EC Declaration of Conformity

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

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

Brand: Widex
Series: MENU
Description: Hearing Aid

Model	Variant(s)	Type	GMDN Code	GMDN Term
ME-SP	T-VC-RC	BTE	34671	Behind-the-ear air-conduction hearing aid

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 2011/65/EU (RoHS 2)

Conformity assessment procedure	MDD	: Annex II of 93/42/EEC
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. Standard versions valid on the date when this DoC is issued.	MDD RoHS 2	: EN 1041, EN 10993-1, EN 13485, EN 14155, EN 14971, EN 15223-1, EN 50419, EN 60118-0, EN 60118-6, EN 60118-7, EN 60118-13, EN 62366 : EN 50581, EN 62321

Lynge, 18-08-2020
Place and date of issue

Director, Global Regulatory Affairs — Kristine Klitgaard Pedersen