



Sysmex Europe SE - Bornbarch 1 - 22848 Norderstedt - Germany

To whom it may concern

Sysmex Europe SE

Bornbarch 1

22848 Norderstedt, Germany

Phone +49 40 527 26-0

Fax +49 40 527 26-100

info@sysmex-europe.com

LETTER OF AUTHORIZATION

Whereas Sysmex Europe SE ("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 Wakino-hama-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 the company

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

with Reagents, Accessories, Software and Spare Parts

(the "Products")

In the territory 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.

Company Location Norderstedt
Registered AG Kiel
HRB 24262 KI
VAT-ID DE 118 687 842
WEEE/ElektroG Reg. Nr. DE 159 56 453

Chairman of the
Supervisory Board:
Iwane Matsui

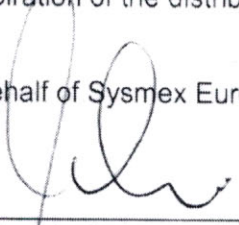
Management Board:
Alain Baverel (CEO)
Alberto Bonacini
Yuki Hyogu
Stefanie Schaal
Matthias Völkel

COMMERZBANK AG, Hamburg
IBAN DE20 2004 0000 0287 1879 00
SWIFT/BIC Code COBADEFFXXX

www.sysmex-europe.com

This declaration is valid until 31 March 2024 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 SE



Matthias Voelkel
Senior Executive Officer



Date: 08 March 2023
Place: 22848 Norderstedt, Germany

Sysmex Europe SE
Bornbarch 1
22848 Norderstedt



Certificate

Standard **ISO 9001:2015**

Certificate Registr. No. **09 100 89004**

Certificate Holder: **SYSMEX CORPORATION**
1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan
including the locations according to annex

Scope: Development, design, production, sales and servicing of in-vitro diagnostic medical devices, laboratory equipment, reagents and laboratory information system, and development, design, production and sales of customized recombinant protein

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

Validity: The certificate is valid from 2022-05-13 until 2024-07-31.
First certification 1998

2022-05-13



TÜV Rheinland Cert GmbH
Am Grauen Stein · 51105 Köln

www.tuv.com



Deutsche
Akkreditierungsstelle
D-ZM-16031-01-00



TÜVRheinland®
Precisely Right.

Certificate



Quality Management System
EN ISO 13485:2016

Registration No.: SX 1254782-1

Organization: SYSMEX CORPORATION
1-5-1 Wakinohama-Kaigandori,
Chuo-ku, Kobe
651-0073 Japan

Scope: Design and development, manufacture, distribution, installation and service of blood analyzer, urine analyzer, related reagents and accessories
Product categories: analyzers and reagents for hematological test, blood coagulation test, immune serum test, biochemical test, genetic test, bacteriological test and urine test

In accordance with EN ISO 13485:2016 Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)

The Certification Body of TÜV Rheinland LGA Products GmbH certifies that the organization has established and applies a quality management system for medical devices. Proof has been furnished that the requirements specified in the abovementioned standard are fulfilled. The quality management system is subject to yearly surveillance.

Report No.: 150258788-301

Effective date: 2022-04-28

Expiry date: 2024-07-31

Issue date: 2022-04-28



M. Aihara

Michiaki Aihara
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

EU Declaration of Conformity

Product identification:

Product name:	CELLPACK DCL
Model name:	N/A
REF code:	CT-661-628, CU-228-496
BUDI-DI:	4987562CELLPACKDCLX9
Intended Purpose	See attachment

Manufacturer:

Name:	SYSMEX CORPORATION
Single Registration Number:	JP-MF-000014037
Address:	1-5-1 Wakinoama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised representative:

Name:	SYSMEX EUROPE SE
Single Registration Number:	DE-AR-000022333
Address:	Bornbarch 1, 22848 Norderstedt, Germany

SYSMEX CORPORATION, as the manufacturer of the device, take sole responsibility for and hereby declare that the above mentioned device meets the provisions of the following Regulation:

- ☒ Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices
☐ Other Regulation(s)/Directive(s) as applicable for the device(s):

Risk class:

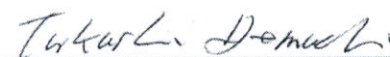
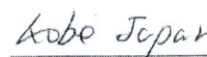
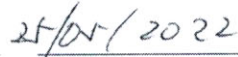
- ☒ A ☐ B ☐ C ☐ D

Conformity route:

Annex I+II+III according to Article 48 (10) of EU 2017/746

Common Specification:

