

EC Declaration of Conformity



(MDR 2017/745 (UE))

The Manufacturer: CBS Medical Srl with legal headquarters in Schio (VI) 36015,
via Cappuccini 129 and operating office in Marano Vicentino (VI) 36035, via
dell'Industria, 50

DECLARES

that the product: Bactericidal disposable mat with Biomaster antimicrobial agent,
complies with MDR 2017/745 (UE), concerning medical devices.

DEVICE IDENTIFICATION:

Non-invasive **Class 1** device
RDM Italian Ministry of Health: **1876007**
CND: **V9099**

ID Eudamed APP000016192. REGULATION (EU) 2017/745 OF THE EUROPEAN
PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices

SRN: **IT-MF-000020073**

Basic UDI-ID: 805571310biomaster-mats2L

It is registered as Medical Device (class 1) at the Italian Ministry of Health
(**N.1876007**).

Our products with **Biomaster** antimicrobial are tested according to the standard
ISO 22196: 2011, with Certificates No. 1038185.40 / 13224; 22LA00323; 22LA00324;
22LA00325; 22LA00326; 22LA00327
ISO 21702: 2019, with Certificate No. 20200030/01

Product codes: 3L49BIO; 3L41BIO; 3L69BIO; 3L61BIO; 3L91BIO; 3L115BIO; 3L150BIO;
3L240BIO

Reference our technical sheet: REV-01-2022

This EU declaration of conformity is issued under the sole responsibility of the
manufacturer;

The device that is covered by the present declaration is in conformity with this
Regulation and with any other relevant Union legislation that provides for the issuing of
an EU declaration of conformity.

Date: 13/02/2023

Place: Schio, Italy

Signature

Mirko Nannini – CEO