ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-IVDR-099

Product Service

EU Quality Management System Certificate (IVDR)

Pursuant to Regulation (EU) 2017/746 on in Vitro Diagnostic Medical Devices, Annex IX Chapters I and III (Class C and B Devices excluding self-/near-patient-testing and Companion Diagnostics)

No. V12 074544 0021 Rev. 00

Manufacturer:

URIT Medical Electronic Co., Ltd.
No. D-07
Information Industry District
High-tech Zone
541004 Guilin, Guangxi
PEOPLE'S REPUBLIC OF CHINA

SRN Manufacturer - CN-MF-000011840

Authorized Representative:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80, 20537 Hamburg, GERMANY

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (8) of the Regulation (EU) 2017/746 on in Vitro Diagnostic Medical Devices. Details on devices covered by the quality management system are described on the following page(s).
The Report referenced below summarizes the result of the assessment and includes reference to relevant CS, harmonized standards, audit and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.
The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.
The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment includes an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples.All applicable requirements of the Testing, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with.
For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:V12 074544 0021 Rev. 00

Report No.: GZ2216303

Valid from: 2024-06-04
Valid until: 2029-06-03

Digitally signed by Stoianova Elena
Date: 2025.07.10 12:11:39 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Marta Carnielli
Head of Certification IVD

Issue date: 2024-06-04

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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No. V12 074544 0021 Rev. 00

Classification:	Class B
Device Group:	W0101 - CLINICAL CHEMISTRY
Intended Purpose:	IVR 0608 - Devices intended to be used for screening, determination or monitoring of physiological markers

Classification:	Class B
Device Group:	W0202 - HEMATOLOGY / HISTOLOGY / CYTOLOGY INSTRUMENTS
Intended Purpose:	IVR 0608 - Devices intended to be used for screening, determination or monitoring of physiological markers

The validity of this certificate depends on conditions and/or is limited to the following:	-none-

Revision History:

Rev.	Dated	Report	Description
00	2024-06-04	GZ2216303	Initial issuance

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