## Welch Allyn Declaration of Conformity

(in accordance with ISO/IEC 17050-1)

SAP DIR No .:

80020006

Version:

B

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 1999/5/EC of the European Parliament and of the Council of 9 March 1999 on radio equipment and telecommunications terminal equipment and the mutual recognition of their conformity
- 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

Welch Allyn, Inc.

Name and

4341 State Street Road

Business Address:

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:

Welch Allyn Spot Vision Screener

REF

901029 Vision Screener

#

VS100

Medical Device

Conformity

Assessment Route

Annex:

Medical Device

I(m)

II

Classification:

Medical Device

Class I according to Annex IX, Rule 12 with measurement function (m)

Classification

Rules:

GMDN Code and

46390 Visual Screening Analyser

Term:

**UMDNS** Code

46390 Visual Screening Analyser

and Term

Notified Body:

DQS Medizinprodukte GmbH,

(CE 0297)

August-Schanz-Str.21, 60433 Frankfurt am Main

EC-certificate No. 314505 MR2

## WelchAllyn\* DECLARATION OF CONFORMITY (in accordance with ISO/IEC 17050-1)

SAP DIR No.:	80020006	Version:	В
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.
	EN 60601-1-2	2	Medical electrical equipment – Part 1-2: General requirements for safety – Collateral Standard Safety requirements for medical electrical systems.
	ISO 15004-2		Ophthalmic instruments Fundamental requirements and test methods: Part 2:Light hazard Protection

Authorised Signatory:

Fiona Butler, Manager Regulatory Affairs {EU Authorised Representative}

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Place of Issue