



SCOPE OF ACCREDITATION TO ISO/IEC 17025:2017

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ELECTRICAL

Valid to: February 28, 2025

Certificate Number: 3331.02

In recognition of the successful completion of the A2LA evaluation process (including an assessment of the organization’s compliance with A2LA’s FDA ASCA Accreditation Program ⁴ requirements), accreditation is granted to this laboratory at the location listed above, *as well as the satellite laboratory location listed below*, to perform the following product safety, radio, telecommunications, and electromagnetic compatibility (EMC) tests:

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
WiFi	Wi-Fi Alliance 802.11 with WPA2, WPA, and WEP System Interoperability Test Plan with ASD Test Engine for IEEE 802.11a, b, & g Devices; Wi-Fi Alliance Wi-Fi 802.11n System Interoperability Test Plan; Wi-Fi Alliance WMM System Interoperability Test Plan; Wi-Fi Alliance WMM Power Save System Interoperability Test Plan; Wi-Fi Alliance Wi-Fi WPS Test Plan; Wi-Fi Alliance Protected Management Frames; Wi-Fi Alliance Miracast; Wi-Fi Alliance Tunnel Direct Link Setup; Wi-Fi Alliance Passpoint; Wi-Fi Alliance Wi-Fi Direct; Wi-Fi Alliance 802.11 a/b/g interoperability, including WPA/WPA2-PSK, WPA/WPA2-Enterprise, and WEP security; Wi-Fi Alliance Wi-Fi Multimedia “WMM” interoperability; Wi-Fi Alliance 802.11n, including legacy a/b/g and WMM interoperability; Wi-Fi Alliance 802.11ac Interoperability; Wi-Fi Alliance Wi-Fi Protected Setup “WPS” version 1.0; Wi-Fi Alliance Wi-Fi Protected Setup “WPS” version 2.0; Wi-Fi Alliance Protected Management Frames; Wi-Fi Alliance Passpoint “Hotspot2.0” release 1; Wi-Fi Alliance Passpoint “Hotspot2.0” release 2; Wi-Fi Alliance WMM-Powersave; Wi-Fi Alliance WMM-Admission Control; Wi-Fi Alliance Voice Enterprise; Wi-Fi Alliance Wi-Fi Direct “P2P”; Wi-Fi Alliance Wi-Fi Display “Miracast” interoperability;

