



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 17 12 33250 011

Manufacturer: **CLINI-LAB s.r.l.**
VIA 2° STRADA 14 Z.I.
35026 Conselve (PD)
ITALY



Facility(ies): CLINI-LAB s.r.l.
VIA 2° STRADA 14 Z.I., 35026 Conselve (PD), ITALY

Product Category(ies): **Sterile disposable gloves
for medical examination**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: ITA1009165

Valid from: 2018-02-13
Valid until: 2023-02-12



Date, 2018-01-11

Stefan Preiß

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

Page 1 of 1