DECLARATION OF CONFORMITY

(in accordance with ISO/IEC 17050-1)

SAP DIR No.:

80017196

Version:

C

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.

Manufacturer's

Welch Allyn, Inc.

Name and

4341 State Street Road

Business Address:

Skaneateles Falls, NY 13153

U.S.A.

EC REP

Regulatory Affairs Representative

Welch Allyn Limited Navan Business Park

Dublin Road

Navan, County Meath Republic of Ireland

Product Name:

Ear Wash System

REF

29350, 29360, 293600, 29350CAN-D, 29350C-I, 29350DAN-I, 29350D-I, 29350E-I, 29350E-ID, 29350FCAN-D, 29350F-I, 29350G-I, 29350I-I,

29350P-I, 29350P-ID, 29350S-D, 29350S-I, 29350S-ID, 29350SW-I,

293600SCHP, 29352SCHP and 29360SCHP.

Annex:

VII

Classification:

I

Classification

5

Rules:

GMDN Code and

35025 - Syringe, irrigating, ear

Term:

35348 - Speculum, ear

UMDNS Code

23526 - Irrigators, Ear Canal

and Term

Standards Applied:

EN ISO 14971

Medical Devices - Application of Risk Management to

Medical Devices

EN ISO 10993-1

Biological Evaluation of Medical Devices - Part 1:

Evaluation and Testing within a Risk Management Process

Authorised Signatory:

Fiona Butler, Manager Regulatory Affairs

{EU Authorised Representative}

Date Navan
Place of Iss