<u>Test Description:</u>	<u>Test Method(s)^{2,3:}</u>
WiFi (cont.)	Wi-Fi Alliance Wi-Fi Direct Services; Wi-Fi Alliance Tunneled Direct Link Setup; Wi-Fi Alliance Certified 6
<i>Product Safety</i>	
Information Technology	EN/IEC/CSA/UL 60950-1; EN/IEC/CSA/UL 60950-21; EN/IEC/CSA/UL 60950-23; EN/IEC 62040; IEC/UL 62368-1
Audio/Video	EN/IEC/CSA/UL 60065
Lab, Test & Measurement	IEC/EN/CSA/UL 61010-1; IEC/EN/CSA/UL 61010-2-010; IEC/EN/CSA/UL 61010-2-020; IEC/EN/CSA/UL 61010-2-081; IEC/EN/CSA/UL 61010-2-101
Laser	IEC/EN 60825-1
Household	EN/IEC/CSA/UL 60335-1; General Requirements; EN/IEC/CSA 60335-2-2; Vacuum Cleaners and Water Suction Cleaning; EN/IEC/CSA/UL 60335-2-8; Shavers, Hair Clippers; EN/IEC/CSA 60335-2-9; Grills, Toasters, and Similar Equipment; EN/IEC/CSA 60335-2-10; Floor Treatment and Wet Scrubbing Machines; EN/IEC/CSA 60335-2-14; Kitchen Appliances; EN/IEC/CSA 60335-2-15; Appliances for Heating Liquids; EN/IEC/CSA 60335-2-23; Appliances for Skin or Hair Care; EN/IEC/CSA 60335-2-28; Sewing Machines; EN/IEC/CSA 60335-2-29; Battery Chargers; EN/IEC/CSA 60335-2-30; Room Heaters; EN/IEC/CSA 60335-2-32; Massage Appliances; EN/IEC/CSA 60335-2-41; Pumps; EN/IEC/CSA 60335-2-42; Commercial Forced Convection Ovens, Steam Cookers and Steam-Convection Ovens; EN/IEC/CSA 60335-2-43; Clothes Dryers and Towel Rails; EN/IEC/CSA 60335-2-44; Ironers; EN/IEC/CSA 60335-2-45; Portable Heating Tools and Similar Equipment; EN/IEC/CSA 60335-2-52; Oral Hygiene Appliances; EN/IEC/CSA 60335-2-60; Whirlpool Baths; EN/IEC/CSA 60335-2-64; Commercial Electric Kitchen Machines; EN/IEC/CSA 60335-2-65; Air Cleaning Appliances; EN/IEC/CSA 60335-2-75; Commercial Dispensing Appliances and Vending; EN/IEC/CSA 60335-2-80; Fans; EN/IEC/CSA 60335-2-81; Foot Warmers and Heating Mats; EN/IEC/CSA 60335-2-82; Amusement Machines and Personal Service Machines; EN/IEC/CSA 60335-2-84; Toilets; EN/IEC/CSA 60335-2-95; Drives for Vertically Moving Garage Doors; EN/IEC/CSA 60335-2-97; Drives for Rolling Shutters, Awnings, Blinds;

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
Medical ³	IEC 60601-1-2 ⁴ ; KS C IEC 60601-1-2:2007; EN 60601-1-2; EN 60601-2-2; IEC 60601-1 (Ed. 2); IEC 60601-1 (Ed. 3); IEC 60601-1 (Ed 3.1); IEC 60601-1 (Ed 3.2); IEC 60601-1; ANSI/AAMI 60601-1; CAN/CSA C22.2 No. 60601-1; IEC 60601-1-1 (Medical electrical systems); IEC 60601-1-6 (Usability); IEC 60601-1-8 ⁴ ; IEC 60601-1-11 ⁴ (Home Healthcare); IEC 60601-2-2 ⁴ (High frequency surgical equipment); IEC 60601-2-4 (Cardiac defibrillators); IEC 60601-2-5 ⁴ (Ultrasonic physiotherapy equipment); IEC 60601-2-6 ⁴ (Microwave therapy equipment); IEC 60601-2-8 ⁴ ; IEC 60601-2-10 ⁴ (Nerve and muscle stimulators); IEC 60601-2-11 ⁴ (Gamma beam therapy equipment); IEC 60601-2-12 (Lung ventilators; Critical care ventilators); IEC 60601-2-13 (Anaesthetic systems); IEC 60601-2-16 ⁴ (Haemodialysis, haemodiafiltration, and haemofiltration equipment); IEC 60601-2-17 ⁴ (Automatically controlled brachytherapy after-loading equipment); IEC 60601-2-18 ⁴ (Endoscopic equipment); IEC 60601-2-19 ⁴ (Baby incubators); IEC 60601-2-20 ⁴ (Transport incubators); IEC 60601-2-21 ⁴ (Infant radiant warmers); IEC 60601-2-22 ⁴ (Diagnostic and therapeutic laser equipment); IEC 60601-2-23 ⁴ (Transcutaneous partial pressure monitoring equipment); IEC 60601-2-24 (Infusion pumps and controllers); IEC 60601-2-25 ⁴ (Electrocardiographs); IEC 60601-2-26 (Electroencephalographs); IEC 60601-2-27 ⁴ (Electrocardiographic monitoring equipment); IEC 60601-2-28 ⁴ (X-ray source assemblies and X-ray tube assemblies for medical diagnosis); IEC 60601-2-30 (Automatic cycling non-invasive blood pressure monitoring equipment); IEC 60601-2-34 ⁴ (Direct blood pressure monitoring equipment); IEC 60601-2-35 (Blankets, pads and mattresses intended for heating); IEC 60601-2-36 ⁴ (Equipment for extra-corporeally induced lithotripsy); IEC 60601-2-37 ⁴ (Ultrasonic medical diagnostic and monitoring equipment); IEC 60601-2-38 (Electrically operated hospital beds); IEC 60601-2-39 (Peritoneal dialysis equipment); IEC 60601-2-40 (Electromyographs and evoked response equipment); IEC 60601-2-41 (Surgical luminaires and luminaires for diagnosis); IEC 60601-2-43 ⁴ (X-ray equipment for interventional procedures); IEC 60601-2-44 ⁴ (X-ray equipment for computed tomography); IEC 60601-2-45 ⁴ (Mammographic x-ray equipment and mammographic stereotactic devices); IEC 60601-2-46 (Operating tables);



