Declaration of Conformity

This European Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER		
Name of Company	**Address**	**SRN**
SteriLance Medical(Suzhou)Inc.	No.168 PuTuoShan Road,New District,215153 Suzhou,Jiangsu, PEOPLE'S REPUBLIC OF CHINA	CN-MF-000002860

AUTHORIZED REPRESENTATIVE			
Name of Company	**Address**	**SRN**	**Phone/email**
Emergo Europe B.V.	Prinsessegracht 20,2514 AP,The Hague,The Netherlands	NL-AR-000000116	+31.70.345.8570 EmergoEurope@ul.com

PRODUCT IDENTIFICATION		
Product Name	**Type**	**Code / Catalog Number**
Disposable Safety Lancets	Press,Press Plus,Press2, Press2 Plus,Impress,Impress Pro,Lite3,Flex3,UltraSafe,Elite	V010401
Intended Purpose		**Basic UDI-DI**
The safety lancet is used for capillary blood collection.		6945630105BH

RISK CLASS FOR MEDICAL DEVICES		
Device Classification		**Common Specifications / Standards**
Class:	IIa	Medical Devices Regulation (EU) 2017/745
Rule:	6	

NOTIFIED BODY			
Name of Company	**ID Number**	**Conformity Assessment Procedure**	**Certificate Reference(s)**
TÜV SÜD Product Service Gmbh	CE 0123	Medical Devices Regulation (EU) 2017/745 ,Annex IX Chapters I and III	Certificate No.: G10 093119 0001 Rev.00 Issue date: 2022-11-24 Valid until: 2027-11-23

The SteriLance Medical(Suzhou)Inc. declares that the above-mentioned products meet the provision of the following EU legislation:

Medical Devices Regulation (EU) 2017/745

COMPANY REPRESENTATIVE: Zhang Xuanchao

TITLE: Quality Manager **SIGNATURE:** Zhang Xuanchao

PLACE: Suzhou **DATE:** 2022-12-29