



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Ambu A/S

Baltorpbakken 13

Ballerup DK-2750 Denmark

Facility ID Number: F000032

Holds Certificate No: MDSAP 692286

Statement of Conformity: The company listed on this certificate has been audited to and found to conform with the following criteria: ISO 13485:2016 and Australia - Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure; Brasil - RDC ANVISA n. 16/2013, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009; Canada - Medical Devices Regulations - Part 1 - SOR 98/282; Japan - MHLW Ministerial Ordinance 169, Article 4 to Article 68, PMD Act; USA - 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

Please see scope page.

Gary C Stade

For and on behalf of BSI:

Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2019-02-07 Effective Date: 2021-06-19 Expiry Date: 2024-06-18

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MEDICAL DEVICE SINGLE AUDIT PROGRAM
BSI Group America Inc. is an MDSAP authorized auditing organization

...making excellence a habit."

Certificate No: MDSAP 692286

Registered Scope:

Design, development and manufacture of PEEP valves; manual resuscitators and face masks; ventilation upper airway pressure gauges; intubation devices; sterile and non-sterile airway management devices; extrication collars; fixation tape; disposable sterile and non-sterile electrodes and EEG, EMG, and ECG cables. Sterile disposable endoscopes for airways management, gastrointestinal endoscopy and for cystoscopy and endoscope monitor units; clip connector for ECG electrodes. Sterile IVD Sample Container. The distribution of CO2 detectors.



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