UL 61010-2-101

STANDARD FOR

Safety Requirements for Electrical Equipment for Measurement, Control and Laboratory Use – Part 2-101: Particular Requirements for In Vitro Diagnostic (IVD) Medical Equipment

UL Standard for Safety Requirements for Electrical Equipment for Measurement, Control and Laboratory Use – Part 2-101: Particular Requirements for In Vitro Diagnostic (IVD) Medical Equipment, UL 61010-2-101

Third Edition, Dated July 31, 2019

Summary of Topics

Adoption of IEC 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 (third edition, issued by IEC October 2018) as a new IEC-based UL standard, UL 61010-2-101 with No US Differences.

The new requirements are substantially in accordance with Proposal(s) on this subject dated May 3, 2019.

No Text on This Page

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Standard for Safety Requirements for Electrical Equipment for Measurement, Control and Laboratory Use – Part 2-101: Particular Requirements for In Vitro Diagnostic (IVD) Medical Equipment

Second Edition – August 2015

Third Edition

July 31, 2019

This ANSI/UL Standard for Safety consists of the Third Edition.

The most recent designation of ANSI/UL 61010-2-101 as an American National Standard (ANSI) occurred on July 31, 2019. ANSI approval for a standard does not include the Cover Page, Transmittal Pages, Title Page, or Preface. The IEC Foreword is also excluded from the ANSI approval of IEC-based standards.

Comments or proposals for revisions on any part of the Standard may be submitted to UL at any time. Proposals should be submitted via a Proposal Request in UL's On-Line Collaborative Standards Development System (CSDS) at https://csds.ul.com.