

## **EU DECLARATION OF CONFORMITY**

RR Mechatronics Manufacturing B.V.

P.O.Box 225

This declaration of conformity is issued under the sole responsibility

1620 AE HOORN

of the manufacturer:

The Netherlands

De Corantijn 13

RR Mechatronics Manufacturing B.V.

1689 AN ZWAAG

De Corantijn 13, 1689 AN Zwaag, The Netherlands

The Netherlands

SRN: NL-MF-000023105

T+31 229 291 129

www.rrmechatronics.com

We declare that:

The whole blood diluent:

Trade name/ Model:

Starrsed Saline

Variants:

| Variant:        | Product-ID (REF): | UDI-DI         |
|-----------------|-------------------|----------------|
| Starrsed Saline | QRR010933         | 08719189137200 |

Basic UDI-DI (BUDI-DI): 8719189137ZL8SAEUW

EMDN-code:

W0103010105 Cbc-reagents (cleaning-/diluting-/lysing-/sheat-fluids)

Classification IVDR:

Class A

Intended purpose:

Product for the automatic cleaning of the needle and fill-nozzle assembly in Starrsed

ESR analyzers

is in conformity with the requirements of the following EU legislation(s):

**Regulation (EU) 2017/746** 

In vitro diagnostic medical devices

(conformity assessment according Article 48, §10 of this regulation)

DOC-2074 F4401 DoC Starrsed Saline.v1

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Conditions: By the ORGALIME GENERAL CONDITIONS S2012 of March 2012. Any other conditions are herewith explicitly rejected by us. VAT number: NL 8557 65 392 B01 CoC Number 64657353



The following harmonized standards have been applied:

EN ISO 13485:2016 Medical devices – Quality management systems – Requirements for regulatory purposes EN ISO 14971:2012 Medical devices – Application of risk management to medical devices EN-ISO 15223-1:2021 Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements EN ISO 18113-1:2011 In vitro medical devices – Information supplied by the manufacturer (labelling) – Part 1: Terms, definitions and general requirements In vitro diagnostic medical devices – Information supplied by the manufacturer (labelling) – EN ISO 18113-2:2011

In vitro medical devices – Evaluation of stability of in vitro diagnostic reagents

Part 2: In vitro diagnostic reagents for professional use

The CE mark was applied for the first time on this type of IVD reagent in 1998.

Place:

EN ISO 23640:2015

Zwaag, The Netherlands

Date:

May 14, 2024

Signature:

Stamp:

Name:

M.S van der Meer

Function title: CEO

RR Mechatronics Manufacturing B.V.



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