bsi.

By Royal Charter

EC Certificate - Production Quality Assurance

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

No. **CE 01488**

Issued To: **GC Europe N.V.**
Research Park
Interleuvenlaan 33
Leuven
B-3001
Belgium

In respect of:

The manufacture of dental glass ionomer restorative systems, luting cements, fibre reinforcements, temporary filling, crown and bridge materials, conditioners, coating agents, composite resin and adhesive systems for direct and indirect restorations, temporary crown and bridge material, relining and periodontal dressing materials, ceramics, alloys and adhesives for dental applications.

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 2797):

Gary E Slack

Gary E Slack, Senior Vice President Medical Devices

First Issued: **1996-12-17** Date: **2019-04-25** Expiry Date: **2024-05-06**

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