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)

Saseem® – mouth spray

Intended Purpose: Saseem® 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

G. Pohl-Boskamp GmbH & Co. KG Pharmaceuticals since 1835 Kieler Str. 11 25551 Hohenlockstedt Germany

I. Rowedder

R. Gl

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

Robert Losch
Advisor Medical Device Registration