33 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| CAN/CSA C22.2 No. 60601-2-33+AMD1 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| CAN/CSA C22.2 No. 60601-2-33+AMD2 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| CAN/CSA C22.2 No. 60601-2-37 | Medical electrical equipment - part 2-37: particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment |
| CAN/CSA C22.2 No. 60601-2-37+AMD1 | Medical electrical equipment - part 2-37: particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment |
| CAN/CSA C22.2 No. 60601-2-43 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| CAN/CSA C22.2 No. 60601-2-43+AMD1 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| CAN/CSA C22.2 No. 60601-2-43+AMD2 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| CAN/CSA C22.2 No. 60601-2-44 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| CAN/CSA C22.2 No. 60601-2-44+AMD1 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| CAN/CSA C22.2 No. 60601-2-44+AMD2 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| CAN/CSA C22.2 No. 60601-2-45 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| CAN/CSA C22.2 No. 60601-2-45+AMD1 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| CAN/CSA C22.2 No. 60601-2-45+AMD2 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| CAN/CSA C22.2 No. 60601-2-54 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |

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| CAN/CSA C22.2 No. 60601-2-54+AMD1 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |
| CAN/CSA C22.2 No. 60601-2-54+AMD2 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |
| CSA C22.2 NO. 80601-2-60 | Medical electrical equipment – Part 2-60: particular requirements for the basic safety and essential performance of dental equipment |
| CAN/CSA C22.2 No. 60947-1 | Low-Voltage Switchgear and Controlgear – Part 1: General Rules |
| CAN/CSA C22.2 No. 60947-4-1 | Low-Voltage Switchgear and Controlgear – Part 4-1: Contactors and motor-starters-Electromechanical contactors and motor starters |
| CAN/CSA C22.2 No. 60947-4-2 | Low-Voltage Switchgear and Controlgear – Part 4-2: Contactors and motor-starters-AC semiconductor motor controllers and starters |
| CAN/CSA C22.2 No. 60947-5-1 | Low-Voltage Switchgear and Controlgear – Part 5-1: Control circuit devices and switching elements – Electromechanical control circuit devices |
| CAN/CSA C22.2 No. 60947-5-2 | Low-Voltage Switchgear and Controlgear – Part 5-2: Control circuit devices and switching elements – Proximity Switches |
| CAN/CSA C22.2 60947-7-1 | Low-Voltage Switchgear and Controlgear – Part 7-1: Ancillary equipment – Terminal Blocks for copper conductors |
| CAN/CSA C22.2 60947-7-2 | Low-Voltage Switchgear and Controlgear – Part 7-2: Ancillary Equipment – Protective Conductor Terminal Blocks for Copper Conductors |
| CAN/CSA C22.2 60947-7-3 | Low-Voltage Switchgear and Controlgear – Part 7-3: Ancillary Equipment – Safety Requirements for Fuse Terminal Blocks |
| CAN/CSA C22.2 60947-7-4 | Low-Voltage Switchgear and Controlgear – Part 7-4: Ancillary Equipment – PCB Terminal Blocks for Copper Conductors |
| CAN/CSA C22.2 No. 60950-1 | Information technology equipment - safety - part 1: general requirements |
| CAN/CSA C22.2 No. 60950-1+AMD1 | Information technology equipment - safety - part 1: general requirements |
| CAN/CSA C22.2 No. 60950-1+AMD2 | Information technology equipment - safety - part 1: general requirements |
| CAN/CSA C22.2 No. 61010-031 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 031: safety requirements for hand-held probe assemblies for electrical measurement and test |
| CAN/CSA C22.2 No. 61010-1 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 1: general requirements |
| CAN/CSA C22.2 No. 61010-1+AMD1 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 1: general requirements |
| CAN/CSA C22.2 No. 61010-2-081 | Safety requirements for electrical equipment for measurement, control and laboratory use - part 2-081: particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes |
| CAN/CSA C22.2 No. 61010-2-101 | Safety requirements for electrical equipment for measurement, control and laboratory use - part 2-101: particular requirements for in vitro diagnostic (IVD) medical equipment |
| CAN/CSA C22.2 No. 62304 | Medical device software - software life cycle processes |
| CAN/CSA C22.2 No. 62304+AMD1 | Medical device software - software life cycle processes |