N/A

		
Takashi Demachi	Kobe Japan	25/05/2022
Executive Vice President	Place	Date (DD.MM.YYYY)
Name	Function	

Sysmex Corporation

1-5-1 Wakinoama-Kaigandori, Chuo-ku, Kobe 651-0073, Japan
Tel 81-78-265-0500 Fax 81-78-265-0524



www.sysmex.co.jp

EU Declaration of Conformity

Product identification:

Product name:	SULFOLYSER
Model name:	N/A
REF code:	054-3351-4, 904-1131-7, AS788212, 904-1141-4
BUDI-DI:	4987562SULFOLYSERBV
Intended Purpose	See attachment

Manufacturer:

Name:	SYSMEX CORPORATION
Single Registration Number:	JP-MF-000014037
Address:	1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised representative:

Name:	SYSMEX EUROPE SE
Single Registration Number:	DE-AR-000022333
Address:	Bornbarch 1, 22848 Norderstedt, Germany

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- ☒ Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices
☐ Other Regulation(s)/Directive(s) as applicable for the device(s):

Risk class:

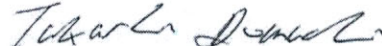
- ☒ A ☐ B ☐ C ☐ D

Conformity route:

Annex I+II+III according to Article 48 (10) of EU 2017/746

Common Specification:

N/A

		
Takashi Demachi	Name	Place
Executive Vice President	Function	Date (DD.MM.YYYY)

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EU Declaration of Conformity

Product identification:

Product name:	Lysercell WDF
Model name:	N/A
REF code:	AL-337-564, BG-689-680, AZ-124-801, AW-993-605
BUDI-DI:	4987562LysercellWDFXJ
Intended Purpose	See attachment

Manufacturer:

Name:	SYSMEX CORPORATION
Single Registration Number:	JP-MF-000014037
Address:	1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised representative:

Name:	SYSMEX EUROPE SE
Single Registration Number:	DE-AR-000022333
Address:	Bornbarch 1, 22848 Norderstedt, Germany

SYSMEX CORPORATION, as the manufacturer of the device, take sole responsibility for and hereby declare that the above mentioned device meets the provisions of the following Regulation:

- ☒ Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices
☐ Other Regulation(s)/Directive(s) as applicable for the device(s):

Risk class:

- ☒ A ☐ B ☐ C ☐ D

Conformity route:

Annex I+II+III according to Article 48 (10) of EU 2017/746

Common Specification:

N/A

		
Takashi Demachi	Kobe Japan	25/05/2022
Executive Vice President	Name	Date
	Function	(DD.MM.YYYY)

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www.sysmex.co.jp

EU Declaration of Conformity

Product identification:

Product name:	Fluorocell WDF
Model name:	N/A
REF code:	AE687941, BY458697, BJ284784, CV-377-552, AA-325-279
BUDI-DI:	4987562FLUOROCELLWDFWE
Intended Purpose	See attachment

Manufacturer:

Name:	SYSMEX CORPORATION
Single Registration Number:	JP-MF-000014037
Address:	1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised representative:

Name:	SYSMEX EUROPE SE
Single Registration Number:	DE-AR-000022333
Address:	Bornbarch 1, 22848 Norderstedt, Germany

SYSMEX CORPORATION, as the manufacturer of the device, take sole responsibility for and hereby declare that the above mentioned device meets the provisions of the following Regulation:

- ☒ Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices
☐ Other Regulation(s)/Directive(s) as applicable for the device(s):

Risk class:

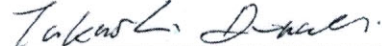
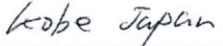
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Conformity route:

Annex I+II+III according to Article 48 (10) of EU 2017/746

Common Specification:

N/A

		
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Executive Vice President	Place	Date
Function		(DD.MM.YYYY)

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www.sysmex.co.jp

EU Declaration of Conformity

Product identification:

Product name:	CELLCLEAN
Model name:	N/A
REF code:	834-0162-1, BU037001
BUDI-DI:	4987562CELLCLEANP7
Intended Purpose	See attachment

Manufacturer:

Name:	SYSMEX CORPORATION
Single Registration Number:	JP-MF-000014037
Address:	1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised representative:

Name:	SYSMEX EUROPE SE
Single Registration Number:	DE-AR-000022333
Address:	Bornbarch 1, 22848 Norderstedt, Germany

SYSMEX CORPORATION, as the manufacturer of the device, take sole responsibility for and hereby declare that the above mentioned device meets the provisions of the following Regulation:

- ☒ Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices
☐ Other Regulation(s)/Directive(s) as applicable for the device(s):

Risk class:

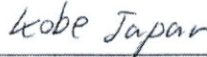
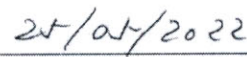
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Conformity route:

Annex I+II+III according to Article 48 (10) of EU 2017/746

Common Specification:

N/A

		
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EC Declaration of Conformity

Application of Council Directive:

- 98/79/EC of 27 October 1998 on In Vitro Diagnostic Medical Devices.

