CERTIFICATE OF ACCREDITATION

This is to attest that

CSA INDIA PRIVATE LIMITED

EINSTEIN BUILDING, BIDARAHALLI HOBLI, WHITEFIELD ASHRAM ROAD (SH 35)
BANGALORE, KA, 560067, REPUBLIC OF INDIA

Testing Laboratory TL-1152

has met the requirements of AC89, *IAS Accreditation Criteria for Testing Laboratories*, 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 July 7, 2023

President

Visit www.iasonline.org for current accreditation information.

SCOPE OF ACCREDITATION

International Accreditation Service, Inc.

3060 Saturn Street, Suite 100, Brea, California 92821, U.S.A. | www.iasonline.org

CSA INDIA PRIVATE LIMITED

www.csagroup.org

Contact Name Wayne Thomas **Contact Phone** +44-7703613034

Accredited to ISO/IEC 17025:2017 *Effective Date July 7, 2023*

IT/AV	
AS/NZS 60065	Audio, video and similar electronic apparatus - Safety requirements Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
CSA C22.2 No. 60065	Audio, video and similar electronic apparatus - Safety requirements Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
CSA C22.2 No. 60950-1	Information technology equipment — Safety Part 1 General requirements Exclusion: 2.10.8.4, 3.2.5.1, 4.2.8, 4.3.12, Annex H, 4.3.13, A.3, Annex U, Annex AA, Annex CC.
CSA C22.2 No. 62368-1	Audio/video Information and Communication Technology Equipment – Part 1: Safety Requirements Exclusion: 5.4.1.10.2, 5.4.1.5.3, 5.4.4.6.5, G.10, Annex R, 10, Annex C, G.5.3.4, G.7.1, G.9, G.13.6, G.15, Annex J, M.7, M.8.2, S.3.2, Annex U, Y.3, Y.4
EN 60065	Audio, video and similar electronic apparatus - Safety requirements Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
EN 60950-1	Information technology equipment — Safety Part 1 General requirements Exclusion : 2.10.8.4, 3.2.5.1, 4.2.8, 4.3.12, Annex H, 4.3.13, A.3, Annex U, Annex AA, Annex CC.
EN 62368-1	Audio/video Information and Communication Technology Equipment – Part 1: Safety Requirements Exclusion: 5.4.1.10.2, 5.4.1.5.3, 5.4.4.6.5, G.10, Annex R, 10, Annex C, G.5.3.4, G.7.1, G.9, G.13.6, G.15, Annex J, M.7, M.8.2, S.3.2, Annex U, Y.3, Y.4
IEC 60065	Audio, video and similar electronic apparatus - Safety requirements Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
IEC 60950-1	Information technology equipment — Safety Part 1 General requirements Exclusion: 2.10.8.4, 3.2.5.1, 4.2.8, 4.3.12, Annex H, 4.3.13, A.3, Annex U, Annex AA, Annex CC.
IEC 62368-1	Audio/video Information and Communication Technology Equipment – Part 1: Safety Requirements Exclusion: 5.4.1.10.2, 5.4.1.5.3, 5.4.4.6.5, G.10, Annex R, 10, Annex C, G.5.3.4, G.7.1, G.9, G.13.6, G.15, Annex J, M.7, M.8.2, S.3.2, Annex U, Y.3, Y.4
IS 616	Audio, video and similar electronic apparatus - Safety requirements

	Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
IS 13252 (Part 1)	Information technology equipment — Safety Part 1 General requirements Exclusion: 2.10.8.4, 3.2.5.1, 4.2.8, 4.3.12, Annex H, 4.3.13, A.3, Annex U, Annex AA, Annex CC.
IS 16333 (Part 3:2022)	Mobile phone handsets: Part 3 Indian language support for mobile phone handsets - Specific requirements (Second Revision)
UL 60065	Audio, video and similar electronic apparatus - Safety requirements Exclusion: 6, 8.17, 8.21, 12.3, 13.6, 13.7, 14.2, 14.3, 14.6.4, 14.7, 16.1, 18, 20.2.4, Annex H
UL 60950-1	Information technology equipment — Safety Part 1 General requirements Exclusion: 2.10.8.4, 3.2.5.1, 4.2.8, 4.3.12, Annex H, 4.3.13, A.3, Annex U, Annex AA, Annex CC.
UL 62368-1	Audio/video Information and Communication Technology Equipment – Part 1: Safety Requirements Exclusion: 5.4.1.10.2, 5.4.1.5.3, 5.4.4.6.5, G.10, Annex R, 10, Annex C, G.5.3.4, G.7.1, G.9, G.13.6, G.15, Annex J, M.7, M.8.2, S.3.2, Annex U, Y.3, Y.4
Luminaires	
EN 61347	d.c or ac Supplied Electronic Controlgear for LED Modules Exclusion: 18.2
EN 61347-2-13	Particular requirements for d.c. or a.c. supplied electronic controlgear for LED modules Exclusion: Annex J, J.7 and Annex J, J.11
EN 62560	Self-ballasted LED-lamps for general lighting services by voltage > 50 V - Safety specifications Exclusion: 17
IEC 60598-02-1	Particular requirements - Fixed general-purpose luminaires Exclusion: 4.24/Annex P, 4.26, 4.34, 4.35, 9.2.9 to .11,12.5.1e), Annex C
IEC 60598-1	Luminaires - Part 1: General requirements and tests Exclusion: 4.24/Annex P, 4.26, 4.34, 4.35, 9.2.9 to .11, 12.5.1e), Annex C
IEC 61347-1	d.c or ac Supplied Electronic Controlgear for LED Modules Exclusion: 18.2
IEC 61347-2-13	Particular requirements for d.c. or a.c. supplied electronic controlgear for LED modules Exclusion: Annex J, J.7 and Annex J, J.11
IEC 62560	Self-ballasted LED-lamps for general lighting services by voltage > 50 V - Safety specifications Exclusion: 17
IS 10322 (Part 1)	Specification for luminaires Part 1 General requirements Exclusion: 4.24/Annex P, 4.26, 4.34, 4.35, 9.2.9 to .11, 12.5.1e), Annex C
IS 10322 (Part 5/Sec 1)	Part 5 Particular requirements Section 1 Fixed General Purpose Luminaires Exclusion: 4.24/Annex P, 4.26, 4.34, 4.35, 9.2.9 to .11, 12.5.1e), Annex C

IS 15885-2-13	Section XIII - Particular requirements for d.c. or a.c. supplied electronic controlgear for LED modules Exclusion: Annex J, J.7 and Annex J, J.11
IS 16102 (Part 1)	Self-ballasted led lamps for general lighting services Part 1 Safety requirements Exclusion: 17
Measuring Instruments	
CAN/CSA C22.2 No. 61010-1	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements Exclusion: 11.7, 12.2.1, 12.3, 12.4, 12.5.1, 12.5.2, 12.6, 13.2.3, 14.8
CAN/CSA C22.2 No. 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 Exclusion: 13.2.101
CAN/CSA C22.2 No. 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 Exclusion: 14.101, 101.2, 101.4
CAN/CSA C22.2 No. 61010-2-051	Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-051: Particular requirements for laboratory equipment for mixing and stirring Exclusion: 13.2.101
CAN/CSA C22.2 No. 61010-2-201	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-201: Particular requirements for control equipment
EN 61010-1	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements Exclusion: 11.7, 12.2.1, 12.3, 12.4, 12.5.1, 12.5.2, 12.6, 13.2.3, 14.8
IEC 61010-1	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements Exclusion: 11.7, 12.2.1, 12.3, 12.4, 12.5.1, 12.5.2, 12.6, 13.2.3, 14.8
IEC 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 Exclusion: 13.2.101
IEC 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 Exclusion: 14.101, 101.2, 101.4
IEC 61010-2-051	Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-051: Particular requirements for laboratory equipment for mixing and stirring Exclusion: 13.2.101
IEC 61010-2-201	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-201: Particular requirements for control equipment

