WIDEX

EU DECLARATION OF CONFORMITY

Manufacturer	WSAUD A/S Nymoellevej 6 DK-3540 Lynge Denmark
Brand:	WIDEX
Product Family:	MOMENT
Type of Device:	Hearing Aid
Basic UDI-DI:	5714880-WSA-65-15-5A
Single registration number:	DK-MF-000015974
GMDN Code:	47169 Receiver-in-canal air-conduction hearing aid
EMDN Code:	Y2145060102
Product Identification:	See below

We declare under our sole responsibility that above products are in conformity with the following Regulations and Directives:

REGULATION (EU) 2017/745 (MDR) OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

Conformity assessment procedure:	Annex IX of Regulation (EU) 2017/745
Notified Body:	TÜV SÜD Product Service GmbH, Notified Body No.: NB 0123 Ridlerstr. 65, 80339 München, Germany
Classification of device:	Class IIa (according to Annex VIII Rule 9 to Regulation (EU) 2017/745)

The products meet all applicable standards and the general safety and performance requirements of the Regulation (EU) 2017/745 Annex I. Applicable standards are listed in the respective technical documentation.

Council Directive 2011/65/EU (RoHS) as amended by Dir. 2017/2102/EC (RoHS2)

Relevant Harmonized Standards:	EN IEC 63000

Council Directive 2014/53/EU (RED)

Relevant Harmonized Standards: Standard versions as listed in the respective technical documentation.	EN 62479, EN 301 489-1, EN 301 489-3, EN 301 489-17, EN 300 330, EN 300 328

Product Identification	Type of Device
MRR4D (-, 440, 330, 220, 110, DEMO)	RIC (Receiver In the Canal) Hearing Aid

This Declaration of Conformity includes all hearing aid components and spare parts of the products listed above.

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Place and valid from date	Lynge, June 22, 2022
Name	Hans-Otto Bindeballe Global Regulatory Affairs Manager
Signature	

This declaration will be renewed on any significant change of product, product range, standards and laws.