EU DECLARATION OF CONFORMITY (EU MDR 2017/745)

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1. Manufacturer

Name: Medbric AS
Registered place of business: Krambugata 2, 7011 Trondheim, Norway
Contact: Jon Espen Ingvaldsen, jonespen@medbric.com

2. Declaration issued under sole responsibility

We, Medbric AS, declare under our sole responsibility that the device(s) covered by this declaration conform(s) to Regulation (EU) 2017/745 on medical devices (MDR).

3. Device(s) covered by this Declaration

Item	Details
Device / Trade name	Medbric Software
Device type	Medical Device Software (MDSW) – Software-only
Configuration / modules included	Transcription + dictation + summarization + chatbot (non-clinical documentation/workflow support)

Item	Details
Software version	v0.1.0
Basic UDI-DI	PP144331912202519
UDI-PI	(1T)v01(16D)251219
Risk class (MDR)	Class I (Rule 11)
Intended users	Healthcare professionals
eIFU location	https://medbric.com/

4. Conformity assessment route

Notified Body involvement: Not applicable (Class I device; no certificate issued).
Notified Body name/number & certificate(s): N/A

5. Union legislation

The device covered by this declaration is in conformity with Regulation (EU) 2017/745 (MDR).

6. Common Specifications (CS)

CS used: None

7. Additional information

This declaration is continuously updated as required.

10. Place and date of issue

Place: Oslo, Norway
Date: 2025-12-19

11. Signature

Name: Jorunn Thaulow
Function: CEO
Signature: