ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Production Quality Assurance

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 12 05 69185 004

Manufacturer:

Panasonic Healthcare Co., Ltd.
247 Fukutake-ko
Saijo, Ehime
793-8510 JAPAN

EC-Representative:

Panasonic Marketing Europe GmbH
Panasonic Testing Centre
Winsbergring 15,
22525 Hamburg
GERMANY

Product Category(ies):

Dental Porcelains

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: **OAQ235005335A**

Valid from: **2012-07-23**
Valid until: **2017-07-22**

Date, 2012-07-26

Hans-Heiner Junker

522169

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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No. G2 12 05 69185 004

Facility(ies):

Panasonic Healthcare Co., Ltd. Medical Instruments & Hospital System Business Unit
247 Fukutake-ko, Saijo, Ehime, 793-8510 JAPAN