UL 61010-1	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements Exclusion: 11.7, 12.2.1, 12.3, 12.4, 12.5.1, 12.5.2, 12.6, 13.2.3, 14.8
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 Exclusion: 13.2.101
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 Exclusion: 14.101, 101.2, 101.4
UL 61010-2-051	Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-051: Particular requirements for laboratory equipment for mixing and stirring Exclusion: 13.2.101
UL 61010-2-201	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-201: Particular requirements for control equipment
Household	
EN 60335-1	Household and similar electrical appliances - Safety - Part 1: General requirements Exclusion: 19.11.4.1, 19.11.4.2, 19.11.4.3, 19.11.4.4, 19.11.4.5, 19.11.4.6, 19.11.4.7, 22.16, 22.32, Annex H, Annex J, Annex R, Annex T, Annex U
EN 60335-2-14	Household and similar electrical appliances - Safety - Part 2-14: Particular requirements for kitchen machines Exclusion: 19.11.4.1, 19.11.4.2, 19.11.4.3, 19.11.4.4, 19.11.4.5, 19.11.4.6, 19.11.4.7, 22.16, 22.32, Annex H, Annex J, Annex R, Annex T, Annex U
IEC 60335-1	Household and similar electrical appliances - Safety - Part 1: General requirements Exclusion: 19.11.4.1, 19.11.4.2, 19.11.4.3, 19.11.4.4, 19.11.4.5, 19.11.4.6, 19.11.4.7, 22.16, 22.32, Annex H, Annex J, Annex R, Annex T, Annex U
IEC 60335-2-14	Household and similar electrical appliances - Safety - Part 2-14: Particular requirements for kitchen machines Exclusion: 19.11.4.1, 19.11.4.2, 19.11.4.3, 19.11.4.4, 19.11.4.5, 19.11.4.6, 19.11.4.7, 22.16, 22.32, Annex H, Annex J, Annex R, Annex T, Annex U
IS 302-1	Safety of household and similar electrical appliances, Part 1: General Requirements Exclusion: 19.11.4.1, 19.11.4.2, 19.11.4.3, 19.11.4.4, 22.16, 22.32, Annex H, Annex J, Annex R
Fire (Material Testing)	
ANSI/IEC 60529	Degrees of protection provided by enclosures (IP Code)
ASTM D635-18	Standard Test Method for Rate of Burning and/or Extent and Time of Burning of Plastics in a Horizontal Position
ASTM D3801	Standard Test Method for Measuring the Comparative Burning Characteristics of Solid Plastics in a Vertical Position

CSA C22.2 No. 0.17	Evaluation of Properties of Polymeric Materials (Inclusion from Cl. No 5.1 to 5.2.4 only)
CSA C22.2 No. 60529:16	Degrees of protection provided by enclosures (IP Code)
IEC 60529	Degrees of protection provided by enclosures (IP Code)
IEC 60695-2-11	Fire hazard testing – Part 2-11: Glowing/hot-wire based test methods – Glow-wire flammability test method for end products (GWEPT)
IEC 60695-10-2	Fire hazard testing - Part 10-2: Abnormal heat - Ball pressure test method
IEC 60695-11-5	Fire hazard testing - Part 11-5: Test flames - Needle-flame test method - Apparatus, confirmatory test arrangement and guidance
IEC 60695-11-10	Fire hazard testing - Part 11-10: Test flames - 50 W horizontal and vertical flame test methods
IS/IEC 60034: Part 5	Rotating electrical machines - Part 5: Degrees of protection provided by the integral design of rotating electrical machines (IP code) - Classification
IS/IEC 60529	Degrees of protection provided by enclosures (IP Code)
UL 94	Tests for Flammability of Plastic Materials for Parts in Devices and Appliances (Included Cl. No 7, 8 and 9 only)
UL 746A	Standard for polymeric materials - short term property evaluation (inclusion Cl. No 35 only)
Medical	
ANSI/AAMI ES60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance Exclusion: 10.1.1, 10.1.2, 10.2, 10.3
ANSI/AAMI HA60601-1-11	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment Exclusion: 10 Construction of ME Equipment
ANSI/AAMI/IEC 60601-1-8	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems Exclusion:6.3.3 * Auditory Alarm Signals
ANSI/AAMI/IEC 60601-2-19	Medical electrical equipment - Part 2-19: Requirements for the basic safety and essential performance of infant incubators Exclusion: 202 Electromagnetic disturbances – Requirements and tests
ANSI/AAMI/IEC 60601-2-21	Medical electrical equipment - Part 2-21: Requirements for the basic safety and essential performance of infant radiant warmers Exclusion: 202 Electromagnetic disturbances – Requirements and tests
ANSI/AAMI/IEC 60601-2-25	Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests

