

## **Quality System Approval Certificate**Medical Devices Directive 93/42/EEC

The National Standards Authority of Ireland as a duly designated Notified Body, (identification number **0050**), for the purposes of the European Communities (Medical Devices) Regulations (S.I. No. 252 of 1994)

APPROVES THE QUALITY SYSTEM APPLIED BY

## **Smooth-Bor Plastics**

23322 del Lago Dr Laguna Hills CA 92653 USA

to the Product Family

## **SmoothBor Plastics - Air Supply Tubing**

**GMDN Code: 61833** 

on the basis of examination under the requirements of Directive 93/42/EEC on Medical Devices, Annex II, excluding (4)

The use of the NSAI Notified Body identification number **0050** in conjunction with CE Marking of Conformance for this product family is hereby authorised.

Registration Number: 252.1238
Original Approval: 12 May 2021
Last Amended on: 12 May 2021
Remains valid until: 26 May 2024

**Signed:** 

Approved by: Dr. Caroline Dore Geraghty Director, Medical Devices Approved by: Dr. Elaine Darcy

European Medical Device Operations Manager

This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner.

Details of the operational locations included within the scope of this approval can be obtained from NSAI

In the case of a Class III device, this certificate must be supported by a valid design examination certificate

National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.