Means of conformity:

The following products are in conformity with

- Directive 98/79/EC based on the conformity assessment procedures in accordance with Annex III.

Product identification:

Product name: XN-L CHECK

Classification: Other device (except Annex II and self-testing devices)

List of Applied Standards:

- Harmonised Standards used for conformity assessment are listed in the technical documentation.

Legal Manufacturer:

Name: SYSMEX CORPORATION

Address: 1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised officer: Takashi Demachi Date: Apr. 28, 2022
Takashi Demachi, Executive Vice President

Authorised representative:

Name: SYSMEX EUROPE SE

Address: Bornbarch 1, 22848 Norderstedt, Germany

Authorised officer: Sinem Yaman Date: 28.04.2022
Sinem Yaman, Vice President, Head of Regulatory Affairs,
Quality Assurance, Quality Control

This declaration of conformity is issued under the sole responsibility of the manufacturer and is valid until the end of transitional period stipulated in REGULATION (EU) 2017/746 & its related regulations. The Classification of this product under REGULATION (EU) 2017/746 is Class B.



EC Declaration of Conformity

Application of Council Directive:

- 98/79/EC of 27 October 1998 on In Vitro Diagnostic Medical Devices.

Means of conformity:

The following products are in conformity with

- Directive 98/79/EC based on the conformity assessment procedures in accordance with Annex III.

Product identification:

Product name: XN CAL

Classification: Other device (except Annex II and self-testing devices)

List of Applied Standards:

- Harmonised Standards used for conformity assessment are listed in the technical documentation.

Legal Manufacturer:

Name: SYSMEX CORPORATION

Address: 1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised officer: Takashi Demachi Date: Apr. 28, 2022
Takashi Demachi, Executive Vice President

Authorised representative:

Name: SYSMEX EUROPE SE

Address: Bornbarch 1, 22848 Norderstedt, Germany

Authorised officer: Sinem Yaman Date: 28.04.2022
Sinem Yaman, Vice President, Head of Regulatory Affairs,
Quality Assurance, Quality Control

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EC Declaration of Conformity

Application of Council Directive:

- 98/79/EC of 27 October 1998 on In Vitro Diagnostic Medical Devices,
- 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment and its related amending directives including the (EU) 2015/863 amending Annex II of the Directive 2011/65/EU,
- 2014/53/EU of 16 April 2014 on the harmonisation of the laws of the Member States relating to the making available on the market of radio equipment.

Means of conformity:

- The following products are in conformity with
- Directive 98/79/EC based on the conformity assessment procedures in accordance with Annex III,
 - Directive 2011/65/EU based on the conformity assessment procedure in accordance with EN IEC 63000: 2018,
 - Directive 2014/53/EU based on the conformity assessment procedures using harmonized standards EN 50364:2010, EN 61010-2-101:2002, EN 61326-2-6:2006, EN 301 489-1 V2.2.0, EN 301 489-3 V2.1.1 and EN 300 330 V2.1.1 in accordance with Annex II Conformity Assessment Module A.

Product identification:

Product name: Automated Hematology Analyzer XN-L series
Model name: XN-550, XN-530, XN-450, XN-430, XN-350, XN-330
Accessories: Touch Panel Monitor TM104-SYX01
Classification: Other device (except Annex II and self-testing devices)

List of Applied Standards:

- Harmonised Standards used for conformity assessment are listed in the technical documentation.

Legal Manufacturer:

Name: SYSMEX CORPORATION
Address: 1-5-1 Wakinohama-Kaigandori, Chuo-ku, Kobe 651-0073 Japan

Authorised officer: Takashi Demachi Date: Apr. 28, 2022
Takashi Demachi, Executive Vice President

Authorised representative:

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Sinem Yaman, Vice President, Head of Regulatory Affairs,
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Sysmex Corporation

