

Certificate

mdc medical device certification GmbH
certifies that

botiss biomaterials GmbH
Hauptstraße 28
15806 Zossen
Germany

with the location

Ullsteinstraße 108, Aufgang D
12109 Berlin

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 13485:2016 + AC:2016 - ISO 13485:2016

Valid from	2021-12-16
Valid until	2024-12-15
Registration no.	D1323300055
Report no.	P21-01034-208630
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Head of Certification Body



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