<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Product Safety (cont.)</i>	
Medical ³ (cont.)	IEC 60601-2-47 ⁴ (Ambulatory electrocardiographic systems); IEC 60601-2-49 (Multifunction patient monitoring equipment); IEC 60601-2-50 ⁴ (Infant phototherapy equipment); IEC 60601-2-51 (Recording and analysing single channel and multichannel electrocardiographs); IEC 60601-2-52 ⁴ (Medical Beds); IEC 60601-2-54 ⁴ (Particular requirements for the basic safety and essential performance of X-ray equipment for radiography and radioscopy); IEC 60601-2-57 (Non-laser Light Source eq.); IEC 60601-2-62 ⁴ (High intensity therapeutic ultrasound (HITU) equipment); IEC 60601-2-63 ⁴ (Dental extra-oral x-ray equipment); IEC 60601-2-64 ⁴ (Light ion beam medical electrical equipment); IEC 60601-2-65 ⁴ (Dental intra-oral x-ray equipment); IEC 60601-2-66 (Particular requirements for the basic safety and essential performance of hearing instruments and hearing instrument systems); IEC 60601-2-68 ⁴ (X-ray-based image-guided radiotherapy equipment for use with electron accelerators, light ion beam therapy equipment and radionuclide beam therapy equipment); IEC 60601-2-83 (Home Light Therapy Equipment); IEC 80601-2-30 ⁴ (Automated Non-invasive BP); IEC 80601-2-49 (Multifunction Patient Monitoring Equipment); IEC 80601-2-58 (Vitrectomy); IEC 80601-2-61; ISO 80601-2-61 ⁴ ; IEC 80601-2-77
Lighting	IEC 60598-1:2003; IEC / EN 60598-1; IEC 60598-1:2003/AMD1:2006; IEC 60598-1:2008; IEC 60598-1:2014; IEC 60598-1:2014/AMD1:2017; IEC / EN 60598-2-1; IEC 60598-2-1:1979/AMD1:1987; IEC / EN 60598-2-2; IEC 60598-2-2:2011; IEC / EN 60598-2-3; IEC 60598-2-3:2002/AMD1:2011; IEC / EN 60598-2-4; IEC 60598-2-4:1997; IEC 60598-2-4:2017; IEC / EN 60598-2-5; IEC 60598-2-5:1998; IEC 60598-2-5:2015; IEC / EN 60598-2-7; IEC 60598-2-7:1982/AMD1:1987/AMD2:1994; IEC / EN 60598-2-17; IEC 60598-2-17:1984/AMD1:1987/AMD2:1990;



<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Product Safety (cont.)</i>	
Lighting (cont.)	IEC / EN 60598-2-23; IEC 60598-2-23:1996/AMD1:2000

The below tests are performed using the above Product Safety standards:

