SCOPE OF ACCREDITATION TO ISO/IEC 17025:2017

MEDICAL EQUIPMENT COMPLIANCE ASSOCIATES, LLC.
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ELECTRICAL

Valid to: January 31, 2025 Certificate Number: 3392.01

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 [1] requirements), accreditation is granted to this laboratory to perform the following Product Safety tests:

Test Technology:	**Test Method(s) [2, 3, 4]:**
Medical Electrical Equipment– Part 1: General Requirements for Safety	IEC 60601-1; EN 60601-1; UL 60601-1; CAN/CSA C22 No. 601.1-M90; GB 9706.1:2007 (*excluding clauses:* *29 X-Radiation;* *39 Category AP and APG Testing*)
Medical Electrical Equipment– Part 1: General Requirements for Basic Safety and Essential Performance	IEC 60601-1; EN 60601-1; ANSI/AAMI ES60601-1[1]; CAN/CSA C22.2 No. 60601-1; GB 9706-1:2020; JIS T 0601-1:2012 + A1:2014 (*excluding clauses:* *8.8.4.2 Oxygen Aging for Rubber Insulation Parts;* *9.6.3 Hand Transmitted Vibration;* *10.1 X-Radiation, Annex G [Protection against hazards of ignition of flammable anesthetic mixtures], Annex L [Insulated winding wires for use without interleaved insulation]*)
Safety Requirements for Medical Electrical Systems	IEC 60601-1-1; EN 60601-1-1; CAN/CSA C22.2 No. 60601-1-1

Test Technology:	**Test Method(s) [2, 3, 4]:**
Medical electrical equipment – Electromagnetic disturbances	General Requirements: IEC 60601-1-2:2014 + A1:2020 (excluding ISM, Powerline Harmonics, Voltage Fluctuation and Flicker, Radiated Immunity, Conducted Immunity, Radiated Field in Close Proximity); ANSI/AAMI ES 60601-1-2:2014 + A1:2021; EN 60601-1-2:2014 + A1:2021; CAN/CSA C22.2 No. 60601-1-2:2016; YY 9706.102:2021
Programmable Electrical Medical Systems	IEC 60601-1-4; EN 60601-1-4; CAN/CSA C22.2 No. 60601-1-4
Usability	IEC 60601-1-6[1]; EN 60601-1-6; CAN/CSA C22.2 No. 60601-1-6; CAN/CSA C22.2 No. 60601-1-6:2011; NBR IEC 60601-1-6:2011 + A1:2020; YY/T 9706.106:2021; IEC 62366; EN 62366; ANSI/AAMI/IEC 62366; CAN/CSA IEC 62366; IEC 62366-1; ANSI/AAMI/IEC 62366-1; CAN/CSA-IEC 62366-1; EN 62366-1; JIS T 62366-1:2019
Medical Electrical Equipment and Medical Electrical Systems used in the Home Healthcare Environment	IEC 60601-1-11[1]; IEC 60601-1-11: 2015[1]; ANSI AAMI HA60601-1-11[1]; EN 60601-1-11; EN 60601-1-11:2015; CAN/CSA C22.2 No. 60601-1-11; CAN/CSA C22.2 No. 60601-1-11:2015; NBR IEC 60601-1-11:2021; YY 9706.111:2021
Medical Electrical Equipment and Medical Electrical Systems used in the Emergency Medical Services Environment	IEC 60601-1-12[1]; IEC 60601-1-12[1]; ANSI/AAMI 60601-1-12:2016 + A1:2021[1]; IEC 60601-1-12:2016; CAN/CSA C22.2 No. 60601-1-12; EN 60601-1-12; YY 9706.112:2021

