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## **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 2014/53/EU of the European Parliament and of the Council of 16 April 2014 on the harmonization of the laws of the Member States relating to the making available on the market of radio equipment
- 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)

(10113)					
Manufacturer's Name and Business Address:	Welch Allyn, Inc. 4341 State Street Road Skaneateles Falls, NY				
EC REP	Regulatory Affairs Representative Welch Allyn Limited Navan Business Park, Dublin Road Navan, Co. Meath C15 AW22 Ireland				
Product Name <sup>1,3</sup> :	ELI380				
<b>REF</b> 1,3	The ELI380 comes in different configurations: ELI380-XYYZZ Where "X" and "Y" can be a letter from A to Z whereas "Z" can be a number from 0 to 9 representing the following:				
	Base ELI380- X	Connectivity Y	Option Y	Cable Z	Lead Set Z
# 1,3	901133 ELECTROCA	RDIOGRAPH	I		
Radio equipment <sup>2</sup> :	Laird WB45NBT (DIR 60086189)				
Object of the declaration <sup>2</sup> :	802.11 a/b/g/n Enterprise Wi-Fi + Bluetooth Communications Subsystem				
Radio equipment <sup>2</sup> :	B&B 9373 - WLNN-SP (DHF-THF-89515-00)	-MR551			

<sup>&</sup>lt;sup>1</sup> applicable to the medical device directive, 93/42/EEC

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

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



Object of the declaration <sup>2</sup> :	802.11 a/b/g/n WLAN				
Accessories and components <sup>2</sup> :	Not Applicable				
Medical Device Conformity Assessment Route Annex <sup>1</sup> :	II				
Medical Device Classification <sup>1</sup> :	IIa	IIa			
Medical Device Classification Rules <sup>1</sup> :	10				
GMDN Code and Term <sup>1</sup> :	16231 - Electrocardiograph, professional, multichannel				
Notified Body <sup>1</sup> : (CE 0459)	LNE G-MED, 1, rue Gaston Boissier 75015 Paris France EC-certificate No. 35913				
Standards Applied	Number	Title			
(Standards are applicable to the medical device	EN 50581 <sup>3</sup>	Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances			
directive, unless otherwise	EN/IEC 62304 <sup>1</sup>	Medical Device Software – Software Life Cycle Processes			
indicated):	EN/IEC 60601-1 <sup>1</sup>	Medical Electrical Equipment – Part 1: General Requirements for Basic Safety and Essential Performance			
	EN/IEC 60601-1- 2 <sup>1</sup>	Medical electrical equipment - Part 1-2: General Requirements for Safety – Collateral Standard: Electromagnetic Compatibility – Requirements and Tests			
	EN/IEC 62366 <sup>1</sup>	Medical devices – Application of Usability Engineering to Medical Devices			
	EN/IEC 60601-1-6 <sup>1</sup>	Medical Electrical Equipment – Part 1-6: General Requirements for Safety – Collateral Standard: Usability			
	EN/IEC 60601-2- 25 <sup>1</sup>	Medical Electrical Equipment. Part 2 25: Particular requirements for the basic safety and essential requirements of electrocardiographs			
	ETSI EN 301 489 -1 <sup>2</sup>	Electromagnetic compatibility and Radio Spectrum Matters, Part1: Common technical requirements			

applicable to the medical device directive, 93/42/EEC
 applicable to the radio equipment directive, 2014/53/EU
 applicable to the RoHS directive, 2011/65/EU

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	ETSI EN 300 440 -2 <sup>2</sup>	Electromagnetic compatibility and Radio Spectrum Matters
1	ETSI EN 300 328 <sup>2</sup>	Electromagnetic compatibility and Radio Spectrum Matters, Wideband transmission systems; Data transmission equipment operating in the 2.4 GHz ISM band and using wide band modulation techniques

Authorised Signatory:

Mark Elliott,

Director Quality Assurance

h Allit 07-007-2019
Date

Milwaukee, Wisconsin, USA Place of Issue

 $<sup>^{1}</sup>$  applicable to the medical device directive, 93/42/EEC  $^{2}$  applicable to the radio equipment directive, 2014/53/EU  $^{3}$  applicable to the RoHS directive, 2011/65/EU



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Document Change History

Version	Description	Author	Date
F	Initial Release	Marco Manduchi	2019-08-16
		Steven Co	
		Michael Lippold	
G	Updated EC Rep Address	Michael Lippold	2019-10-01

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