

Certificate

mdc medical device certification GmbH
certifies that

botiss biomaterials GmbH
Hauptstraße 28
15806 Zossen b. Berlin
Germany

with the location listed in the attachment
for the scope

**development, production, storage, shipment, marketing and distribution of
products for bone and tissue regeneration**

has introduced and applies a

Quality Management System

The mdc audit has proven that this quality management system
meets all requirements of the following standard

EN ISO 13485

Medical devices – Quality management systems –
Requirements for regulatory purposes

EN ISO 13485:2012 + AC:2012 - ISO 13485:2003 + Cor. 1:2009

Valid from	2016-12-19
Valid until	2019-12-19
Registration no.	D1323300024
Report no.	P16-01372-81646
Stuttgart	2016-12-19



Head of Certification Body



Attachment of the certificate

No. D1323300024

date 2016-12-19

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Location	Scope
botiss biomaterials GmbH, Ullsteinstraße 108 Aufgang D, 12109 Berlin	development, production, storage, shipment, marketing and distribution of products for bone and tissue regeneration



A handwritten signature in black ink, appearing to be "J. A.", is positioned above a horizontal line.

Head of Certification Body