EU DECLARATION OF CONFORMITY

According to the appendix IV of (EU) 2017/745 Regulation related to the medical devices
We,

HOIST MEDICAL SAS
23 Bis Quai de Rive Neuve 13007 Marseille
+33 4 91 76 73 29 - www.hoist-medical.com

SRN: FR-MF-000026140

Guarantee and declare under our sole responsibility that the medical device:

STARFIT™ NASOGASTRIC FEEDING TUBE WITH GUIDE WIRE

CODE	DESIGNATION	BASIC UDI
NGG	NASOGASTRIC FEEDING TUBE WITH GUIDE WIRE	376034811200.02BA

Invasive Medical Device class IIa intended for nasogastric enteral feeding pursuant to rule 5 of chapter III of appendix VIII of (EU) 2017/745 Regulation
Complies with the applicable requirements of (EU) 2017/745 Regulation and the French Public Health Code.

Notified body involved:
Name: DQS Medizinprodukte GmbH
Addresses: August-Schanz-Straße 21
60433 FRANKFURT AM MAIN Germany
Number: 0297
Assessment Evaluation Procedure: Annex IX (conformity assessment based on a quality management system and on assessment of technical documentation)

Certificate number: 551177 MDR2017Q

EMDN CODE: G02020101

Common specifications: No common specifications are used in relation to which conformity is declared

Marseille, November 1st, 2024

Bernard DAURELLE, CEO and PRRC