Test Technology:	**Test Method(s) [2, 3, 4]:**
Requirements/Guidelines for Alarms in Medical Electrical Equipment	IEC 60601-1-8[1]; IEC 60601-1-8:2012[1]; EN 60601-1-8; EN 60601-1-8:2007 + A1:2013; CAN/CSA C22 No. 60601-1-8; CAN/CSA C22 No. 60601-1-8:2008; ANSI/AAMI/IEC 60601-1-8; ANSI AAMI IEC 60601-1-8:2006 + A1:2012 + A2:2021[1]; YY 9706.108:2021
Requirements for Physiological Closed-Loop Controllers	IEC 60601-1-10[1]; IEC 60601-1-10: 2013[1]; EN 60601-1-10; CAN/CSA C22.2 No. 60601-1-10; YY 9706.110:2021
High Frequency Surgical Equipment	IEC 60601-2-2; IEC 60601-2-2 Edition 6.0 2017-03[1]; ANSI AAMI IEC 60601-2-2:2017[1]; IEC 60601-2-2:1998; IEC 60601-2-2:2009; ANSI/AAMI/IEC 60601-2-2:2009; EN 60601-2-2:2009 + A11:2011; EN 60601-2-2:2018; CAN/CSA C22.2 No. 60601-2-2:2008; CAN/CSA C22.2 No. 60601-2-2:2009; GB 9706.4:2009 (*excluding:* *Ed. 2, 3, 4 Clause 59.104.7;* *Ed. 5 Clause 201.15.101.7* *[Neutral Electrode Adhesion]*)
Cardiac Defibrillators, Defibrillator-Monitors	IEC 60601-2-4; IEC 60601-2-4:2002; IEC 60601-2-4:2010 + A1:2018; ANSI/AAMI/IEC 60601-2-4:2010 + A1:2018; EN 60601-2-4:2011; EN 60601-2-4:2003; CAN/CSA C22.2 No. 60601-2-4:2004; CAN/CSA C22.2 No. 60601-2-4:2012; GB 9706.8:2009
Nerve and Muscle Stimulators	IEC 60601-2-10[1]; IEC 60601-2-10:1987 + A1:2001; IEC 60601-2-10:2016; CSA C22.2 No. 601.2.10:1992; CAN/CSA C22.2 No. 60601-2-10:2014; EN 60601-2-10:2000 + A1:2001; EN 60601-2-10:2015 + A1:2016; YY 0607:2007; YY 9706.210:2021

Test Technology:	**Test Method(s) [2, 3, 4]:**
Hemodialysis Equipment	IEC 60601-2-16 [1]; ANSI AAMI IEC 60601-2-16:2018[1]; IEC 60601-2-16:1998; IEC 60601-2-16:2012; ANSI/AAMI/IEC 60601-2-16:2012; EN 60601-2-16:1998; EN 60601-2-16; CAN/CSA C22.2 No. 60601-2-16; CAN/CSA C22.2 No. 60601-2-16:2001; CAN/CSA C22.2 No. 60601-2-16:2009; CAN/CSA C22.2 No. 60601-2-16:2014; GB 9706.2:2003; GB 9706.216:2021
Endoscopic Equipment	IEC 60601-2-18[1]; IEC 60601-2-18:1996 + A1:2000; EN 60601-2-18; EN 60601-2-18:1996 + A1:2000; CAN/CSA C22.2 No. 60601-2-18; CAN/CSA C22.2 No. 60601-2-18:2001; GB 9706.19:2000
Infusion Pumps and Controllers	IEC 60601-2-24; IEC 60601-2-24:1998; EN 60601-2-24; EN 60601-2-24:1998; CAN/CSA C22.2 No. 60601-2-24; CAN/CSA C22.2 No. 60601-2-24:2001; GB 9706.24:2005
Electrocardiographs	IEC 60601-2-25[1]; ANSI AAMI IEC 60601-2-25[1]; IEC 60601-2-25:1993 + A1:1999; EN 60601-2-25; EN 60601-2-25:1995 + A1:1999; CAN/CSA C22.2 No. 601.2.25; CAN/CSA C22.2 No. 60601-2-25:1994
Electroencephalographs	IEC 60601-2-26; IEC 60601-2-26:2002; IEC 80601-2-26; EN 60601-2-26; EN 60601-2-26:2003; CAN/CSA C22.2 No. 60601-2-26; CAN/CSA C22.2 No. 60601-2-26:04; GB 9706.26:2005; GB 9706.226:2021

