EU Certificate

Quality Management System
REGULATION (EU) 2017/745 on Medical Devices
Annex IX Chapter I, Section 2 and 3 and Chapter III

Registration No.:	HZ 1200911-1
Manufacturer:	Procter & Gamble Technical Centres Limited The Heights, Brooklands Weybridge Surrey KT13 0XP United Kingdom
EUDAMED Single Registration No.:	GB-MF-000017862
Products:	Products of class IIa: Q010280 - Devices for Prosthetic Dentistry – Accessories: - Denture Adhesive Creams
Authorized representative(s):	Procter & Gamble Service GmbH Sulzbacher Str. 40 65824 Schwalbach am Taunus Germany DE-AR-000016615

The Notified Body hereby declares that the requirements of Annex IX, Chapter I, Section 2 and 3 of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.
If class III devices or class IIb implantable devices referred to in the second subparagraph of Article 52(4) are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.9 is required before placing them on the market.

Report No.:	1184800-600
Effective date:	2025-04-21
Expiry date:	2030-04-20
Issue date:	2025-04-16

Dr. Thomas Kießling
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

This certificate can be validated on https://www.certipedia.com

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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Certificate history		
Revision:	Description:	Issue date:
7	Re-certification. Replaces certificate HZ 1200911-1 Rev. 6 issued 2025-03-28	2025-04-16

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