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# **EU** Certificate

# Quality Management System REGULATION (EU) 2017/746 on In Vitro Diagnostic Medical Devices Annex IX Chapters I and III

Registration No.: HX 1483000-1
Manufacturer: EUROIMMUN

Medizinische Labordiagnostika AG

Seekamp 31 23560 Lübeck Germany

EUDAMED Single Registration No.:

DE-MF-000005296

Products:

Products of class B:

IMMUNOCHEMISTRY (IMMUNOLOGY)

IVR 0602: Devices intended to be used for screening, determination or monitoring of physiological markers for a

specific disease

W01021001 - AUTOIMMUNE CONNECTIVE TISSUE

**DISEASES** 

W01021002 - NEURO-AUTO-IMMUNE DISEASE W01021090 - VARIOUS AUTO-IMMUNE DISEASE W01021112 - ANTI-CYCLIC CITRULLINATED PEPTIDE

The Notified Body hereby declares that the requirements of Annex IX, Chapter I of the REGULATION (EU) 2017/746 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.

If class D devices are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4 is required before placing them on the market.

If class B, C or D devices for self-testing or near-patient testing are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.1 is required before placing them on the market. If companion diagnostics are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.2 is required before placing them on the market.

 Report No.:
 1195791-20

 Effective date:
 2025-08-20

 Expiry date:
 2028-05-09

 Issue date:
 2025-08-20

V. Ull

Dr. Volker Schlüter TÜV Rheinland LGA Products GmbH Tillystraße 2 · 90431 Nürnberg · Germany

This certificate can be validated on https://www.certipedia.com

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/746 concerning medical devices with the identification number 0197.





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Products:

W01021520 - CONTROLS - IMMUNOCHEMISTRY

W01021199 - RHEUMATOID / INFLAMMATORY DISEASE

MARKERS - OTHER

W01020190 - OTHER SPECIFIC PROTEINS

IVR 0603: Devices intended to be used for screening, confirmation/determination, or monitoring of allergies and

intolerances

W01020201 - IMMUNOGLOBULIN E - TOTAL

W01020204 - IMMUNOGLOBULIN E – MONOTEST/PLURIRESULT-MULTI AG

IVR 0608: Devices intended to be used for screening, determination or monitoring of physiological markers

W01020190 - OTHER SPECIFIC PROTEINS

W01020702 - VITAMINES

W01021520 - CONTROLS - IMMUNOCHEMISTRY

W01029099 - IMMUNOCHEMISTRY REAGENTS - OTHER

### INFECTIOUS DISEASES

IVR 0503: Devices intended to be used to detect the presence of, or exposure to an infectious agent including sexually transmitted agents

W01050106 - LYME BORRELIOSIS

W01050117 - OTHER BACTERIOLOGY IMMUNOASSAYS

W01050404 - EPSTEIN BARR VIRUS

W01050405 - OTHER VIROLOGY - NA REAGENTS W01050406 - OTHER VIROLOGY ANTIGEN/ANTIBODY

**DETECTION** 

W01050407 - MEASLES VIRUS





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Products:

W01050502 - MISCELLANEOUS PARASITOLOGY W01050808 - CONTROLS - INFECT, IMMUNOLOGY

IVR 0504: Devices intended to be used to determine the infectious load, to determine infective disease status or immune status and devices used for infectious disease staging W01050117 - OTHER BACTERIOLOGY IMMUNOASSAYS

CHEMISTRY / IMMUNOCHEMISTRY INSTRUMENTS IVR 0503: Devices intended to be used to detect the presence of, or exposure to an infectious agent including sexually transmitted agents

W0201020192 - AUTOMATED IMMUNOCHEMISTRY ANALYSERS – IVD MEDICAL DEVICE SOFTWARE

Products of class C:

IMMUNOCHEMISTRY (IMMUNOLOGY)

IVR 0602: Devices intended to be used for screening, determination or monitoring of physiological markers for a specific disease

W01021001 - AUTOIMMUNE CONNECTIVE TISSUE

DISEASES

W01021090 - VARIOUS AUTO-IMMUNE DISEASE

W01050501 - TOXOPLASMA

### **INFECTIOUS DISEASES**

IVR 0501: Devices intended to be used for pre-natal screening of women in order to determine their immune status towards transmissible agents





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Products:

W01050101 - CHLAMYDIA W01050401 - RUBELLA VIRUS W01050402 - CYTOMEGALOVIRUS W01050501 - TOXOPLASMA

IVR 0503: Devices intended to be used to detect the presence of, or exposure to an infectious agent including sexually transmitted agents

W01050405 - OTHER VIROLOGY - NA REAGENTS W01050107 - MYCOBACTERIA GENUS + SPECIES

W01050501 - TOXOPLASMA

W01050103 – SYPHILIS (TREPONEMA PALLIDUM)

W01050409 - VARICELLA ZOSTER VIRUS

### **GENETIC TESTING**

IVR 0402: Devices intended to be used to predict genetic

disease/disorder risk and prognosis

W01060101 - MONOGENETIC DISORDERS

W01060104 - POLYMORPHISMS

### NUCLEIC ACID TESTING INSTRUMENTS

IVR 0402: Devices intended to be used to predict genetic

disease/disorder risk and prognosis

W02050292 - MICRO-ARRAY INSTRUMENTS - IVD

MEDICAL DEVICE SOFTWARE

### CHEMISTRY / IMMUNOCHEMISTRY INSTRUMENTS

IVR 0501: Devices intended to be used for pre-natal screening of women in order to determine their immune status towards transmissible agents





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Products:

W0201020192 - AUTOMATED IMMUNOCHEMISTRY ANALYSERS - IVD MEDICAL DEVICE SOFTWARE

Authorized representative(s): N/A





# **Quality Management System** REGULATION (EU) 2017/746 on In Vitro Diagnostic Medical Devices **Annex IX Chapters I and III**

Registration No.: HX 1483000-1 Manufacturer: **EUROIMMUN** 

Medizinische Labordiagnostika AG

Seekamp 31 23560 Lübeck Germany

**EUDAMED Single** Registration No.:

DE-MF-000005296

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2023-05-10
1	Scope extension, EUROI_PDQ2_HX_2023-07- 12_2_20230822_extsigned.pdf	2023-08-22
2	Scope extension: Products of class B (W01050405, W01021199, W01020204, W01020190, W01021520) Scope reduction: Products of class B (W01020299) and class C (W01050403, W01050705) EUROI_PDQ2_HX_2023-12-15_2024-03-26_extsigned.pdf	2024-03-26
3	Scope extension: Products of class B (W01050117, W01021001), Products of class C (W01021001), EUROI_PDQ2_HX_2024-03-06_exsigned.pdf	2024-05-17
4	Scope extension: Product of class D (W01050406) and class C and B EUROI PDQ2 HX 2024-08-07	2024-08-19
5	Scope extension: Products of class B (W01050407, W01029099), EUROI PDQ2 HX 2024-12-18.pdf	2024-12-18
6	Scope extension: Products of class C (W01050501) Scope reduction: Products of class B (W01050106) EUROI PDQ2 HXIX 2025-03-26.pdf	2025-03-31
7	Scope extension: Products of class B (W01050106, W01021002) and class C (W01050101, W01050401) Scope reduction: Products of class D (W01050406)	2025-06-13
8	Scope extension: Products of class B (W01020190) and class C (W01050103, W01050409)	2025-08-20