Test Technology:	**Test Method(s) [2, 3, 4]:**
Electrocardiographic Monitoring Equipment	IEC 60601-2-27[1]; ANSI AAMI IEC 60601-2-27[1]; EN 60601-2-27; GB 9706.227:2021; IEC 60601-2-27:1994; EN 60601-2-27; EN 60601-2-27:1994; CAN/CSA C22.2 No. 60601-2-27; GB 9706.25:2005
Automated Non-Invasive Sphygmomanometers	IEC 60601-2-30; CAN/CSA C22.2 No. 60601-2-30; EN 60601-2-30; YY 0667:2008; IEC 80601-2-30[1]; IEC 80601-2-30 Edition 1.1 2013-07[1]; ANSI AAMI IEC 80601-2-30:2009 & A1:2013 (R2016)[1] ANSI AAMI IEC 80601-2-30:2018[1]; EN ISO 80601-2-30; CAN/CSA C22.2 No. 80601-2-30
Invasive Blood Pressure Monitoring Equipment	IEC 60601-2-34; IEC 60601-2-34:2000; EEN 60601-2-34; EN 60601-2-34:2004; CAN/CSA C22.2 No. 60601-2-34; CAN/CSA C22.2 No. 60601-2-34:2002; YY 0783:2010
Heating Devices Using Blankets, Pads, and Mattresses	IEC 60601-2-35; IEC 60601-2-35:1996; EN 60601-2-35; EN 60601-2-35:1996; YY 0834:2011; IEC 80601-2-35[1]; ANSI AAMI IEC 80601-2-35; EN 80601-2-35; CAN/CSA C22.2 No. 80601-2-35; (*excluding: Ed.1 Clause 59.2.101 [Spark Ignition Test]*)
Extracorporeally Induced Lithotripsy	IEC 60601-2-36[1]; IEC 60601-2-36:1997; EN 60601-2-36; EN 60601-2-36:1997; CAN/CSA C22.2 No. 60601-2-36; CAN/CSA C22.2 No. 60601-2-36:1998; GB 9706.22:2003
Ultrasonic Monitoring and Diagnostic Equipment	IEC 60601-2-37 [1]; CSA C22.2 No. 60601-2-37; EN 60601-2-37; GB 9706.237:2020

Test Technology:	**Test Method(s) [2, 3, 4]:**
Electrically Operated Hospital Beds	IEC 60601-2-38; EN 60601-2-38; CAN/CSA C22.2 No. 60601-2-38
Peritoneal Dialysis Equipment	IEC 60601-2-39; IEC 60601-2-39:1999; EN 60601-2-39; EN 60601-2-39:1999; CAN/CSA C22.2 No. 60601-2-39; CAN/CSA C22.2 No. 60601-2-39:2002; GB 9706.239:2021; GB 9706.36:2008; JIS T 0601-2-39:2013
Electromyographs and Evoked Response Equipment	IEC 60601-2-40; IEC 60601-2-40:1998; EN 60601-2-40; EN 60601-2-40:1998; CAN/CSA C22.2 No. 60601-2-40; CAN/CSA C22.2 No. 60601-2-40:2001
Operating Tables	IEC 60601-2-46; IEC 60601-2-46:1998; EN 60601-2-46; EN 60601-2-46:1998; CAN/CSA C22.2 No. 60601-2-46; CAN/CSA C22.2 No. 60601-2-46:2001; YY 0570:2013
Ambulatory Electrocardiographic Monitors	IEC 60601-2-47[1]; ANSI AAMI IEC 60601-2-47[1]; IEC 60601-2-47:2001; EN 60601-2-47; EN 60601-2-47:2001; CAN/CSA C22.2 No. 60601-2-47; CAN/CSA C22.2 No. 60601-2-47:2003; YY 0885:2013
Multiparameter Patient Monitoring Equipment	IEC 60601-2-49; IEC 60601-2-49:2001; EN 60601-2-49; EN 60601-2-49:2001; CAN/CSA C22.2 No. 60601-2-49; CAN/CSA C22.2 No. 60601-2-49:2004; YY 0668:2008; IEC 80601-2-49; EN IEC 80601-2-49

