

# CERTIFICATE

Number: 2194749

The management system of the organization(s) and locations mentioned on the addendum belonging to:

## HTL-Strefa S.A.

Ul. Adamówek 7  
95-035 Ozorków  
Poland

Manufacturer Facility Identifier F000578

Conforms with the following standard and regulatory requirements:

## ISO 13485:2016

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002 and Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure  
Brazil: RDC ANVISA N. 16/2013, 23/2012 and 67/2009  
Canada: Medical Devices Regulations - Part 1- SOR 98/282  
Japan: MHLW Ministerial Ordinance 169, Article 4 to Article 68 and PMD Act  
United States: 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D and 21 CFR 820

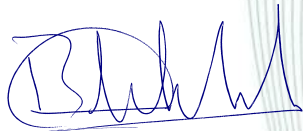
### Scope:

Design and development, manufacture and distribution of sterile single use blood lancets, pen needles and lancing devices.

Certificate expiry date: 2023-01-01  
Certificate effective date: 2022-01-24  
Certified since: 2019-01-28

This certificate is valid for the organization(s) and/or locations mentioned on the addendum.

DEKRA Certification B.V.



B.T.M. Holtus  
Managing Director



J.A. van Vugt  
Certification Manager

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The validation of the validity of this certificate can be checked through DEKRA's website using the following link:  
<https://www.dekra-product-safety.com/en/certified-organizations>

DEKRA Certification B.V. is recognized under the Medical Devices Single Audit Program.



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# ADDENDUM

To certificate: 2194749

The management system of the organization(s) and/or location(s) of:

## HTL-Strefa S.A.

Ul. Adamówek 7  
95-035 Ozorków  
Poland

Certified organization(s) and/or locations:

Different scope

HTL-Strefa S.A  
Ul. Lotnicza 21h  
99-100 Leczyca  
Poland

Manufacturer Facility Identifier F000578

Design and development, manufacture and distribution of sterile single use blood lancets, pen needles and lancing devices.

Addendum expiry date: 2023-01-01

Addendum effective date: 2022-01-24