DECLARATION OF CONFORMITY

(in accordance with ISO/IEC 17050-1)

SAP DIR No.: 80016490 Version: G

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. 4341 State Street Road Skaneateles Falls, NY 13153, USA
EC REP	Regulatory Affairs Representative Welch Allyn Limited Navan Business Park Dublin Road Navan, County Meath Republic of Ireland
Product Name:	Retinoscope
REF	901024, RETINOSCOPE
#	18240, 18245, 18300, 18210-BI, 18342-VC
Medical Device Conformity Assessment Route Annex:	VII
Medical Device Classification:	I
Medical Device Classification Rules:	12
GMDN Code and Term:	32712 - Retinoscope, battery-powered
UMDNS Code and Term	17840 - Retinoscope

Standards Applied:	EN 50581	Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances
	EN 60601-1	Medical Electrical equipment – Part 1: General requirements for Safety

WelchAllyn® DECLARATION OF CONFORMITY

(in accordance with ISO/IEC 17050-1)

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EN 60601-1-2	Medical electrical equipment – Part 1-2: General requirements for safety – Collateral Standard Safety requirements for medical electrical systems

Authorised Signatory:

Fiona Butler 2016-01-07 Navan

Fiona Butler, Manager Regulatory Affairs {EU Authorised Representative} | Date | Place of Issue