



## Declaration of Conformity

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**Manufacturer:**

ResMed Pty. Ltd.  
1 Elizabeth Macarthur Drive  
Bella Vista  
NSW 2153  
Australia

**Authorized Representative:**

ResMed SAS  
Parc Technologique de Lyon  
292 Allée Jacques Monod  
69791 Saint Priest Cedex  
France

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**Product:** Air10 Oximeter Adapter

**Intended Use:**

The Air10 Oximeter Adapter is intended to be used with AirSense™ 10 / AirCurve™ 10 / Lumis™ devices. The oximeter adapter connects a NONIN Xpod™ oximeter to a device. It is intended for home and hospital use.

**Classification:** I according to Rule 13

**EMDN:** Z1203019080 Various Instruments for Anaesthesia and Pulmonary Ventilation Support - Hardware Accessories

**Conformity Assessment Route:** Annex II and III, Regulation EU 2017/745

**Basic UDI-DI:** 619498EC1326A

**Common Specification:** N/A

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We herewith declare that the above mentioned products are in conformity with the Council Regulation 2017/745 for medical devices.

Compliance is applicable from the date listed below. All supporting documentation is retained at the premises of the manufacturer. This declaration is issued under the sole responsibility of ResMed Pty. Ltd.

**SRN:** AU-MF-000011753

Signed at Sydney, Australia on: 09 January 2024

DocuSigned by:

*Nicole Wilson*

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Nicole Wilson  
Person Responsible for Regulatory Compliance (PRRC)  
ResMed Pty. Ltd.

**EC132.3**

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