

EC DECLARATION OF CONFORMITY

Regulation (EU) 2017/745 of the European Parliament and of the Council on
medical devices

Manufacturer:

UAB SOFTNETA
K. Barsausko str. 59B
LT-51423 Kaunas
Lithuania

Product: Stand-alone software medical device

Model: «MedDream»

Types: «MedDream»

Version: 8.3.1

UDI-DI: (01)04779049590105(10)MDSY8310

Notified body: TÜV Rheinland LGA Products GmbH

Class IIb active medical device according to MDR 2017/745 Annex VIII
Chapter III, Rule 11.

We hereby declare that the above mentioned device meets the applicable provisions of Regulation (EU) 2017/745 of the European Parliament and of the Council on medical devices. Route of compliance according Annex IX Chapter I, Section 2 and 3 and Chapter III is applied. Issued certificate: registration No. HZ 1992126-1. All supporting technical documentation is retained at the premises of the manufacturer. Manufacturer is exclusively responsible for the declaration of conformity.

Date of issue:

2023-12-15

Director of Softneta
Vytautas Baublys

Place: Kaunas, Lithuania