



EU Quality Management System Certificate

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

MDR 738262 R000

Manufacturer: exocad GmbH

Address: Rosa-Parks-Str.2 64295 Darmstadt, Germany

Single Registration Number: DE-MF-000007341

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 and Class IIb implantable devices an 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: 2022-04-08 Starting Validity Date: 2023-01-11

Current Issue Date: **2023-01-11** Expiry Date: **2027-04-07**

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Page 1 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.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80 Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK. A Member of the BSI Group of Companies.





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

Device(s)	Risk Classification	
Dental implant planning software / Dental restoration design	Class IIa	
software		



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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
2022-04-08	3313725	Issued
Current	3795514	Supplemented - Addition of "Dental implant planning software". Amended – Removal of critical subcontractor/crucial supplier page. Addition of SRN. Change of legal manufacturer address. Amendment of "Dental Restoration Devices" to "Dental restoration design software" in Device Schedule.

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