ANSI/AAMI/IEC 60601-2-27	Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment 202 Electromagnetic compatibility – Requirements and tests
ANSI/AAMI/IEC 60601-2-47	Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems Exclusion: 202 Electromagnetic compatibility – Requirements and tests
ANSI/AAMI/IEC 60601-2-50	Medical electrical equipment - Part 2-50: Particular requirements for the basic safety and essential performance of infant phototherapy equipment Exclusion: 202 Electromagnetic disturbances – Requirements and tests
ANSI/AAMI/IEC 62304	Medical device software - Software life cycle processes
ANSI/AAMI/IEC 80601-2-30	Medical electrical equipment - Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers Exclusion: 201.15.3.5.101 * Shock and vibration (robustness) 202 Electromagnetic disturbances – Requirements and tests
ANSI/AAMI/IEC MP80601-2-49	Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitors Exclusion: 202.7 Electromagnetic Emissions requirements for ME Equipment and ME Systems
CAN/CSA-C22.2 No. 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance Exclusion: 10.1.1, 10.1.2, 10.2, 10.3
CAN/CSA-C22.2 No. 60601-2-19	Medical electrical equipment - Part 2-19: Requirements for the basic safety and essential performance of infant incubators Exclusion: 202 Electromagnetic disturbances – Requirements and tests
CAN/CSA-C22.2 No. 60601-2-21	Medical electrical equipment - Part 2-21: Requirements for the basic safety and essential performance of infant radiant warmers Exclusion: 202 Electromagnetic disturbances – Requirements and tests
CAN/CSA-C22.2 No. 60601-2-25	Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests

CAN/CSA-C22.2 No. 60601-2-26	Medical electrical equipment - Part 2-26: Particular requirements for the basic safety and essential performance of electroencephalographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests
CAN/CSA-C22.2 No. 60601-2-27	Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment 202 Electromagnetic compatibility – Requirements and tests
CAN/CSA-C22.2 No. 60601-2-47	Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems Exclusion: 202 Electromagnetic compatibility – Requirements and tests
CAN/CSA-C22.2 No. 60601-2-49	Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitoring equipment Exclusion: 202.7 Electromagnetic Emissions requirements for ME Equipment and ME Systems
CAN/CSA-C22.2 No. 60601-2-50	Medical electrical equipment - Part 2-50: Particular requirements for the basic safety and essential performance of infant phototherapy Exclusion: 202 Electromagnetic disturbances – Requirements and tests
CAN/CSA-C22.2 No. 62304	Medical device software - Software life cycle processes
CAN/CSA-C22.2 No. 80601-2-61	Medical electrical equipment - Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment Exclusion: 201.15.3.5.101.1 * Shock and vibration 202 Electromagnetic disturbances – Requirements and tests
CSA C22.2 No. 60601-1-6	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
CSA C22.2 No. 60601-1-8	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems Exclusion:6.3.3 * Auditory Alarm Signals
CSA C22.2 No. 60601-1-11	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment Exclusion: 10 Construction of ME Equipment
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 Exclusion: 202.6 Electromagnetic Compatibility
CSA C22.2 No. 80601-2-30	Medical electrical equipment - Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers Exclusion: 201.15.3.5.101 * Shock and vibration (robustness) 202 Electromagnetic disturbances – Requirements and tests

