EC Certificate

mdc medical device certification GmbH
Notified Body 0483
herewith grants

botiss biomaterials GmbH
Hauptstraße 28
15806 Zossen
Germany

for the scope

collagen matrix for implantation (dental) mucoderm®
(see attachment)

the

EC Design Examination Certificate

The examination of the design of the product by mdc has proven
that the design meets the requirements according to

Annex II – Section 4
of the Council Directive 93/42/EEC

of 14 June 1993 concerning medical devices.

This certificate is only valid in connection with a valid
mdc certificate according to Annex II – excluding section 4 for the
above mentioned products.

Valid from	2019-11-21
Valid until	2024-05-26
Registration no.	D1323300032
Report no.	P19-00782-148638
Stuttgart	2019-10-29

Head of Certification Body

medical device certification mdc

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-246.10.06

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Kriegerstraße 6
D-70191 Stuttgart, Germany
Phone: +49-(0)711-253597-0
Fax: +49-(0)711-253597-10
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Attachment of the certificate

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Product category	Product	Class
collagen matrix for implantation (dental)	mucoderm® Größe 15x20 mm REF 701520	III
	mucoderm® Size 20x30 mm REF 702030	III
	mucoderm® Size 30x40 mm REF 703040	III
	mucoderm® Größe 10x30 mm REF 701030	III
	mucoderm® Größe 20x50 mm REF 702050	III
	mucoderm® Größe 20x100 mm REF 702000	III
	mucoderm® Größe 30x5 mm REF 703005	III
	mucoderm® Größe Ø8 mm REF 710208	III
	mucoderm® Size Ø10 mm REF 710210	III

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