- Input current / Power Input
- Durability of Markings
- Access to Live Parts
- Energy Hazards
- Capacitance Discharge
- TNV Circuits, Limits, Connections
- Voltages Generated Externally
- SELV Circuits
- Torque
- Telecommunication Network
- Separation and Protection
- Limited Current Circuits, Values
- Limited Power Sources
- Resistances of Earthing Conductors
- GND Continuity Test
- Humidity Conditioning
- Creepage Distances, Clearances
- Working Voltage
- Thermal Cycling and Thermal Aging
- Mechanical Strength / Impact
- Enclosed and Sealed Parts
- Steady Force Test
- Drop Test
- Stress Relief
- Wall or Ceiling Mounted Equipment
- Handles and Manual Controls
- Battery Overcharge/Discharge and Reverse Current
- Measurements
- Spillage Tests
- Protection against Hazardous Moving Parts
- Thermal Requirements / Ball Pressure Test
- Temperature Rise
- Resistance to Abnormal Heat
- Touch Current and Protective Conductor Current / Leakage
- Dielectric Strength / Hipot
- Component Failure and Abnormal Operation
- Power Supply Output/transformer/accessible Connector Overload
- Voltage Surge / Impulse
- Stability
- Sound Pressure Level
- Resistance to Fire

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
Energy Efficiency	IEC/EN 62301; CAN/CSA C381.1-08; CAN/CSA C802.2

EPA ENERGY STAR Testing

<u>Product Family Guidelines:</u>	<u>Supporting Test Method(s) ²:</u>
Computers	ENERGY STAR Program Requirements for Computers, Version 8.0; IEC 62301; EPRI Generalized Test Protocol for Calculating the Energy Efficiency of Internal A.C.-D.C. and D.C.-D.C. Power Supplies



<u>Product Family Guidelines:</u>	<u>Supporting Test Method(s) ²:</u>
Enterprise Servers	ENERGY STAR Program Requirements for Computer Servers, Version 3.0; IEC 62301; EPRI Generalized Test Protocol for Calculating the Energy Efficiency of Internal AC-DC and DC-DC Power Supplies
Data Center Storage	ENERGY STAR Program Requirements for Data Center Storage, Version 2.1; ENERGY STAR Test Method for Data Center Storage Equipment, Rev. Mar 2014
Imaging Equipment	ENERGY STAR Program Requirements for Imaging Equipment, Version 3.2; IEC 62301
Uninterruptable Power Supplies	ENERGY STAR Program Requirements for Uninterruptable Power Supplies, Version 2.0
Displays	ENERGY STAR Program Requirements for Displays, Version 8.0
Televisions	ENERGY STAR Program Requirements for Televisions, Version 8.0
Audio/Video Equipment	ENERGY STAR Program Requirements for Audio/Video Equipment, Version 3.0

The below tests are performed using the above EPA Energy Star methods:

- Voltage
- Luminance
- Current

¹ *This accreditation covers testing performed at the main laboratory listed above, and at the satellite laboratory indicated below:*

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<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Emissions</i>	
Radiated & Conducted <i>(10m semi-anechoic chamber, up to 220 GHz)</i>	47 CFR, FCC Part 15B (using ANSI C63.4:2014); ANSI C63.4:2009; 47 CFR, FCC Part 18 (using MP-5:1986); IEC/CISPR 11; EN 55011; KS C 9811:2019; EN/IEC 55012; CISPR 12; EC/EN 55013; CISPR 13; CNS 13439 <i>(only associated equipment without antenna connection)</i> ; CISPR 14-1; IEC/EN 55014-1; KS C 9814-1:2020; CISPR 15; IEC/EN 55015; KS C 9815:2019;



