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## Declaration of Conformity

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<b>Manufacturer:</b>	<b>Authorized Representative:</b>	<b>Notified Body:</b>
ResMed Corp. 9001 Spectrum Center Blvd. San Diego CA 92123 USA	ResMed SAS Parc Technologique de Lyon 292 Allée Jacques Monod 69791 Saint Priest Cedex France	TÜV SÜD Product Service GmbH Ridlerstraße 65 80339 München Germany

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**Product:** EasyCare Tx 2

**Intended Use:**

EasyCare Tx 2 software is intended to be used with ResMed compatible therapy devices in a clinical environment. EasyCare Tx 2 provides therapy device setting changes and displays real-time data and treatment settings from the compatible ResMed therapy device when used together with the Tx Link 2 module.

**Classification:** IIa according to Rule 11

**EMDN:** Z1203019092 Various Instruments for Anaesthesia and Pulmonary Ventilation Support - Medical Device Software

**Conformity Assessment Route:** Annex IX (excluding Chapter II), Regulation EU 2017/745

**Basic UDI-DI:** 8401934EC205L7

**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 Corp.

**EC Certificate Number:** G10 083904 0009

**SRN:** US-MF-000011805

Signed at San Diego CA, USA on: 21.01.2026

Signed by:

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Jason Gorman

Director of Global Product Regulatory Affairs  
ResMed Corp.

**EC205.1**

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