

Number: 2262711CE02

# EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

Manufacturer:

**Nutricia Medical Devices B.V.**

Taurusavenue 167

2132 LS Hoofddorp

The Netherlands

SRN ID.: NL-MF-000012729

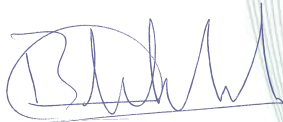
DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

**0344**

**Supplement to certificate: 93928CN**

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer/ authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.



B.T.M. Holtus  
Managing Director



J.M.A. McKenzie  
Principal Certification Manager

First Issued: 2 November 2022

Date: 2 November 2022

Expiry date: **1 November 2027**

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DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands  
T +31 88 96 83000 www.dekra.nl Company registration 09085396



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This certificate covers the following device(s) / groups of device(s):

G02020101, Nasogastric tubes (Class IIb)	
Flocare PUR nasogastric feeding tubes	<i>Intended Purpose:</i> Flocare nasogastric feeding tubes are intended for nutrition administration of enteral feeding into the stomach
Flocare PURSoft nasogastric feeding tubes	
Flocare Nutrisoft nasogastric feeding tubes	
G02020102, Nasojejunal/nasoduodenal tubes (Class IIb)	
Flocare Bengmark nasointestinal feeding tubes	<i>Intended Purpose:</i> Flocare nasointestinal feeding tubes are intended for nutrition administration of enteral feeding into the intestine

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Conditions for or limitations to the validity of this certificate:

- N/A

## Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Revision	Date of Issue certificate	Certification Notice Reference	Action
0	2 November 2022	93928CN46	First issue 2262711CE02
1			
2			

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