



EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

No. CE 01220
Issued To: **Kerr Italia S.r.l.**
Via Passanti, 332
Scafati (SA)
84018
Italy

In respect of:

The manufacture of dental adhesives, pit and fissure sealants, cavity lining materials, non-medicated pulp capping materials, dental cements, materials for temporary crowns and bridges, temporary filling materials, composite dental materials for direct insertion, fixed and removable prostheses, and endodontic sealers.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Gary Fenton, Global Assurance Director

First Issued: **29/02/1996**

Date: **26/02/2014**

Expiry Date: **28/02/2019**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.