SCOPE OF ACCREDITATION TO ISO/IEC 17025:2017

DEVICE CONFORMITY AND TESTING, LLC
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ELECTRICAL

Valid To: June 30, 2028 Certificate Number: 6136.01

In recognition of the successful completion of the A2LA evaluation process (including an assessment of the organization's compliance with A2LA's R256 Specific Program Accreditation requirements), accreditation is granted to this laboratory to perform the following tests:

Test Technology:	**Test Method(s) [1,2,3]:**
Information Technology	IEC 60950-1:2005/AMD2:2013
Information and Communication Technology Equipment	IEC 62368-1:2018[4]; IEC 62368-1 Ed. 4 2023-05[4]
Medical Equipment	IEC 60601-1:1988/AMD2:1995; IEC 60601-1:2005+A1:2012; IEC 60601-1:2005+A1:2012+A2:2020; ES60601-1:2005/(R)2012 and A1:2012 C1:2009/(R)2012 and A2:2010/(R)2012 (Consolidated Text); IEC 60601-1-4:1996/AMD1:1999; IEC 60601-1-6:2010+A1:2013+A2:2020; IEC 60601-1-6 Edition 3.1 2013-10; IEC 60601-1-6 Edition 3.2 2020-07 CONSOLIDATED VERSION; IEC 60601-1-8:2006+A1:2012+A2:2020; IEC 60601-1-8 Edition 2.1 2012-11; IEC 60601-1-8 Edition 2.2 2020-07 CONSOLIDATED VERSION; IEC 60601-1-11:2015+A1:2020; IEC 60601-1-11 Edition 2.0 2015-01; IEC 60601-1-11 Edition 2.1 2020-07 CONSOLIDATED VERSION; IEC 60601-2-2 Edition 6.0 2017-03; IEC 60601-2-2 Edition 6.1 2023-02; IEC 60601-2-18:2009-08

Test Technology:	Test Method(s) [1,2,3]:
Medical Equipment (Continued)	IEC 60601-2-64:2014-09; IEC 60601-2-68 Edition 1.0 2014-09
Electrical Equipment for Measurement, Control, and Laboratory Use	IEC 61010-1:2010+A1:2016; IEC 61010-1 Edition 3.1 2017-01 CONSOLIDATED VERSION; IEC 61010-2-010:2019; IEC 61010-2-051:2018; IEC 61010-2-101:2018

[1] 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 test method, per Annex A, Part C of A2LA's *R101 - General Requirements: Accreditation of Conformity Assessment Bodies.*

[2] 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.

[3] For any test method requiring an IP rating, the CAB is limited to IPx0, where x = 0, 1, 2, 3 or 4.

[4] Excluding:
Annex B – Excluding equipment that connects to the wired telephone network.
Annex B – Excluding equipment intended for building-in or rack mounting.
Annex E – Any equipment with audio amplifiers.
Annex G – Components.
Annex J – Insulated winding wires.
Annex L – Disconnect devices.
Annex S – Tests for resistance to heat and fire.

Accredited Laboratory

A2LA has accredited

DEVICE CONFORMITY AND TESTING, LLC

Devens, MA

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 27th day of July 2026.

Mr. Trace McInturff, Vice President, Accreditation Services
For the Accreditation Council
Certificate Number 6136.01
Valid to June 30, 2028

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