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Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
www.zlgm.de  
BS-MDR-099



Product Service

## EU Technical Documentation Assessment Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapter II  
(Implantable Class IIb Devices and Class III Devices)

**No. G70 078611 0123 Rev. 01**

### Manufacturer:

**Cochlear Limited**

1 University Avenue  
Macquarie University NSW 2109  
AUSTRALIA

### Authorized Representative:

Cochlear Deutschland GmbH & Co. KG  
Karl-Wiechert-Allee 76A, 30625 Hannover, GERMANY

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has drawn up and presented a Technical Documentation according to Annex II and III of the Regulation (EU) 2017/745 on medical devices. Details on devices covered by the Technical Documentation are described on the following page(s). The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The technical documentation assessment included an assessment of the clinical evaluation assessment. The conformity assessment has been carried out according to Annex IX chapter II of this regulation with a positive result.

Changes to the approved device, where such changes could affect the safety and performance of the device or the conditions prescribed for use of the device, shall require approval from the notified body TÜV SÜD Product Service GmbH. In order to place the devices on the market with CE-marking, an EU Quality Management System Certificate pursuant to Annex IX chapters I and III is necessary in addition to this EU Technical Documentation Assessment Certificate. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with.

For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G70 078611 0123 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:G70 078611 0123 Rev. 01)

**Report No.:** 713198997

**Preceding Certificate No.:** G70 078611 0123 Rev. 00

**Valid from:** 2021-05-18

**Valid until:** 2026-01-21

**Date of Initial Issuance:** 2021-01-22

Christoph Dicks  
Head of Certification/Notified Body

**Issue date:** 2021-05-18



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Classification:	III						
Device Group:	J0380 - AUDITORY ACTIVE-IMPLANTABLE DEVICES - ACCESSORIES						
Basic UDI-DI:	9321502CP1000PU3T						
Intended Purpose:	<p>The processing unit is intended to be used in combination with other devices as part of a hearing implant system to provide hearing sensation. The processing unit converts sounds into electrical signals, which it sends, via a coil, to an implant. The processing unit also provides power to the implant.</p> <p>When used in combination with an audio receiver, the sound processor also delivers sound to the ear canal in recipients with residual hearing.</p>						
Device(s):	<p>Nucleus® 7 Processing Unit, model CP1000. Available in the following variants:</p> <p>Nucleus® 7 Processing Unit, CP1000 - Black, Platinum Detail</p> <p>Nucleus® 7 Processing Unit, CP1000 - Brown</p> <p>Nucleus® 7 Processing Unit, CP1000 - Grey</p> <p>Nucleus® 7 Processing Unit, CP1000 - Sand</p> <p>Nucleus® 7 Processing Unit, CP1000 - Black, Golden Detail</p> <p>Nucleus® 7 Processing Unit, CP1000 - White</p>						
The validity of this certificate depends on conditions and/or is limited to the following:	-None-						
Revision History:	<table><tr><td>Rev.</td><td>Dated</td><td>Report</td></tr><tr><td>00</td><td>2021-01-22</td><td>713184969</td></tr></table>	Rev.	Dated	Report	00	2021-01-22	713184969
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00	2021-01-22	713184969					