<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Emissions (cont.)</i>	
Radiated & Conducted (10m semi-anechoic chamber, up to 220 GHz) (cont.)	IEC/CISPR 22; EN 55022; AS/NZS CISPR 22:2009 + A1:2010; EN 55032; CISPR 32; KS C 9832:2019; AS/NZS CISPR 32; ICES-001; ICES-003, Issue 6; ICES-003, Issue 7; VCCI-CISPR 32:2016 (up to 6 GHz); CNS 13803; CNS 13783-1; CNS 13438 (up to 6 GHz); CNS 13439; QCVN 96:2015/BTTTT; QCVN 118:2018/BTTTT; JIS C 1806-1; SANS 222
Radiated & Conducted (3m semi-anechoic chamber, up to 220 GHz)	47 CFR, FCC Part 15B (using ANSI C63.4:2014); ANSI C63.4:2014; 47 CFR, FCC Part 18 (using MP-5:1986); IEC/CISPR 11; EN 55011; KS C 9811:2019; EN/IEC 55012; CISPR 12; EC/EN 55013; CISPR 13; CNS 13439 (only associated equipment without antenna connection); CISPR 14-1; IEC/EN 55014-1; KS C 9814-1:2020; CISPR 15; IEC/EN 55015; KS C 9815:2019; IEC/CISPR 22; EN 55022; AS/NZS CISPR 22:2009 + A1:2010; EN 55032; CISPR 32; AS/NZS CISPR 32; ICES-001; ICES-003, Issue 6; ICES-003, Issue 7; VCCI-CISPR 32:2016 (up to 6 GHz); CNS 13803; CNS 13783-1; CNS 13438 (up to 6 GHz); CNS 13439; QCVN 118:2018/BTTTT; JIS C 1806-1
Current Harmonics	IEC/EN 61000-3-2; SANS 61000-3-2
Flicker and Fluctuations	IEC/EN 61000-3-3; SANS 61000-3-3
<i>Immunity</i>	
Electrostatic Discharge	IEC 61000-4-2; EN 61000-4-2; SANS 61000-4-2
Radiated (20V/m 80% AM modulated @ 6 GHz)	IEC 61000-4-3; EN 61000-4-3; SANS 61000-4-3
Electrical Fast Transient / Burst	IEC 61000-4-4; EN 61000-4-4; SANS 61000-4-4
Surge	IEC 61000-4-5; EN 61000-4-5; SANS 61000-4-5

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Immunity (cont.)</i>	
Conducted	IEC 61000-4-6; EN 61000-4-6; SANS 61000-4-6
Power Frequency Magnetic Field	IEC 61000-4-8; EN 61000-4-8; SANS 61000-4-8
Voltage Dips, Short Interrupts, and Voltage Variations	IEC 61000-4-11; EN 61000-4-11; SANS 61000-4-11
Ring Wave	IEC 61000-4-12; SANS 61000-4-12
<i>Generic / Product Specific EMC Standards</i>	<p>EN/IEC 61000-6-1; EN/IEC 61000-6-2; EN/IEC 61000-6-3; EN/IEC 61000-6-4; KS C 9610-6-1:2019; KS C 9610-6-2:2019; KS C 9610-6-3:2017; KS C 9610-6-4:2017; IEC/EN 61204-3; EN/IEC 60601-1-2; KS C IEC 60601-1-2:2007; EN/IEC 61547; KS C 9547:2020; ISO 11451-4; EN/IEC 12895; EN/IEC 13309; EN 12015; EN 12016; EN/ISO 13766; EN/ISO 14982; EN 50121-3-2; EN 50121-2; EN 50121-3-1; EN 50121-4; EN 62233; EN 55103-1; EN 55103-2; EN/IEC 61326-1; EN/IEC 61326-2-6; EN/IEC 61326-3-2; EN/IEC 61800-3; KS C 9800-3:2017; CISPR 24; EN 55024; CISPR 35: EN 55035; EN 50121-1; EN 50130-4; EN 55103-2; EN 50121-4; EN 50121-3-2; EN/IEC 50155; EN 50270; EN 50293; EN/IEC 55014-2; IEC/CISPR 14-2; KS C 9814-2:2020; EN 50370-1; EN 50370-2; EN 50361; EN 50364; EN 50371; KS C 9815:2019; KS C 9835: 2019 (up to 12V/m, <i>excluding broadcast and TV receivers</i>); ETSI EN 301 489-1; ETSI EN 301 489-3; ETSI EN 301 489-4; ETSI EN 301 489-5; ETSI EN 301 489-6; ETSI EN 301 489-7; ETSI EN 301 489-8; ETSI EN 301 489-9; ETSI EN 301 489-10; ETSI EN 301 489-12; ETSI EN 301 489-15; ETSI EN 301 489-16; ETSI EN 301 489-17; ETSI EN 301 489-18; ETSI EN 301 489-19; ETSI EN 301 489-20; ETSI EN 301 489-23; ETSI EN 301 489-24; ETSI EN 301 489-25; ETSI EN 301 489-26; ETSI EN 300 386 V1.5.1/ V1.6.1; KS X 3124:2020; KS X 3125:2020; KS X 3126:2020; AIM Standard 7351731; AAMI TIR69; ANSI C63.27:2017</p>

