



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 07 73936 001

Manufacturer: Copan Italia S.p.a.
Via F. Perotti, 10
25125 Brescia
ITALY

Facility(ies): Copan Italia S.p.a.
Via F. Perotti, 10, 25125 Brescia, ITALY

Product Category(ies): Medical Swabs and applicators
for sample collection and preservation
and microbiology investigations
with or without transport medium

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: ITA 204097

Valid until: 2015-09-01

Hans-Heiner Junker



Date, 2010-09-02

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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