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: 07392130Sorbact1DS
EMDN: M040416

Intended Purpose: Cutimed Sorbact Contact® 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 Contact® can be used on both superficial and deep wounds.

Trade and Product Name	**Catalogue number (REF)**
Cutimed Sorbact Contact®	Healthcare: 72164-30, 72164-31, 72164-32, 72164-33, 72165-60, 72165-61, 72165-62, 72165-63, 72693-11, 72693-12

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	28620115063-02, 28 June 2026

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.

2024-08-12
Mölndal, date of issue

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

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