

Medical Device Full Quality Assurance System Certificate  
GB23/00000089



The management system of

## Armstrong Medical Ltd

Wattstown Business Park Newbridge Road Coleraine N. Ireland BT52 1BS United Kingdom

has been assessed and certified as meeting the requirements of

**Part II of The Medical Devices Regulations 2002, Annex II excluding  
section 4 [as modified by Part 2 of Schedule 2A to The Medical  
Devices Regulations 2002]**

For the following products

**The Scope of Registration appears on page 2 of this certificate**

This certificate is valid from 04 August 2025 until 12 May 2028 and remains valid subject to satisfactory surveillance audits.

Issue 7. Certified since 17 February 2023

Authorised by  
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## Armstrong Medical Ltd

### Part II of The Medical Devices Regulations 2002, Annex II excluding section 4 [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

Issue 7

Anaesthetic Circuits;  
Ventilator Circuits;  
CPAP Circuits;  
Resuscitation Sets;  
Humidification Chambers,  
Breathing Filters;  
Catheter Mounts;  
Respiratory Tubing;  
Respiratory Face Masks;  
Adapters; Air Valves;  
Water Traps;  
Breathing Bags; Port Caps;  
Gas Sampling Lines;  
Carbon Dioxide Absorbents;  
Absorbent Canister Adapters,  
Non sterile Laryngeal Airways.  
Nasal Interfaces,  
Gas Flow Driver

Where the above scope includes class III medical device(s), a valid Design Examination Certificate according to Annex II (Section 4) [as modified by Part 2 of Schedule 2A of The MDR 2002] is a mandatory requirement for each device in addition to this certificate to place that device on the market

Certification is based on reports numbered GB/PC/08349

Previous certificate number: N/A

Change in between this certificate and previous one: Addition of Non sterile Laryngeal Airways to the scope.

