

EC Declaration of Conformity

We, manufacturer **Widex A/S**
Nymoellevej 6
DK-3540 Lyngø
Denmark

Declare under our sole responsibility that the following **PRODUCTS** starting with Serial no. 10.000

Brand: **Vogue**
 Description: **Hearing Aid**

Model	Variant(s)	Type	GMDN code	GMDN Term
VPA E	RC	RIC	47169	Receiver-in-canal air-conduction hearing aid
VF2 E	T-RC	RIC, RITE		
VFS E	T-RC			
VXP E	T-RC	ITE	34672	In-the-ear air-conduction hearing aid
VCIC E	RC-R, RC-L	CIC	41209	Canal air-conduction hearing aid
VCIC E TR	RC-R, RC-L			

are in conformity with the essential requirements and other applicable provisions of the following **EU Directives**:

Council Directive 93/42/EEC as amended by Dir. 2007/47/EC (MDD)
Directive 2014/53/EU (RED)
Directive 2011/65/EU (RoHS 2)

Conformity assessment procedure	MDD : Annex II of 93/42/EEC RED : Annex II of 2014/53/EU
Notified Body	MDD : TÜV SÜD Product Service GmbH, Notified Body No.: NB 0123 Ridlerstr. 65, 80339 München, Germany
EC-Certificate	MDD : No. G1 093461 0011
Classification of Device	MDD : Class IIa, Rule 9 according to 93/42/EEC Annex IX
Applied standards and specifications in conformity assessment.	MDD : EN 1041, EN 10993-1, EN 13485, EN 14155, EN 14971, ISO 15223-1, EN 50419, EN 60118-0, EN 60118-6, EN 60118-7, EN 60118-13, EN 60601-1-2, EN 62304, EN 62366-1
Standard versions valid on the date when this DoC is issued.	RED : EN 60950-1, EN 62479, EN 301 489-1, EN 301 489-3, EN 301 489-17, EN 300 330, EN 300 328 RoHS 2 : EN 50581, EN 62321

Lyngø, 24-09-2020
 Place and date of issue


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