POHL BOSKAMP

Declaration of Conformity

We, the company
G. Pohl-Boskamp GmbH & Co. KG
Kieler Strasse 11
25551 Hohenlockstedt, Germany
Single Registration Number: DE-MF-000010001

declare on our own responsibility, that the risk class IIa medical device (rule 21, dash 3 of annex VIII of the regulation mentioned below)

GeloSitin® Nasal Care with

Intended Purpose: GeloSitin® Nasal Care is used for physical treatment and moisturization of dry or damaged nasal mucosa.
Product code: 575
Basic-UDI-DI: 40291250032L

which is subject of this declaration, complies with the requirements of

Annex I

and this Declaration of Conformity complies with the

Annex IV

of the REGULATION (EU) 2017/745 of 5th of April, 2017 on medical devices (MDR).
The Notified Body
DNV MEDCERT GmbH
Pilatuspool 2
20355 Hamburg, Germany
with the Identification Number 0482 was involved in the conformity assessment according to Annex IX of the REGULATION (EU) 2017/745 and has issued the certificate no. 3253GB448250403 valid until 06 December 2026.

This Declaration of Conformity is valid until 06 December 2026.

Hohenlockstedt, 17 April 2025

POHL BOSKAMP

G. Pohl-Boskamp GmbH & Co. KG POHL BOSKAMP Pharmaceuticals since 1855 Kieler Str. 11 25551 Hohenlockstedt Germany

I. Rowedder

Ines Rowedder
Person responsible for regulatory compliance (PRRC) acc. to Article 15 MDR

Lea Dornseiff

Lea Dornseiff
Advisor Medical Device Registration