

SAP DIR No.: 80017159

Version: K

We declare, under our sole responsibility, that the product listed below conforms to the provisions of:

- the European Council Directive 93/42/EEC of 14 June 1993 concerning medical devices
- the Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS)

Manufacturer's Name and Business Address: Welch Allyn, Inc.
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Skaneateles Falls, NY 13153, USA



Regulatory Affairs Representative
Welch Allyn Limited
Navan Business Park
Dublin Road
Navan, County Meath
Republic of Ireland

Product Name: Ophthalmoscopes



901081 OPHTHALMOSCOPE, STANDARD
901082 OPHTHALMOSCOPE, POCKET



11470F, 11710, 11710F, 11720, 11720F, 11720-L, 11720-LF, 11720R, 11721, 11721F, 11722, 11723, 11724, 11730, 11730F, 11730-R, 11731, 11732, 11735, 11735F, 11736, 11736F, 11750, 11750-VBI, 11770, 11770-BI, 11772-BI, 11772-VCI, 11772-VCL, 11790, 11792-SC, 11796-SC, 12800, 12811, 12812, 12820, 12821, 12831, 12850, 12851, 12860, 12861, 12870-BLK, 12870-BLU, 12870-PUR, 12870-WHT, 12880-BLK, 12880-BLU, 12880-PUR, 12880-WHT, 13000, 13010, 19090, 19091, 19092, 19093, 19190

Annex: VII

Classification: I

Classification Rules: 12

GMDN Code and Term: 12817 - Ophthalmoscope, direct

UMDNS Code and Term: 12817 - Ophthalmoscope, direct



DECLARATION OF CONFORMITY

(in accordance with ISO/IEC 17050-1)

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Standards Applied:	EN/IEC 60601-1	Medical Electrical Equipment – General Guidelines for Safety
	EN/IEC 60601-1-2	Medical electrical equipment -- Part 1-2: General requirements for safety - Collateral standard: Electromagnetic compatibility - Requirements and tests
	EN/IEC 60601-1-6	Medical electrical equipment -- Part 1-6: General requirements for safety - Collateral standard: Usability
	EN/IEC 62366	Medical Devices – Application of Usability Engineering to Medical Devices
	EN/ISO 15004-1	Ophthalmic Instruments – Fundamental Requirements and Test Methods – Part 1: General Requirements Applicable to All Ophthalmic Instruments
	EN/ISO 15004-2	Ophthalmic Instruments – Fundamental Requirements and Test Methods – Part 2: Light Hazard Protection
	EN/ISO 10942	Ophthalmic Instruments - Direct Ophthalmoscopes
	EN/IEC 62471	Photobiological Safety of Lamps and Lamp Systems
	EN ISO 10993-1	Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process
	EN 50581	Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances

Authorised Signatory:

Fiona Butler, EU Manager Regulatory Affairs

Date

Navan

Place of Issue