Test Technology:	**Test Method(s) [2, 3, 4]:**
Recording and Analyzing Single and Multichannel Electrocardiographs	IEC 60601-2-51; EN 60601-2-51; CAN/CSA C22.2 No. 60601-2-51; YY 0782:2010 (*excluding clause:* *50.102 Automated ECG interpretation for analyzing electrocardiographs*)
Safety of Medical Beds	IEC 60601-2-52[1]; EN 60601-2-52; EN 60601-2-52:2010 + AC:2011 + A1:2015; CAN/CSA C22.2 No. 60601-2-52
Respiratory Gas Monitors	ISO 80601-2-55[1]; ISO 80601-2-55:2011; EN ISO 80601-2-55; EN 80601-2-55:2011; CAN/CSA C22.2 No. 80601-2-55
Clinical Thermometers for Body Temperature Measurement	ISO 80601-2-56[1]; ISO 80601-2-56:2012[1]; CAN/CSA C22.2 No. 80601-2-56; EN ISO 80601-2-56; EN ISO 80601-2-56:2012
Dental Equipment	ISO 80601-2-60; ISO 80601-2-60:2012; CAN/CSA C22.2 No. 80601-2-60; EN 80601-2-60; GB 9706.260:2020 (*excluding clauses:* *201.10 Protection against excessive radiation hazards*)
Pulse Oximeter Equipment	ISO 80601-2-61[1]; ISO 80601-2-61:2011; EN ISO 80601-2-61; EN ISO 80601-2-61:2011; CAN/CSA C22.2 No. 80601-2-61
Pulse Oximeter Equipment for Medical Use	ISO 9919; CAN/CSA Z9919; YY 0784:2010 (*excluding clause:* *50.101.2 Clinical Determination of SpO2 Accuracy*)

Test Technology:	**Test Method(s) [2, 3, 4]:**
Medical Vehicles and their Equipment	EN 1789; EN 1789:2007 + A1:2010 + A2:2014 (only Clause 6 [Medical Devices]) (*excluding clauses:* *6.1 Provision of medical devices;* *6.2 Medical device storage;* *6.3.5 Fixation of medical devices;* *6.3.8 Gas supply;* *6.5 List of equipment*)
Degrees of Protection Provided by Enclosures	IEC 60529; IEC 60529:1998 + A1:1999 + A2:2013; EN 60529; EN 60529:1991 + A1:2000 + A2:2013 + AC:2016; CAN/CSA C22.2 No. 60529; CAN/CSA C22.2 No. 60529:2005
Software Life-Cycle Processes	IEC 62304; IEC 62304:2006; EN 62304; EN 62304:2006; ANSI/AAMI/IEC 62304; ANSI/AAMI/IEC 62304:2006; CAN/CSA CEI/IEC 62304; YY/T 0664:2020; YY/T 0664:2008
Safety Requirements for Electrical Equipment for Measurement, Control, and Laboratory use – Part 1: General Requirements	IEC 61010-1[1]; IEC 61010-1:2001; IEC 61010-1:2010 + A1:2016 + COR1:2019 (*excluding subclause 16.2 category III & IV*); EN 61010-1; EN 61010-1:2001; CAN/CSA C22.2 No. 61010-1; CAN/CSA C22.2 No. 61010-1:2004; UL 61010-1; UL 61010-1:2004 (*excluding subclause 16.2 category III & IV*)
Equipment for Heating of Materials	IEC 61010-2-010[1]; IEC 61010-2-010:2003; IEC 61010-2-010:2014; EN 61010-2-010; EN 61010-2-010:2003; EN 61010-2-010:2014; CAN/CSA C22.2 No. 61010-2-010; CAN/CSA C22.2 No. 61010-2-010:2004; CAN/CSA C22.2 No. 60601-2-010:2015; UL 61010-2-010; UL 61010-2-010:2004; UL 60601-2-010:2015

