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 I medical device (rule 5, dash 1 of annex VIII of the regulation mentioned below)

GeloMuc

Intended Purpose: Respiratory Therapy Device for the mobilisation of viscous bronchial mucus
Product code: 494
Basic-UDI-DI: 40291250022J

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).

This Declaration of Conformity is valid until 27 July 2031.

Hohenlockstedt, 28 July 2026

POHL BOSKAMP

G. Pohl-Boskamp GmbH & Co. KG
POHL BOSKAMP
Pharmaceuticals since 1845
Kieler Str. 11 25551 Hohenlockstedt Germany

Ines Rowedder Person responsible for regulatory compliance (PRRC) acc. to Article 15 MDR	Lea Dornseiff Advisor Medical Device Registration