

Certificate

mdc medical device certification GmbH
certifies that

curasan

curasan AG
Lindigstraße 4
63801 Kleinostheim
Germany

with the facility

Werk Frankfurt
In der Schildwacht 13
65933 Frankfurt am Main

for the scope

**Development, manufacturing and distribution of
medical devices in the fields of dental implants,
bone substitution and regeneration, titanium foils.**

**Distribution of medical devices in the fields of bone and tissue regeneration (membranes)
wound healing and arthrosis (hyaluronic acid products)**

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

(ISO 13485:2003)

This certificate is valid until:	2011-08-02
Certificate registration no.:	0017.48.01/3
Stuttgart	2007-08-02



Head of
Certification Body



EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith certifies that

**curasan AG
Lindigstraße 4
63801 Kleinostheim
Germany**

for the scope

CERASORB®

**β-tricalcium phosphate ceramic
for filling of bone defects**

has introduced and applies a

Quality System

for the design, manufacture and final inspection.

The mdc audit has proven, that this quality system
meets all requirements according to

**Annex II – Section 3
of the Council Directive 93/42/EEC**

of 14 June 1993 concerning medical devices.

The surveillance will be held as specified in Annex II, Section 5.

This certificate is valid until:	2012-07-11
Certificate registration no:	0017.01.14/0
Stuttgart	2007-07-11



Head of
Certification Body



Akkreditiert durch
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln
und Medizinprodukten
ZLG-ZQ-976.94.05

EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith grants

**curasan AG
Lindigstraße 4
63801 Kleinostheim
Germany**

for the scope

CERASORB® PARO

**β-tricalcium phosphate ceramic
for filling of bone defects**

the

EC Design Examination Certificate

The examination of the design of the product by mdc has proven,
that according to the report of the
file review no. E0017.11/2007-07-11 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 – Section 3 for the
above mentioned products.

This certificate is valid until:	2012-07-12
Certificate registration no:	0017.11.14/0
Stuttgart	2007-07-12



Head of
Certification Body



Akkreditiert durch
Zentralstelle der Länder
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bei Arzneimitteln
und Medizinprodukten
ZLG-ZE-975.94.04

mdc medical device certification GmbH
Kriegerstraße 6
D-70191 Stuttgart, Germany
Phone: +49-(0)711-253597-0
Fax: +49-(0)711-253597-10
Internet: <http://www.mdc-ce.de>

EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith grants

curasan AG
Lindigstraße 4
63801 Kleinostheim
Germany

for the scope

Cerasorb® M

β-tricalcium phosphate ceramic
for filling of bone defects

the

EC Design Examination Certificate

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that according to the report of the
file review no. E0017.11/2007-07-12 the design meets the requirements according to

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Certificate registration no:	0017.11.13/0
Stuttgart	2007-07-12



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EC Certificate

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Notified Body 0483
herewith grants

curasan AG
Lindigstraße 4
63801 Kleinostheim
Germany

for the scope
Cerasorb® M Block Forms
β-tricalcium phosphate ceramic
for filling of bone defects

the

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for the scope

Cerasorb® Granules

β-tricalcium phosphate ceramic
for filling of bone defects

the

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for the scope

Cerasorb® Block Forms

B-tricalcium phosphate ceramic
for filling of bone defects

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