EC Cerificate

EC Directive 93/42/EEC Annex V

Production Quality Assurance

Medical Devices

Registration No.: **DD 60094266 0001**

Report No.: **28106257 001**

Manufacturer: **Colonplus Equipments & Speculums, S.L.**
c/Portuetxe, 13 – 1.C
20018 San Sebastian (Guipúzcoa)
Spain

Scope: **Colon cleaning equipments**

COLONPLUS AUTOMATICA
COLONPLUS MANUAL

Date of Expiry: **29/05/2019**

The Notified Body hereby declares that the requirements of annex V of the Directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned Directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to annex III is required.

Notified Body

Pogliano Milanese (MI) **27/05/2014**

B. Tuzza

TÜV Rheinland Italia S.r.l. - Via Mattei, 3 - 20010 - Pogliano Milanese (MI)
Accrditated by Ministero della Salute and by Ministero dello Sviluppo Economico
with decree of January 09th , 2013 (G.U. n. 32 February 07th 2013)

Notified under No. **1936** to the EC Commission

CE The CE marking may be used if all relevant and effective EC Directives are complied with CE