DECLARATION OF CONFORMITY

Manufacturer	ABIGO Medical AB Vapenvägen 1, SE-696 33 Askersund, Sweden
Device classification and rule (Regulation EU 2017/745 Annex VIII)	IIb, Rule 4
SRN of the Manufacturer:	SE-MF-000000736

Basic UDI-DI: 07392130Sorbact2DU
EMDN: M040416

Intended Purpose: Cutimed® Sorbact Ribbon® is intended for use in management of clean, contaminated, colonized or infected exuding wounds, such as surgical wounds, traumatic wounds, pressure ulcers, diabetic foot ulcers and leg ulcers. Cutimed® Sorbact Ribbon® can be used on superficial and deep wounds such as cavity wounds and fistulas. Cutimed® Sorbact Ribbon® is also intended to treat fungal infections in skin folds (intertrigo).

Trade and Product Name	**Catalogue number (REF)**
Cutimed® Sorbact Ribbon®	Healthcare: 72166-16, 72166-17, 72167-10, 72167-11, 72166-22, 72167-14, 72166-23, 72167-15

Conformity assessment based on a quality management system and on assessment of technical documentation per **Annex IX Chapters I & III of Regulation (EU) 2017/745** has been performed by the following Notified Body:

Name and address	**Notified Body id no**	**EC Certificate no and validity**
Intertek Medical Notified Body AB, Torshamnsgatan 43, Box 1103, SE-164 22 Kista, Sweden	2862	28620115063-03, 28 June 2031

This declaration of conformity is issued under the sole responsibility of ABIGO Medical AB as the manufacturer.
I hereby declare that the above-mentioned devices comply with **Regulation (EU) 2017/745** concerning medical devices.

Askersund, 2026-07-02

Place and date of issue

Marie Skoglund, Regulatory Affairs Director
On behalf of Anna Arvidsson, Managing Director

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