Test Technology:	**Test Method(s) [2, 3, 4]:**
Laboratory Centrifuges	IEC 61010-2-020; IEC 61010-2-020:2006; EN 61010-2-020; EN 61010-2-020:2006; CAN/CSA C22.2 No. 61010-2-020; CAN/CSA C22.2 No. 61010-2-020:2009; UL 61010-2-020 (*excluding 7.3.101 & 7.6*)
Equipment for Mixing and Stirring	IEC 61010-2-051; IEC 61010-2-051:2003; IEC 61010-2-051:2015; EN 61010-2-051; EN 61010-2-051:2003; CAN/CSA C22.2 No. 61010-2-051; CAN/CSA C22.2 No. 61010-2-051:2004; CAN/CSA C22.2 No. 61010-2-051:2015; UL 61010-2-051; UL 61010-2-051:2003; UL 61010-2-051:2015
Automatic & Semi-Automatic Laboratory Equipment for Analysis	IEC 61010-2-081; IEC 61010-2-081:2001 + A1:2003; IEC 61010-2-081:2015; CAN/CSA C22.2 No. 61010-2-081; CAN/CSA C22.2 No. 61010-2-081:2004; CAN/CSA C22.2 No. 61010-2-081:2015; EN 61010-2-081; EN 61010-2-081:2002 + A1:2003; UL 61010-2-081; UL 61010-2-081:2015
IVD Medical Equipment	IEC 61010-2-101; IEC 61010-2-101:2002; IEC 61010-2-101:2015; EN 61010-2-101; EN 61010.2-101:2002; CAN/CSA C22.2 No. 61010-2-101; CAN/CSA C22.2 No. 61010-2-101:2004; CAN/CSA C22.2 No. 61010-2-101:2015; UL 61010-2-101; UL 61010-2-101:2015

On the following products or types of products:
Medical Equipment, Laboratory Equipment

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* [1]:

