



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)

GeloTonsil® Gargle Gel

Intended Purpose: GeloTonsil® Gargle gel is used as a supportive therapy for inflammation of the mucosa of the mouth and throat. GeloTonsil® Gargle Gel relieves sore throat and swallowing difficulties by forming a protective film over the irritated pharyngeal mucosa.

Product code: 585

Basic-UDI-DI: 40291250112K

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

Brooktorkai 18

20457 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.

3253GB448260430 valid until 06 December 2026.

This Declaration of Conformity is valid until 06 December 2026.

Hohenlockstedt, 11 May 2026

POHL BOSKAMP



I. Rowedder

R. Losch

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

Robert Losch
 Advisor Medical Device Registration