

SAP DIR No.: 80016468

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 13153-0220
U.S.A



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

Product Name: KleenSpec® Vaginal Specula



901071 Vaginal Speculum



58600, 58601, 59000, 59001, 59004, 59005, 59006, 58000S, 58001S, 58004S,
59000-B, 59001-B, 59004-B, 590XS, 590XS-B

Annex: VII

Classification: I

Classification Rules: 5 and 12

GMDN Code and Term: 37468 Vaginal Speculum, single use

UMDNS Code and Term: 13666 Speculum, Vaginal

Standards Applied:	EN 50581	Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances
	IEC 60601-1	Medical Electrical Equipment – Part 1: General Requirements for Basic Safety and Essential Performance
	IEC 60601-1-2	Medical Electrical Equipment – Part 1-2: General Requirements for Safety – Collateral Standard: Electromagnetic Compatibility



DECLARATION OF CONFORMITY

(in accordance with ISO/IEC 17050-1)

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IEC 62366	Medical devices – Application of usability engineering to medical devices
IEC 60601-1-6	Medical Electrical Equipment – Part 1-6: General Requirements for Safety – Collateral Standard: Usability
EN ISO 10993-1	Biological Evaluation of Medical Devices - Part 1: Evaluation and Testing

Authorised Signatory:

Fiona Butler, Manager Regulatory Affairs
{EU Authorised Representative}

2014-07-09

Date

Navan

Place of Issue