



Product Service

## Confirmation Statement on validity of EC Certificate (MDD)

pursuant to Directive 93/42/EEC concerning medical devices

**No. GCQ 070143 0005 Rev. 00**

**Manufacturer:** **SAN-O-SUB MBB S.r.l.**  
Via L. da Vinci, 168  
20090 Trezzano sul Naviglio (MI)  
ITALY

**This Confirmation Statement is only valid in combination with the following EC Certificate (MDD):** **G1 070143 0004 Rev. 00**

This Confirmation Statement confirms the validity of the aforementioned EC Certificate (MDD). It considers clarification of scope statements, scope reductions and changes to the manufacturer data initiated 26 May 2021 or later. The conditions laid down in Article 120 (3) of Regulation (EU) 2017/745 on medical devices for placing devices on the market and putting into service apply.

**Report No.:** ITA1934960

**Valid until:** 2024-05-26

Christoph Dicks  
Head of Certification/Notified Body

**Issue Date:** 2022-07-22



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pursuant to Directive 93/42/EEC concerning medical devices

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**Product Category(ies):** Pressure regulators, pressure regulators with integrated cylinder valves, flowmeters, humidifiers for medical gases.

**Description of Change:**

Change in the property of the organization and change of legal name  
from SAN-O-SUB ITALIA s.r.l.  
to SAN-O-SUB MBB s.r.l.