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| CAN/CSA C22.2 No. 62368-1 | Audio/video, information and communication technology equipment - part 1: safety requirements |
| CAN/CSA-IEC 61010-2-201 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 2-201: particular requirements for control equipment |
| CSA C22.2 No. 55 | Special Use Switches |
| CSA C22.2 No. 65 | Wire Connectors |
| CSA C22.2 No. 113 | Fans and Ventilators (Applicable section: Requirements for component fans, clauses 9.1, 9.2, 9.3, 9.4, and 9.5) |
| CSA C22.2 No. 188 | Splicing Wire Connectors |
| CSA C22.2 No. 158 | Terminal Blocks |
| IEC 60601-2-28 | Medical electrical equipment - part 2-28: particular requirements for the basic safety and essential performance of x-ray tube assemblies for medical diagnosis |
| IEC 60601-2-43 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of X-ray equipment for interventional procedures |
| IEC 60601-2-54 | Medical electrical equipment - part 2- 54: particular requirements for the basic safety and essential performance of X-ray equipment for radiography and radioscopy |
| IEC 60601-2-54 Ed.2.0 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of X-ray equipment for radiography and radioscopy |
| IEC/EN 60065 | Audio, video and similar electronic apparatus - safety requirements |
| IEC/EN 60065+AMD1 | Audio, video and similar electronic apparatus - safety requirements |
| IEC/EN 60065+AMD2 | Audio, video and similar electronic apparatus - safety requirements |
| IEC/EN 60601-1 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| IEC/EN 60601-1+AMD1 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| IEC/EN 60601-1+AMD2 | Medical electrical equipment - part 1: general requirements for basic safety and essential performance |
| IEC/EN 60601-1-3 | Medical electrical equipment – part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| IEC/EN 60601-1-3+AMD1 | Medical electrical equipment – part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| IEC/EN 60601-1-3+AMD2 | Medical electrical equipment – part 1-3: General requirements for basic safety and essential performance - collateral standard: radiation protection in diagnostic x-ray equipment |
| IEC/EN 60601-1-6 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |
| IEC/EN 60601-1-6+AMD1 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |
| IEC/EN 60601-1-6+AMD2 | Medical electrical equipment - part 1-6: general requirements for basic safety and essential performance - collateral standard: usability |
| IEC/CSA 60601-1-9 | Medical electrical equipment – Part 1-9: General requirements for basic safety and essential performance – Collateral Standard: Requirements for environmentally conscious design |



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| IEC/CSA 60601-1-9 +AMD1 | Medical electrical equipment – Part 1-9: General requirements for basic safety and essential performance – Collateral Standard: Requirements for environmentally conscious design |
| IEC/CSA 60601-1-9 +AMD2 | Medical electrical equipment – Part 1-9: General requirements for basic safety and essential performance – Collateral Standard: Requirements for environmentally conscious design |
| IEC 80601-2-60 | Medical electrical equipment – Part 2-60: Particular requirements for the basic safety and essential performance of dental equipment |
| IEC/CSA/UL 61010-2-010 | Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-010: Particular requirements for laboratory equipment for the heating of materials |
| IEC/CSA/UL 61010-2-020 | Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-020: Particular requirements for laboratory centrifuges |
| IEC/CSA/UL 61010-2-030 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-030: particular requirements for equipment having testing or measuring circuits |
| IEC/CSA/UL 61010-2-032 | Safety requirements for electrical equipment for measurement, control and laboratory use – Part 2-032: particular requirements for hand-held and hand-manipulated current sensors for electrical test and measurement |
| IEC/CSA/UL 61010-2-033 | Safety requirements for electrical equipment for measurement, control and laboratory use – Part 2-033: particular requirements for hand-held multimeters and other meters for domestic and professional use, capable of measuring mains voltage |
| IEC/CSA/UL 61010-2-034 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-034: Particular requirements for measurement equipment for insulation resistance and test equipment for electric strength |
| IEC/CSA/UL 61010-2-040 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-040: Particular requirements for sterilizers and washer-disinfectors used to treat medical materials |
| IEC/CSA/UL 61010-2-061 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-061: Particular requirements for laboratory atomic spectrometers with thermal atomization and ionization |
| IEC/EN 60601-2-18 | Medical electrical equipment - part 2-18: particular requirements for the basic safety and essential performance of endoscopic equipment |
| IEC/EN 60601-2-18+AMD1 | Medical electrical equipment - part 2-18: particular requirements for the basic safety and essential performance of endoscopic equipment |
| IEC/EN 60601-2-28 | Medical electrical equipment - part 2-28: particular requirements for the basic safety and essential performance of x-ray tube assemblies for medical diagnosis |
| IEC/EN 60601-2-33 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| IEC/EN 60601-2-33+AMD1 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| IEC/EN 60601-2-33+AMD2 | Medical electrical equipment - part 2-33: particular requirements for the basic safety and essential performance of magnetic resonance equipment for medical diagnosis |
| IEC/EN 60601-2-37 | Medical electrical equipment - part 2-37: particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment |



