sysmex

Sysmex Europe GmbH Bornbarch 1 · 22848 Norderstedt Germany

To whom it may concern

Sysmex Europe GmbH
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LETTER OF AUTHORIZATION

Whereas Sysmex Europe GmbH ("Sysmex"), who are established, reputable and authorised representative in Europe, Africa and Middle East (EMEA region), officially announced by the manufacturer Sysmex Corporation, having its principal place of business at 1-5-1 Wakinohama Kaigandoori, Chuo-ku, Kobe 651-0073, Japan, and

having the power to grant authorizations to local representatives within the above mentioned markets, do hereby declare that

ECHIPAMED Plus SRL,
Valea Trandafirilor 24 «B», off. 80, MD-2001 Chisinau, Moldova (the **"COMPANY"**)

is our distributor and local representative for the following Sysmex products:

Sysmex Haematology- and Urine Analysers
Reagents, Accessories, Software and spare parts (the **"PRODUCTS").**

in the territory of Republic of Moldova (the**" TERRITORY").**

The **COMPANY** is therefore authorized to carry out all commercial and support activities for the **PRODUCTS** including sales, marketing, application, registration and field service support in the **TERRITORY.**
The **COMPANY** is aware that this special authorisation is limited to the above listed **PRODUCTS** and does not create any further rights for the **COMPANY.**
We hereby grant our warranty following our general conditions of sale for the **PRODUCTS** delivered, consisting of and limited to:
Free of charge supply of spare parts to the **COMPANY** as replacement for defective new parts for a period of 14 months after B/L - AWB date.
This declaration is valid until 31.03.2020 and may be revoked unilaterally by Sysmex in writing before that date for due cause. Such due cause shall, among others, be the termination or expiration of the distributorship relationship, if any, between Sysmex and the **COMPANY.**

On behalf of Sysmex Europe GmbH

Jan-Willem Schipper
Senior Executive Officer

sysmex
Sysmex Europe GmbH
Bornbarch 1
22848 Norderstedt

Date: April 16, 2019
Place: 22848 Norderstedt

Digitally signed by Iurchevici Valeriu
Date: 2020.03.19 12:36:08 EET
Reason: MoldSign Signature
Location: Moldova

ECHIPAMED-PLUS
REPUBLICA MOLDOVA mun. CHIȘINĂU. SOCIETATEA CU RĂSPUNDERE LIMITATĂ
IDNO 1003600077677

Certificate

Standard **ISO 9001:2015**

Certificate Registr. No. 01 100 110072

Certificate Holder:

SYSMEX EUROPE GmbH
Bornbarch 1
22848 Norderstedt
Germany

including the locations according to annex

Scope: Sales and service of devices, reagents and accessories for in-vitro diagnostics in the area of haematology, urine analytics, coagulation and detection of an epithelial cell marker for the diagnosis of metastases in lymph nodes, as well as of products in the area of laboratory automation and laboratory EDP systems. Design, development and manufacturing of software for in-vitro diagnostic use. Distribution of magnetic sensing devices, probes, associated equipment and sterile magnetic markers. Distribution and servicing of scalp-cooling devices with accessories.

Proof has been furnished by means of an audit that the requirements of ISO 9001:2015 are met.

Validity: The certificate is valid from 2018-09-06 until 2020-07-24.
First certification 2011

2018-09-13

TÜV Rheinland Cert GmbH
Am Grauen Stein · 51105 Köln

www.tuv.com

DAkkS Deutsche Akkreditierungsstelle D-ZM-16031-01-00

TÜVRheinland®
Precisely Right.

TÜVRheinland®

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

SYSMEX EUROPE GmbH
Bornbarch 1
22848 Norderstedt
Deutschland

has establisned and applies a quality management system for medical devices for the following scope:

see attachment

Proof has been furnished that the requirements specified in

EN ISO 13485:2016

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date:	2019-05-17
Certificate Registration No.:	SX 60137613 0001
An audit was performed. Report No.:	21245244 005
This Certificate is valid until:	2022-05-16

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

Date 2019-04-29

Dipl.-Ing. Sven Hoffmann

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail:cert-validity@de.tuv.com http://www.tuv.com/safety

Certificate

Standard **ISO 14001:2015**

Certificate Registr. No. **01 104 110072**

Certificate Holder:

SYSMEX EUROPE GmbH
Bornbarch 1
22848 Norderstedt
Germany

including the location
Sysmex Deutschland GmbH
Bornbarch 1
22848 Norderstedt
Germany

Scope: Sales, marketing and service of in vitro diagnostic medical devices

Proof has been furnished by means of an audit that the requirements of ISO 14001:2015 are met.

Validity: The certificate is valid from 2017-07-25 until 2020-07-24.
First certification 2011

2018-02-12

TÜV Rheinland Cert GmbH
Am Grauen Stein · 51105 Köln

www.tuv.com

DAkkS Deutsche Akkreditierungsstelle D-ZM-16031-01-00

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