

## **CERTIFICATE** OF REGISTRATION

This is to certify that the management system of:

## NISSHA MEDICAL TECHNOLOGIES SAS

Main site: 23-25 Boulevard de la Paix, 95800 Cergy, France

has been registered by Intertek as conforming to the requirements of

## EN ISO 13485:2016

## The management system is applicable to:

Design and development, manufacture and distribution of cables, leadwires and accessories for patient monitoring and diagnosis, cables and accessories for electrosurgery and defibrillation, prewired electrodes for cardiac monitoring and diagnosis, non-invasive blood pressure cuffs and hoses.

Distribution of sterile and non-sterile endoscopy tubing.

Distribution of patient monitors and accessories, vital signs, sensors and related accessories, oxygen therapy devices and related accessories, electrodes (ECG / neutral / defibrillation), electrosurgery devices and related accessories, defibrillators, and related accessories, endoscopy equipment and related accessories, medical chart papers, bandages and tapes. Servicing of patient monitors.

Certificate Number: 0183223 Revision: 02

Initial Certification Date: 6 August 2024

**Date of Certification Decision:** 30 April 2025

**Certificate Valid Date:** 30 April 2025

**Certificate Expiry Date:** 26 December 2027



Certifiering av ledningssystem

ISO/IEC 17021-1



Intertek

Krett

Brian Mather Certification Authority, Intertek Medical Notified Body AB

Intertek Medical Notified Body AB P.O. Box 1103, SE-164 22 Kista, Sweden





In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone.