



EC Certificate

EC Design-Examination Certificate Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD), Annex 2 (4) (Other devices than custom made or intended for clinical investigation)

No. I7 017853 0155 Rev. 00

Manufacturer: MED-EL

Elektromedizinische Geräte GmbH

Fürstenweg 77A 6020 Innsbruck **AUSTRIA**

Product: External components for Cochlear Implant

Systems

and Auditory Brainstem Implant System

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with AIMDD Annex 2 (4). This design of the devices conforms to the requirements of this Directive. For marketing of these devices an additional Annex 2 certificate is mandatory. See also notes overleaf.

Report no.: 713175392

Valid from: 2020-05-07 Valid until: 2024-05-26

2020-05-07 Date.

Christoph Dicks

Head of Certification/Notified Body



EC Certificate

EC Design-Examination Certificate Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD), Annex 2 (4) (Other devices than custom made or intended for clinical investigation)

No. I7 017853 0155 Rev. 00

Model(s):

RONDO 3

Product:

External Components for Cochlear Implant Systems and Auditory Brainstem Implant System

Test Report No.:

713175392

Model:

RONDO 3 Audio Processor

Components:

Processor Unit -RONDO 3 Processor Unit (Me1550, Me1551, Me1552, Me1553)

External Power Supply Options

- -Mini Battery Pack
- -Mini Battery Pack Cable for RONDO 3
- -Charging Cable

External Cables

-MAX Programming Cable for RONDO 2

Covers

- -RONDO 3 Cover
- -RONDO 3 Mini Cover

External Adapter

-Telecoil Adapter

Page 2 of 2 TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123