



EC Certificate - Production Quality Assurance

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

No. CE 51688
Issued To: **GAP Research Co. Limited**
Unit 2 Ebbsfleet Estate
Stonebridge Road
Northfleet
Kent
DA11 9DZ
United Kingdom

In respect of:

The manufacture of temporary tooth filling material and prophylactic cups and brushes for dental professional use.

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: **22 September 1999**Date: **24 September 2014**Expiry Date: **23 September 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.