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| IEC/EN 60601-2-43 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| IEC/EN 60601-2-43+AMD1 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| IEC/EN 60601-2-43+AMD2 | Medical electrical equipment - part 2-43: particular requirements for the basic safety and essential performance of x-ray equipment for interventional procedures |
| IEC/EN 60601-2-44 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| IEC/EN 60601-2-44+AMD1 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| IEC/EN 60601-2-44+AMD2 | Medical electrical equipment - part 2-44: particular requirements for the basic safety and essential performance of x-ray equipment for computed tomography |
| IEC/EN 60601-2-45 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| IEC/EN 60601-2-45+AMD1 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| IEC/EN 60601-2-45+AMD2 | Medical electrical equipment - part 2-45: particular requirements for the basic safety and essential performance of mammographic x-ray equipment and mammographic stereotactic devices |
| IEC/EN 60601-2-54 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |
| IEC/EN 60601-2-54+AMD1 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |
| IEC/EN 60601-2-54+AMD2 | Medical electrical equipment - part 2-54: particular requirements for the basic safety and essential performance of x-ray equipment for radiography and radioscopy |
| IEC/EN 60947-1 | Low-Voltage Switchgear and Controlgear – Part 1: General Rules |
| IEC/EN 60947-4-1 | Low-Voltage Switchgear and Controlgear – Part 4-1: Contactors and motor-starters-Electromechanical contactors and motor starters |
| IEC/EN 60947-4-2 | Low-Voltage Switchgear and Controlgear – Part 4-2: Contactors and motor-starters-AC semiconductor motor controllers and starters |
| IEC/EN 60947-4-3 | Low-Voltage Switchgear and Controlgear – Part 4-3: Contactors and motor-starters-AC semiconductor Controllers and contactors for non-motor loads |
| IEC/EN 60947-5-1 | Low-Voltage Switchgear and Controlgear – Part 5-1: Control circuit devices and switching elements – Electromechanical control circuit devices |
| IEC/EN 60947-5-2 | Low-Voltage Switchgear and Controlgear – Part 5-2: Control circuit devices and switching elements – Proximity Switches |
| IEC/EN 60947-7-1 | Low-Voltage Switchgear and Controlgear – Part 7-1: Ancillary equipment – Terminal Blocks for copper conductors |
| IEC/EN 60947-7-2 | Low-Voltage Switchgear and Controlgear – Part 7-2: Ancillary Equipment – Protective Conductor Terminal Blocks for Copper Conductors |
| IEC/EN 60947-7-3 | Low-Voltage Switchgear and Controlgear – Part 7-3: Ancillary Equipment – Safety Requirements for Fuse Terminal Blocks |



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| IEC/EN 60947-7-4 | Low-Voltage Switchgear and Controlgear – Part 7-4: Ancillary Equipment – PCB Terminal Blocks for Copper Conductors |
| IEC/EN 60950-1 | Information technology equipment - safety - part 1: general requirements |
| IEC/EN 60950-1+AMD1 | Information technology equipment - safety - part 1: general requirements |
| IEC/EN 60950-1+AMD2 | Information technology equipment - safety - part 1: general requirements |
| IEC/EN 61010-031 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 031: safety requirements for hand-held probe assemblies for electrical measurement and test |
| IEC/EN 61010-2-081 | Safety requirements for electrical equipment for measurement, control and laboratory use - part 2-081: particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes |
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| IEC/EN 61010-2-101 | Safety requirements for electrical equipment for measurement, control and laboratory use - part 2-101: particular requirements for in vitro diagnostic (IVD) medical equipment |
| IEC/EN 61010-2-201 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 2-201: particular requirements for control equipment |
| IEC/EN 62304 | Medical device software - software life cycle processes |
| IEC/EN 62304+AMD1 | Medical device software - software life cycle processes |
| IEC/EN 62368-1 | Audio/video, information and communication technology equipment - part 1: safety requirements |
| UL 486A-486B | UL Standard for Safety Wire Connectors |
| UL 486C | UL Standard for Safety Splicing Wire Connectors |
| UL 507 | Standard for Safety Electric Fans (Applicable section: Low Voltage Component Fans, clauses 189, 190, 191, 192, and 193) |
| UL 508 | Industrial Control Equipment |
| UL 1054 | UL Standard for Safety Special-Use Switches |
| UL 1059 | UL standard for Safety Terminal Blocks |
| UL 60065 | Standard for audio, video and similar electronic apparatus - safety requirements |
| UL 60601-1 | Medical electrical equipment, part 1: general requirements for safety |
| UL 60947-1 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 1: General Rules |
| UL 60947-4-1 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 4-1: Contactors and motor-starters-Electromechanical contactors and motor starters |
| UL 60947-5-2 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 5-2: Control circuit devices and switching elements – Proximity Switches |
| UL 60947-7-1 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 7-1: Ancillary equipment – Terminal Blocks for copper conductors |
| UL 60947-7-2 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 7-2: Ancillary Equipment – Protective Conductor Terminal Blocks for Copper Conductors |
| UL 60947-7-3 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 7-3: Ancillary Equipment – Safety Requirements for Fuse Terminal Blocks |



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| U L60947-7-4 | UL Standard for Safety Low-Voltage Switchgear and Controlgear – Part 7-4: Ancillary Equipment – PCB Terminal Blocks for Copper Conductors |
| UL 60950-1 | Information technology equipment - safety - part 1: general requirements |
| UL 60950-1+AMD1 | Information technology equipment - safety - part 1: general requirements |
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| UL 61010-1+AMD1 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 1: general requirements |
| UL 61010-2-101 | Safety requirements for electrical equipment for measurement, control and laboratory use - part 2-101: particular requirements for in vitro diagnostic (IVD) medical equipment |
| UL 61010-2-201 | Safety requirements for electrical equipment for measurement, control, and laboratory use - part 2-201: particular requirements for control equipment |
| UL 62368-1 | Audio/video, information and communication technology equipment - part 1: safety requirements |

