



## 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)

### **Saseem<sup>®</sup> – mouth spray**

Intended Purpose: Saseem<sup>®</sup> mouth spray is a medical device that is used for the intensive moisturisation of the oral mucosa when saliva production is impaired and in patients suffering from xerostomia of various aetiologies.

Product code: 660

Basic-UDI-DI: 40291250042N

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

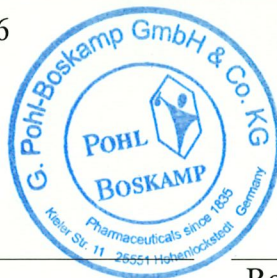
20457 Hamburg, Germany

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, 04 May 2026

POHL BOSKAMP



*I. Rowedder*

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

*R. Losch*

Robert Losch  
Advisor Medical Device Registration