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
<i>Radio Communications</i> <i>(up to 220 GHz, excluding HAC)</i>	
Australia / New Zealand	AS/NZS 4268:2017 ACMA Radiocommunications (Short Range Devices) Standard 2014
Unlicensed Radio - FCC	CFR 47, FCC Part 2; 47 CFR, FCC Part 15, Subpart C (using ANSI C63.10:2013); 47 CFR, FCC Part 15, Subpart D (using ANSI C63.17:2013); 47 CFR, FCC Part 15, Subpart E (using ANSI C63.10:2013 and FCC KDB 905462 D02 (v02)); ANSI C63.10:2013
Licensed Radio - FCC	47 CFR, FCC Part 2; 47 CFR, FCC Parts 22, 24, 25, 27, 74, 80, 87, 90, 95, 97, 101 (using ANSI/TIA-603-E)
Canada	RSS-102 Measurement (RF Exposure); RSS-102 Measurement (NS); SPR-002; Radio Scope 1 RSS-Gen; RSS-102; RSS-210; RSS-211; RSS-213; RSS-215; RSS-216; RSS-220; RSS-222; RSS-236; RSS-238; RSS-243; RSS-244; RSS-246; RSS-247; RSS-248; RSS-251; RSS-252; RSS-287; RSS-288; RSS 310; Radio Scope 2 RSS-Gen; RSS-102; RSS-112; RSS-130; RSS-132; RSS-133; RSS-134; RSS-139; RSS-170; Radio Scope 3 RSS-Gen; RSS-102; RSS-111; RSS-119; RSS-123; RSS-125; RSS-127; RSS-131; RSS-135; RSS-137; RSS-140; RSS-197; RSS-199; Radio Scope 4 RSS-Gen; RSS-102; RSS-117; RSS-141; RSS-181; RSS-182; Radio Scope 5 RSS-Gen; RSS-102; RSS-142; RSS-191; RSS-192; RSS-194; RSS-195; RSS-196
Europe (EU)	ETSI EN 300 220-1; ETSI EN 300 220-2; ETSI EN 300 220-3; ETSI EN 300 220-4; ETSI EN 300 328; ETSI EN 300 330-1; ETSI EN 300 330-2; ETSI EN 300 440-1; ETSI EN 300 440-2; ETSI EN 302 208-1; ETSI EN 302 208-2; ETSI EN 300 113-1; ETSI EN 300 113-2; ETSI EN 300 133-1; ETSI EN 300 133-2; ETSI EN 300 422-2;

