# *EU Declaration of Conformity*

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|---|---|
| **Manufacturer Name:** | AescuBrands UG (haftungsbeschränkt) ("AescuBrands") |
| **Manufacturer Address:** | Im Gewerbegebiet 5, 91093 Heßdorf, Deutschland |
| **SRN (Single Registration Number):** | DE-MF-000011001 |
| **Basic UDI-DI:** | 42604428101CAUZ |
| **Name of the Device (s):** | SomnoSept |
| **Classification:** | Class I Device (Annex VIII Regulation (EU) 2017/745) |
| **Conformity assessment route:** | AescuBrands uses the following procedure for the CE-labeling of their products according to Regulation (EU) 2017/745:<br><br>Class 1: EU conformity declaration according to Annex IV |

This declaration of conformity is issued under the sole responsibility of AescuBrands. We hereby declare that the medical devices specified above meet the provisions of the Regulation (EU) MDR 2017/745 for medical devices. All supporting documentation is retained at the premises of the manufacturer.

Signature:

Dr. Daniel Grätz
Managing Director

Place and date of issue:

Heßdorf, 15.12.2021