



## MANUFACTURER'S DECLARATION OF CONFORMITY AUSTRALIAN THERAPEUTIC GOODS (MEDICAL DEVICES) REGULATIONS 2002 DECLARATION OF CONFORMITY PROCEDURES

SAP DIR No.: 80021366 Version: This is a declaration of conformity made under clause 6.6 of Schedule 3 to the Therapeutic Goods (Medical Devices) Regulations 2002. Manufacturer's name: Welch Allyn, Inc. Business address: 4341 State Street Road Skaneateles Falls, NY 13153-0220 U.S.A. Product name: Fiber-optic Laryngoscope Blades 901038 LARYNGOSCOPE REF 68060, 68061, 68062, 68063, 68064, 68065, 69061, 69062, 69063, 69064, 69211, 69212, 69213, 69214. Classification: GMDN code and term: 46827 - Laryngoscope blade, reusable Scope of application: **A11** Each kind of medical device to which the technical documentation applies complies with the applicable

provisions of the essential principles and the classification rules before being supplied.

| Standards applied: | EN ISO/ISO 13485 | 2012/2ED 2003  | MEDICAL DEVICES - QUALITY MANAGEMENT SYSTEMS - REQUIREMENTS FOR REGULATORY PURPOSES  |
|--------------------|------------------|--|--|
|                    | EN ISO/ISO 14971 | 2012/2ED 2007<br>CORRECTED   | MEDICAL DEVICES - APPLICATION OF RISK MANAGEMENT TO MEDICAL DEVICES  |
|                    | EN ISO/ISO 7376  | 2009/2ED 2009  | ANAESTHETIC AND RESPIRATORY EQUIPMENT - LARYNGOSCOPES FOR TRACHEAL INTUBATION  |
|                    | EN/IEC 60601-1   | 1990, AMD 1, AMD 2,<br>AMD 11, AMD 12, AMD<br>13/2ED 1988, AMD 1 1991,<br>AMD 2 1995 | MEDICAL ELECTRICAL EQUIPMENT - PART 1:<br>GENERAL REQUIREMENTS FOR BASIC<br>SAFETY AND ESSENTIAL PERFORMANCE   |
|                    | EN/IEC 60601-1   | 2006 AMD 11 2011, AMD<br>1 2013/3ED 2005, AMD 1<br>2012                              | MEDICAL ELECTRICAL EQUIPMENT - PART 1:<br>GENERAL REQUIREMENTS FOR BASIC<br>SAFETY AND ESSENTIAL PERFORMANCE   |
|                    | EN/IEC 60601-1-2 | 2007/3ED 2007  | MEDICAL ELECTRICAL EQUIPMENT - PART 1-<br>2: GENERAL REQUIREMENTS FOR BASIC<br>SAFETY AND ESSENTIAL PERFORMANCE -<br>COLLATERAL STANDARD:<br>ELECTROMAGNETIC COMPATIBILITY -<br>REQUIREMENTS AND TESTS |



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| SAP DIR No | .: 80021366      | Version:      | A   |
|------------|------------------|---------------|---|
| F          | EN/IEC 60601-1-6 | 2004/1ED 2004 | MEDICAL ELECTRICAL EQUIPMENT - PART 1-<br>6: GENERAL REQUIREMENTS FOR BASIC<br>SAFETY AND ESSENTIAL PERFORMANCE -<br>COLLATERAL STANDARD: USABILITY |
| F          | EN/IEC 60601-1-6 | 2010/3ED 2010 | MEDICAL ELECTRICAL EQUIPMENT - PART 1-<br>6: GENERAL REQUIREMENTS FOR BASIC<br>SAFETY AND ESSENTIAL PERFORMANCE -<br>COLLATERAL STANDARD: USABILITY |
| I          | STA 1A           | 2001          | PACKAGED-PRODUCTS 150 LB (68 KG) OR<br>LESS   |

Authorised Signatory:

2016-03-

Skaneateles Falls, NY USA

Kevin Crosson, Director Global Regulatory Affairs Date

This authorisation is given in the signatory's capacity as representative of the "Manufacturer" (as recorded on page 1 of this declaration)