<u>Test Description:</u>	<u>Test Method(s) ^{2,3}:</u>
Europe (EU) (<i>cont.</i>)	ETSI EN 301 839-1; ETSI EN 301 839-2; ETSI EN 301 893; ETSI EN 302 502; ETSI EN 303 413
Hong Kong	HKCA 1039; HKCA 1041; HKCA 1042; HKCA 1049
Singapore	IMDA TS SRD; IMDA TS UWB
Taiwan	DGT LP0002 (2020); DGT IP0001 (2020)
Japan	ARIB Standard STD-T66; ARIB Standard STD-T67; ARIB Standard STD-T70; STD-T71, STD-T82, STD-T90, STD-T106, STD-T107, STD-T108
Vietnam	QCVN 54:2011/BTTTT; QCVN 55:2011/BTTTT; QCVN 118:2018/BTTTT
Cellular Wireless	EN 302 511; EN 301-908-1; EN 301-908-2; EN 301-908-13;

Republic of Korea Technical Regulations

Regulations on Radio Equipment (Enforcement decree MSIP No 78);
 Unlicensed Radio Equipment Established Without Notice (MSIT Public Notification 2020-59, Oct 16, 2020);
 Technical Requirements for Radio Equipment for Telecommunication Services
 (RRA Public Notification 2019-9, Jun 3, 2019))
 Conformity Assessment Procedure of Radio Equipment (RRA Announce 2015-135 KS X 3123);
 Technical Requirements for Measurement of Electromagnetic Field Strength
 (RRA Public Notification 2019-3, March 4, 2019)

² When the date, edition, version, etc. is not identified in the scope of accreditation, laboratories may use the version that immediately precedes the current version for a period of one year from the date of publication of the standard measurement method, per part C., Section 1 of A2LA R101 - *General Requirements- Accreditation of ISO-IEC 17025 Laboratories*.

³ The laboratory is only accredited for testing activities outlined within the test methods listed above. Reference to any other activity within these standards, such as risk management or risk assessment, does not fall within the laboratory's accredited capabilities.

Testing Activities performed under the scope of the U.S FDA ASCA Pilot Program Specifications: *Basic Safety and Essential Performance of Medical Electrical Equipment, Medical Electrical Systems, and Laboratory Medical Equipment – Standards Specific Information for the Accreditation Scheme for Conformity Assessment (ASCA) Pilot Program* published on September 25th, 2020, and in accordance with all requirements of A2LA R256 *Specific Requirements- FDA ASCA Program* ⁴:

Standards:

ANSI AAMI ES60601-1:2005/(R)2012 and A1:2012, C1:2009/(R)2012 and A2:2010/(R)2012
(Consolidated Text)

