

**DECLARATION OF CONFORMITY**

(in accordance with ISO/IEC 17050-1)

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
<b>EC REP</b>	Regulatory Affairs Representative Welch Allyn Limited Navan Business Park Dublin Road Navan, County Meath Republic of Ireland
Product Name <sup>1,3</sup> :	Retinoscope
<b>REF</b> <sub>1,3</sub>	901024, RETINOSCOPE
<b>#</b> <sub>1,3</sub>	18240, 18245, 18300, 18342-VC, 18344-V
Radio equipment <sup>2</sup> :	Not Applicable
Object of the declaration <sup>2</sup> :	Not Applicable
Accessories and components <sup>2</sup> :	Not Applicable
Medical Device Conformity Assessment Route Annex <sup>1</sup> :	VII
Medical Device Classification <sup>1</sup> :	I
Medical Device Classification Rules <sup>1</sup> :	12
GMDN Code and Term <sup>1</sup> :	32712 - Retinoscope, battery-powered

<sup>1</sup> applicable to the medical devices directive, 93/42/EEC

<sup>2</sup> applicable to the radio equipment directive, 2014/53/EU

<sup>3</sup> applicable to the RoHS directive, 2011/65/EU

UMDNS Code and Term<sup>1</sup>: 17840 - Retinoscope

Standards Applied (Standards are applicable to the medical device directive, unless otherwise indicated):	Number	Title
	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
	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*

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Fiona Butler, Manager Regulatory Affairs  
{EU Authorised Representative}

*2019-07-25*

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Date

*Navan*

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Place of Issue

<sup>1</sup> applicable to the medical devices directive, 93/42/EEC

<sup>2</sup> applicable to the radio equipment directive, 2014/53/EU

<sup>3</sup> applicable to the RoHS directive, 2011/65/EU