EN 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance Exclusion: 10.1.1, 10.1.2, 10.2, 10.3
EN 60601-1-6	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
EN 60601-1-8	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems Exclusion:6.3.3 * Auditory Alarm Signals
EN 60601-1-11	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment. Exclusion: 10 Construction of ME Equipment
EN 60601-2-19	Medical electrical equipment - Part 2-19: Requirements for the basic safety and essential performance of infant incubators Exclusion: 202 Electromagnetic disturbances – Requirements and tests
EN 60601-2-21	Medical electrical equipment - Part 2-21: Requirements for the basic safety and essential performance of infant radiant warmers Exclusion: 202 Electromagnetic disturbances – Requirements and tests
EN 60601-2-25	Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests
EN 60601-2-27	Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment 202 Electromagnetic compatibility – Requirements and tests
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 Exclusion: 202.6 Electromagnetic Compatibility
EN 60601-2-47	Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems Exclusion: 202 Electromagnetic compatibility – Requirements and tests
EN 60601-2-50	Medical electrical equipment - Part 2-50: Particular requirements for the basic safety and essential performance of infant phototherapy equipment Exclusion: 202 Electromagnetic disturbances – Requirements and tests
EN 62304	Medical device software - Software life cycle processes
EN 80601-2-26 / EN 60601-2-26	Medical electrical equipment - Part 2-26: Particular requirements for the basic safety and essential performance of electroencephalographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests

EN 80601-2-49 / EN 60601-2-49	Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitors Exclusion: 202.7 Electromagnetic Emissions requirements for ME Equipment and ME Systems
EN IEC 80601-2-30	Medical electrical equipment - Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers Exclusion: 201.15.3.5.101 * Shock and vibration (robustness) 202 Electromagnetic disturbances – Requirements and tests
EN ISO 80601-2-61	Medical electrical equipment - Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment Exclusion: 201.15.3.5.101.1 * Shock and vibration 202 Electromagnetic disturbances – Requirements and tests
IEC 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance Exclusion: 10.1.1, 10.1.2, 10.2, 10.3
IEC 60601-1-6	Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability
IEC 60601-1-8	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems Exclusion: 6.3.3 * Auditory Alarm Signals
IEC 60601-1-11	Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment. Exclusion: 10 Construction of ME Equipment
IEC 60601-2-19	Medical electrical equipment - Part 2-19: Requirements for the basic safety and essential performance of infant incubators Exclusion: 202 Electromagnetic disturbances – Requirements and tests
IEC 60601-2-21	Medical electrical equipment - Part 2-21: Requirements for the basic safety and essential performance of infant radiant warmers Exclusion: 202 Electromagnetic disturbances – Requirements and tests
IEC 60601-2-25	Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests
IEC 60601-2-27	Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment 202 Electromagnetic compatibility – Requirements and tests

IEC 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 Exclusion: 202.6 Electromagnetic Compatibility
IEC 60601-2-47	Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems Exclusion: 202 Electromagnetic compatibility – Requirements and tests
IEC 60601-2-50	Medical electrical equipment - Part 2-50: Particular requirements for the basic safety and essential performance of infant phototherapy equipment Exclusion: 202 Electromagnetic disturbances – Requirements and tests
IEC 62304	Medical device software - Software life cycle processes
IEC 80601-2-26 / IEC 60601-2-26	Medical electrical equipment - Part 2-26: Particular requirements for the basic safety and essential performance of electroencephalographs Exclusion: 202 Electromagnetic disturbances – Requirements and tests
IEC 80601-2-30	Medical electrical equipment - Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers Exclusion: 201.15.3.5.101 * Shock and vibration (robustness) 202 Electromagnetic disturbances – Requirements and tests
IEC 80601-2-49 / IEC 60601-2-49	Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitors Exclusion: 202.7 Electromagnetic Emissions requirements for ME Equipment and ME Systems
IS 13450-1	Medical electrical equipment, part 1: general requirements for safety Exclusion: 10.1.1, 10.1.2, 10.2, 10.3
ISO 80601-2-61	Medical electrical equipment - Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment Exclusion: 201.15.3.5.101.1 * Shock and vibration 202 Electromagnetic disturbances – Requirements and tests