IEC 60601-1-2 Edition 4.1 2020-09 CONSOLIDATED VERSION

IEC 60601-1-8 Edition 2.2 2020-07 CONSOLIDATED VERSION

IEC 60601-1-11 Edition 2.1 2020-07 CONSOLIDATED VERSION

ANSI AAMI IEC 60601-2-2:2017

IEC 60601-2-2 Edition 6.0 2017-03

IEC 60601-2-5 Edition 3.0 2009-07

IEC 60601-2-6 Edition 2.1 2016-04

IEC 60601-2-8 Edition 2.1 b:2015

IEC 60601-2-10 Edition 2.1 2016-04

IEC 60601-2-11 Edition 3.0 2013-01

IEC 60601-2-16 Edition 5.0 2018-4

IEC 60601-2-17 Edition 3.0 2013-11

IEC 60601-2-18 Edition 3.0 2009-08

ANSI AAMI IEC 60601-2-19:2009/(R)2014 & A1:2016

IEC 60601-2-20 Edition 2.1 2016-04

IEC 60601-2-21 Edition 2.1 2016-04

IEC 60601-2-22 Edition 3.1 2012-10

IEC 60601-2-23 Edition 3.0 2011-02

IEC 60601-2-25 Edition 2.0 2011-10

IEC 60601-2-27 Edition 3.0 2011-03

IEC 60601-2-28 Edition 3.0 2017-06

IEC 60601-2-34 Edition 3.0 2011-05

IEC 60601-2-36 Edition 2.0 2014-04

IEC 60601-2-37 Edition 2.1 2015

IEC 60601-2-43 - Ed. 2.0 2010-03

IEC 60601-2-43 Edition 2.2 2019-10 CONSOLIDATED VERSION

IEC 60601-2-44 Edition 3.2 2016

IEC 60601-2-45 Edition 3.1 2015

IEC 60601-2-47 Edition 2.0 2012-02

IEC 60601-2-50 Edition 3.0 2020-09

IEC 60601-2-52 Edition 1.0 2009-12

IEC 60601-2-54 Edition 1.2 2018-06 CONSOLIDATED VERSION

IEC 60601-2-57 Edition 1.0 2011-01

IEC 60601-2-62 Edition 1.0 2013-07

IEC 60601-2-63 Edition 1.1 2017-07 CONSOLIDATED VERSION

IEC 60601-2-64 Edition 1.0 2014-09

IEC 60601-2-65 Edition 1.1 2017-05 CONSOLIDATED VERSION

IEC 60601-2-68 Edition 1.0 2014-09

IEC 80601-2-30 Edition 2.0 2018-03

ISO 80601-2-61 Second Edition 2017-12 (Corrected Version 2018-02)

⁴ These methods have been assessed by A2LA according to A2LA's FDA ASCA Program requirements. Accreditation by A2LA does not imply FDA ASCA-Accreditation. All ASCA-accreditation decisions for testing laboratory applications are made solely by the FDA, a list of approved laboratories can be found at FDA.gov.

Testing Activities Performed in Support of FCC Certification in Accordance with 47 Code of Federal Regulations and FCC KDB 974614, Appendix A, Table A.1 ⁵ :		
Rule Subpart/Technology	Test Method	Maximum Frequency
Unintentional Radiators		
Part 15B	ANSI C63.4:2014	220000 MHz
Industrial, Scientific, and Medical Equipment		
Part 18	FCC MP-5 (February 1986)	220000 MHz
Intentional Radiators		
Part 15C	ANSI C63.10:2013	220000 MHz
Unlicensed Personal Communication Systems Devices		
Part 15D	ANSI C63.17:2013	220000 MHz
U-NII without DFS Intentional Radiators		
Part 15E	ANSI C63.10:2013	220000 MHz
U-NII with DFS Intentional Radiators		
Part 15E	FCC KDB 905462 D02 (v02)	220000 MHz
Commercial Mobile Services (FCC Licensed Radio Service Equipment)		
Parts 22 (cellular), 24, 25 (below 3 GHz), and 27	ANSI/TIA-603-E; TIA-102.CAAA-E	220000 MHz
General Mobile Radio Services (FCC Licensed Radio Service Equipment)		
Parts 22 (non-cellular), 90 (below 3 GHz), 95, 97, and 101 (below 3 GHz)	ANSI/TIA-603-E; TIA-102.CAAA-E	220000 MHz

⁵Accreditation does not imply acceptance to the FCC equipment authorization program. Please see the FCC website (<https://apps.fcc.gov/oetcf/eas/>) for a listing of FCC approved laboratories.



Accredited Laboratory

A2LA has accredited

TUV RHEINLAND OF NORTH AMERICA, INC.

Pleasanton, CA

for technical competence in the field of

Electrical Testing

This laboratory is accredited in accordance with the recognized International Standard ISO/IEC 17025:2017 *General requirements for the competence of testing and calibration laboratories*. This laboratory also meets A2LA R256 - *Specific Requirements - FDA ASCA Program*. This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory quality management system (refer to joint ISO-ILAC-IAF Communiqué dated April 2017).



Presented this 31st day of May 2023.

A blue ink signature of Mr. Trace McInturff, written over a horizontal line.

Mr. Trace McInturff, Vice President, Accreditation Services
For the Accreditation Council
Certificate Number 3331.02
Valid to February 28, 2025

For the tests to which this accreditation applies, please refer to the laboratory's Electrical Scope of Accreditation.