

Certificate

mdc medical device certification GmbH
certifies that

botiss biomaterials GmbH
Hauptstraße 28
15806 Zossen b. Berlin
Germany

with the locations listed in the attachment

for the scope

**development, production, storage, shipment 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	2015-04-15
Valid until	2018-09-05
Registration no.	D1323300015
Report no.	P15-00552-44849
Stuttgart	2015-04-15



Head of Certification Body



Attachment of the certificate

No. D1323300015

date 2015-04-15

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Location	Scope
botiss biomaterials GmbH, Emser Straße 36, 10719 Berlin	storage and shipment of products for bone and tissue regeneration
botiss biomaterials GmbH, Uhlandstraße 20 -20/II, 10623 Berlin	development, production, shipment and distribution of products for bone and tissue regeneration
botiss biomaterials GmbH, Hauptstraße 28, 15806 Zossen b. Berlin	development, production, storage, shipment and distribution of products for bone and tissue regeneration



Head of Certification Body