EC Certificate

mdc medical device certification GmbH
Notified Body 0483
herewith grants

botiss biomaterials GmbH
Hauptstraße 28
15806 Zossen b. Berlin
Germany

for the scope

mucoderm - composed of porcine collagen
used solely or in combination with appropriate augmentation materials
(e.g. autogenic bone, allogeneic, xenogeneic or alloplastic bone replacement materials)
indicated in dental surgery for immediate or
delayed guided tissue and bone regeneration,
Sizes 15 x 20 mm, 10 x 30 mm, 20 x 30 mm, 20 x 50 mm,
20 x 100 mm, 30 x 5 mm, 30 x 40 mm, 8 mm, Ø 10 mm

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	2015-04-15
Valid until	2019-11-20
Registration no.	D1323300021
Report no.	P15-00552-44849
Stuttgart	2015-04-15

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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