

# 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 1803402-1  
Manufacturer: APOLLO IMPLANT COMPONENTS SP. Z O.O.  
SPÓŁKA KOMANDYTOWA  
ul. Konopna 16  
95-200 Pabianice  
Poland  
EUDAMED Single Registration No.: PL-MF-000023711  
Products: Products of class IIb:  
P01020101 - DENTAL IMPLANTS AND ACCESSORIES  
DENTAL IMPLANTS  
Limited to: prosthetic abutments for dental implants and related screws, healing abutments.  
Authorized representative(s): Not applicable

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2023-12-20

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.: 84958606-60  
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Expiry date: 2028-12-19  
Issue date: 2023-12-20



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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.

