Human
Gesellschaft für Biochemica
und Diagnostica mbH

Max-Planck-Ring 21
D-65205 Wiesbaden
Germany

Digitally signed by Iurchevici Valeriu
Date: 2024.02.14 17:41:06 EET
Reason: MoldSign Signature
Location: Moldova

Certificate

We, HUMAN GmbH, Max-Planck-Ring 21, 65 205 Wiesbaden, Germany,
hereby certify that the company

Echipamed Plus SRL

str. Valea Trandafirilor, 24B
MD-2001, Chisinau
Moldova

is the exclusive distributor of

HUMAN GmbH

in the territory of Moldova Republic,
effective by October 11, 2000.

Wiesbaden, February 21, 2007

HUMAN GmbH

Ralph Neuberger
Director of Sales & Marketing

Human
Gesellschaft für Biochemica
und Diagnostica mbH
Max-Planck-Ring 21
65205 Wiesbaden-Delkenheim
Germany

ECHIPAMED-PLUS
IDNO 1003600077677

Certificate

mdc medical device certification GmbH
certifies that

Human

Gesellschaft für Biochemica und Diagnostica mbH
Max-Planck-Ring 21
65205 Wiesbaden
Germany

with the location

Stegelitzer Straße 3
39126 Magdeburg

for the scope

development, manufacturing and distribution of in vitro diagnostic products and analyzers

has introduced and applies a

Quality Management System

The mdc audit has proven that this quality management system meets all requirements of the following standard

EN ISO 9001

Quality management systems – Requirements

(ISO 9001:2015)

Valid from	2022-03-12
Valid until	2025-03-11
Registration no.	D1030000083
Report no.	P21-01793-219728
Stuttgart	2022-03-10

Head of Certification Body

medical device certification mdc

DAkkS Deutsche Akkreditierungsstelle D-ZM-16002-01-01

mdc medical device certification GmbH
Kriegerstraße 6
D-70191 Stuttgart, Germany
Phone: +49-(0)711-253597-0
Fax: +49-(0)711-253597-10
Internet: http://www.mdc-ce.de

Certificate

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Human

Gesellschaft für Biochemica und Diagnostica mbH
Max-Planck-Ring 21
65205 Wiesbaden
Germany

with the location

Stegelitzer Straße 3
39126 Magdeburg

for the scope

development, manufacturing and distribution of reagents, reagent products, calibrators and control materials for clinical chemistry, haematology, haemostasis, immune chemistry as well as analyzers for in-vitro diagnosis.
distribution of test kits for molecular biology analyses as well as rapid tests for the diagnosis of infectious diseases.

has introduced and applies a

Quality Management System

The mdc audit has proven that this quality management system meets all requirements of the following standard

EN ISO 13485

Medical devices – Quality management systems – Requirements for regulatory purposes

EN ISO 13485:2016 + AC:2016 - ISO 13485:2016

Valid from	2022-03-12
Valid until	2025-03-11
Registration no.	D1030000084
Report no.	P21-01793-219732
Stuttgart	2022-03-10

Head of Certification Body

DAkkS
Deutsche Akkreditierungsstelle
D-ZM-16002-06-00

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