Declaration of Conformity

Legal Manufacturer:	Cochlear Limited 1 University Avenue Macquarie University NSW 2109 Australia
EC Representative:	Cochlear Deutschland GmbH & Co. KG Karl-Wiechert-Allee 76A 30625 Hannover Germany
Product:	Refer to Appendix I
GMDN Code & Term	47374 - Cochlear implant system sound processor
Classification:	Active Implantable Medical Device (AIMD)

I herewith declare that the products listed in Appendix I conform to the provisions of:

- The Active Implantable Medical Devices Directive 90/385/EEC Annex 1 and the conformity assessment route of Annex 2 including Part 4:
  - **Notified Body**: TÜV SÜD Product Service GmbH
Ridlerstraβe 65, 80339 München
Germany
  - **Notified Body No**: 0123

AND

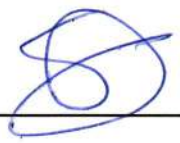
- Council Directive 2014/53/EU on Radio Equipment (RED) and the conformity assessment route of Annex II.

AND

- Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment.

I declare that Cochlear Limited is exclusively responsible for this Declaration of Conformity.

Authorised Signatory:



Date: 08 MAY 2020.

Steven Kennedy

Global Head of Regulatory Affairs

Place: Sydney, Australia

Valid to: 26May2024

APPENDIX I

Product	Reference No.
Kanso® 2 Sound Processor - Black	P1320278
Kanso® 2 Sound Processor - Chocolate Brown	P1320276
Kanso® 2 Sound Processor - Sandy Blonde	P1320274
Kanso® 2 Sound Processor - Silver	P1320275
Kanso® 2 Sound Processor - Slate Grey	P1320277