Standards:
ANSI/AAMI ES 60601-1:2005/(R2012) and A1:2012, C1:2000/(R2012) and A2:2010/(R2012) – FDA Recognition Number: 19-4; ANSI AAMI ES60601-1:2005/(R)2012 and A1:2012, C1:2009/(R)2012 and A2:2010/(R)2012 (Consolidated Text) [Incl. AMD2:2020] – FDA Recognition Number: 19-46; ANSI AAMI HA60601-1-11:2015 [Including AMD1:2021] – FDA Recognition Number: 19-47
*IEC 60601-1-2:2014 + A1:2020 – FDA Recognition Number: 19-36; *IEC 60601-1-2:2014 – FDA Recognition Number: 19-8; *ANSI/AAMI/IEC 60601-1-2:2014 + A1:2021 – FDA Recognition Number: 19-36; *ANSI/AAMI/IEC 60601-1-2:2014 – FDA Recognition Number: 19-8; ***Exclusion from the laboratory scope of accreditation in the non-FDA ASCA section of this scope are listed above for some clauses of the 60601 methods (EDIT THIS TO MATCH THE EXCLUSION TECHNOLOGY ABOVE, Test Technology Titled: Medical electrical equipment – Electromagnetic disturbances** IEC 60601-1-6 Edition 3.2 2020-07 CONSOLIDATED VERSION – FDA Recognition Number: 5-132; IEC 60601-1-6 Edition 3.1 2013-10 – FDA Recognition Number: 5-89; IEC 60601-1-8 Edition 2.2 2020-07 CONSOLIDATED VERSION – FDA Recognition Number: 5-131; IEC 60601-1-8 Edition 2.1 2012-11 – FDA Recognition Number: 5-76 ANSI AAMI IEC 60601-1-8:2006 and A1:2012 [Including AMD2:2020] – FDA Recognition Number: 5-131 ANSI/AAMI/IEC 60601-1-8:2006 and A1:2012 – FDA Recognition Number: 5-76; IEC 60601-1-10 Edition 1.2 2020-07 CONSOLIDATED VERSION – FDA Recognition Number: 19-37; IEC 60601-1-10 Edition 1.1 2013-11 – FDA Recognition Number: 19-9; IEC 60601-1-11 Edition 2.1 2020-07 CONSOLIDATED VERSION – FDA Recognition Number: 19-38; IEC 60601-1-11 Edition 2.0 2015-01 – FDA Recognition Number: 19-14; ANSI AAMI HA60601-1-11:2015 – FDA Recognition Number:19-16; IEC 60601-1-12 Edition 1.1 2020-07 CONSOLIDATED VERSION – FDA Recognition Number: 19-39
Standards:
ANSI/AAMI/IEC 60601-1-12:2016 + A1:2021 – FDA Recognition Number: 19-39; IEC 60601-1-12 Edition 1.0 2014-06 – FDA Recognition Number: 19-15; ANSI/AAMI/IEC 60601-1-12:2016 – FDA Recognition Number: 19-15
IEC 60601-2-2 Edition 6.0 2017-03 – FDA Recognition Number: 6-389; ANSI AAMI IEC 60601-2-2:2017 – FDA Recognition Number: 6-389; IEC 60601-2-10 Edition 2.1 2016-04 – FDA Recognition Number: 17-16; IEC 60601-2-16 Edition 5.0 2018-4 – FDA Recognition Number: 9-121; ANSI AAMI IEC 60601-2-16:2018 – FDA Recognition Number: 9-121; IEC 60601-2-18: Edition 3.0 2009-08 – FDA Recognition Number: 9-114; IEC 60601-2-25 Edition 2.0 2011-10 – FDA Recognition Number: 3-105; ANSI AAMI IEC 60601-2-25:2011/(R)2016 – FDA Recognition Number: 3-105; IEC 60601-2-27 Edition 3.0 2011-03 – FDA Recognition Number: 3-126; ANSI AAMI IEC 60601-2-27:2011(R)2016 – FDA Recognition Number: 3-126; IEC 60601-2-34 Edition 3.0 2011-05 – FDA Recognition Number: 3-115; IEC 60601-2-36 Edition 2.0 2014-04 – FDA Recognition Number: 9-119; IEC 60601-2-37:2007 + A1:2015 – FDA Recognition Number: 12-293; IEC 60601-2-47 Edition 2.0 2012-02 – FDA Recognition Number: 3-155; ANSI AAMI IEC 60601-2-47:2012/(R)2016 - FDA Recognition Number: 3-155; IEC 60601-2-52 Edition 1.0 2009-12 – FDA Recognition Number: 6-321

Standards:
IEC 61010-1 Edition 3.1 2017-01 – FDA Recognition Number: 19-34; ANSI/UL 61010-1 3rd Ed, dated May 12, 2012 with revision through July 19, 2019 – FDA Recognition Number: 19-41
IEC 80601-2-30 Edition 2.0 2018-03 – FDA Recognition Number: 3-123; ANSI AAMI IEC 80601-2-30:2018 – FDA Recognition Number: 3-123; IEC 80601-2-35 Edition 2.1 2016-04 – FDA Recognition Number: 6-390; ANSI/AAMI/IEC 80601-2-35:2009 + A1:2016 – FDA Recognition Number: 6-390
ISO 80601-2-55 Second Edition 2018-02 – FDA Recognition Number: 1-140; ISO 80601-2-56 Second Edition 2017-03 – FDA Recognition Number: 6-421; ISO 80601-2-61 Second Edition 2017-12 (Corrected Version 2018-02) – FDA Recognition Number: 1-139

[1] 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 www.FDA.gov.

[2] Broadband random vibration, Mechanical shock, and any requirements associated with Clinical Data and/or Clinical Trials are not included within the testing listed on the scope of accreditation.

[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.

[4] 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*.

Accredited Laboratory

A2LA has accredited

MEDICAL EQUIPMENT COMPLIANCE ASSOCIATES, LLC.

Oak Creek, Wisconsin

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 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 12th day of January 2023.

Mr. Trace McInturff, Vice President, Accreditation Services
For the Accreditation Council
Certificate Number 3392.01
Valid to January 31, 2025

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