

EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 772747 R000

Manufacturer: AB Dental Germany UG

Address:

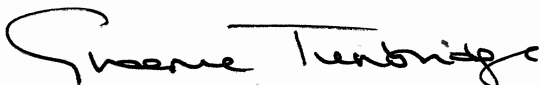
Friedrichstrasse 95
Berlin
10117
Germany

Single Registration Number: DE-MF-000016161

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III devices, and Class IIb implantable devices that are not considered well-established technologies as specified in Article 52(4) an additional Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Medical Devices

First Issue Date: **2023-01-18**

Current Issue Date: **2023-08-24**

Starting Validity Date: **2023-08-24**

Expiry Date: **2028-01-17**

...making excellence a habit.™

EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 772747 R000

Device Schedule: Class III and Class IIb devices

Class IIb, Implantable, Well-established technologies	Intended purpose
Abutments	Intended for use to support single and multiple tooth prosthesis, in the mandible or maxilla. The prosthesis can be screw retained or cement retained to the abutment.
Implant Accessories	Intended for use to connect sleeves, abutments and implants; support single or multiple tooth prostheses; and cover implant and abutments. The accessories can be cement retained to the abutment/sleeves.

Device Schedule: Class IIa, Custom-made and other devices

Device(s)	Risk Classification
Reusable Dental Instruments Connected to an Active Device	Class IIa

First Issue Date: **2023-01-18**

Current Issue Date: **2023-08-24**

Starting Validity Date: **2023-08-24**

Expiry Date: **2028-01-17**

...making excellence a habit.™

EU Quality Management System Certificate

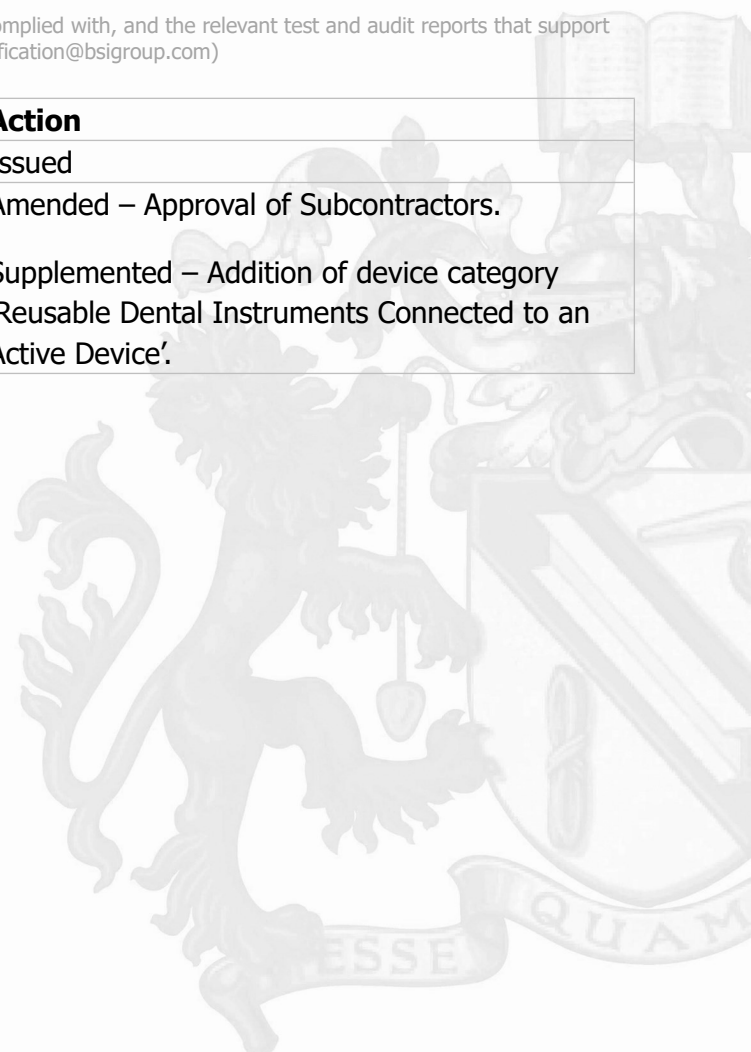
Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 772747 R000

Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)

Date	Reference Number	Action
2023-01-18	3699107	Issued
Current	3913475	Amended – Approval of Subcontractors. Supplemented – Addition of device category 'Reusable Dental Instruments Connected to an Active Device'.



First Issue Date: **2023-01-18**

Current Issue Date: **2023-08-24**

Starting Validity Date: **2023-08-24**

Expiry Date: **2028-01-17**

...making excellence a habit.™

Page 3 of 3

Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.