Certificate

TÜVRheinland®

Certificate no. CU 72180312 01

License Holder:
Euroimmun Medizinische
Labordiagnostika AG
Seekamp 31
23560 Lübeck
Germany

Manufacturing Plant:
Euroimmun Medizinische
Labordiagnostika AG
Werkstrasse 2-22
23942 Dassow
Germany

Test report no.: USA- 31880266 002 **Client Reference:** Mrs. Hammoud-Schuett

Tested to: UL 61010-1:2012 R4.16
CAN/CSA-C22.2 NO. 61010-1-12 + GI1 + GI2 (R2017)
UL 61010-2-101:2015
CAN/CSA-C22.2 NO. 61010-2-101:15

Certified Product: Work Station IFA **License Fee - Units**

Model Designation: EUROLabWorkstation IFA 7

Ratings: AC 120 V/230 V, 50/60 Hz, 850 VA
Protection Class: I

7

Appendix: 1, 1-6

Licensed Test mark:

TÜVRheinland
C US

Date of Issue (day/mo/yr)
11/09/2018

TÜVRheinland®
LGA
Precisely Right.

Date : 2019/07/16

Euroimmun Medizinische
Labordiagnostika AG
Seekamp 31
23560 Lübeck
Germany
Attn: Stefanie Hammoud-Schuett

Re. : CU US + Canada Certificate

Type of Equipment	:	Work Station IFA
Model Designation	:	See Certificate
Certificate No.	:	CU 72180312 0002
File No.	:	31880266 003
Engineer/Contact	:	Volker Ebinghaus
Standard(s)	:	UL 61010-1:2012 R4.16
		CAN/CSA-C22.2 NO. 61010-1-12 + GI1 + GI2 (R2017)
		UL 61010-2-101:2015
		CAN/CSA-C22.2 NO. 61010-2-101:15

Dear Madame or Sir,

The above referenced technical equipment has been tested and was found to be in compliance with the listed test requirement(s). Enclosed, please find the TUV Rheinland approval document No. CU 72180312 0002.
It authorizes you to label the listed product(s) with the TUV Rheinland Mark identified in the approval document.
For compliance, the Test Mark must be on the approved unit.

Your product is subject to regular factory follow-up inspections as well as annual certificate and factory registration fees.

In using the TUV Rheinland Mark you are obligated to comply with the TUV Rheinland of North America Service Agreement.

If we can be of any further assistance to you, please do not hesitate to contact us.

Sincerely yours,
Certification Body

Gregor Stupp
QA Certification Officer

Enclosure

TÜV Rheinland
of North America, Inc.
North American Headquarters

12 Commerce Road
Newtown, CT 06470

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Member of
TÜV Rheinland Group

Certificate

TÜVRheinland®

Certificate no. CU 72180312 02

License Holder:
Euroimmun Medizinische
Labordiagnostika AG
Seekamp 31
23560 Lübeck
Germany

Manufacturing Plant:
EUROIMMUN Medizinische
Labordiagnostika AG
An der Trave 1
23923 Selmsdorf
Germany

Test report no.: USA- 31880266 003 **Client Reference:** Mrs. Hammoud-Schuett

Tested to:
UL 61010-1:2012 R4.16
CAN/CSA-C22.2 NO. 61010-1-12 + GI1 + GI2 (R2017)
UL 61010-2-101:2015
CAN/CSA-C22.2 NO. 61010-2-101:15

Certified Product: Work Station IFA **License Fee - Units**

Product as described on page 01.

Change:
Address of manufacturing plant changed,
see above. (K757960)

Licensed Test mark:

TÜVRheinland
C US

Date of Issue (day/mo/yr)
16/07/2019

TÜV Rheinland of North America, Inc., 12 Commerce Road, Newton, CT 06470, Tel (203) 426-0888 Fax (203) 426-4009