



EC-CERTIFICATE

(Full quality assurance system)



This is to certify that the company

METOXIT AG

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8240 Thayngen
Switzerland

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Device	Class
Sterile ceramic dental implants	Ib
Dental Instruments	Im and IIa
Ceramic dental abutments	Ib
Ceramic blanks and corresponding dyeing liquids for the manufacturing of dental prosthetic restorations	IIa

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no.	066501 MR2
Certificate unique ID	170585351
Effective date	2017-09-22
Expiry date	2022-09-21
Frankfurt am Main	2017-09-19

DQS Medizinprodukte GmbH

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DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.

