



EC Certificate

Full Quality Assurance System according to Medical Devices Directive 93/42/EEC Annex-II Section 3

Certificate Number: 1984-MDD-19-562

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive.

Organization:

SCHÖNN MEDIZINTECHNIK GmbH

Head Office: Helena Rubinstein Strasse 4-G, 40699 Erkrath, Düsseldorf, Germany

Factory: Herforder Strasse 46, 32602 Vlotho, Germany

Products: Medical Vacuum Supply System, AGSS Supply System, Medical Compressed Gas Supply Systems (for O₂, N₂, N₂O, CO₂ and Air), Medical Air Supply System, Medical Gas Terminal Units and Probes, Oxygen Flowmeters, Vacuum Regulators, Zone Wall Units, Medical Alarm Panels, Operation Room Surgical Control Panels, Intensive Care Units, Pendants, Bed Head Units, Medical Copper Tubes and Fittings

The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa for details.

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Kiwa Belgelendirme Hizmetleri A.Ş. is Notified Body under Council Directive 93/42/EEC concerning medical devices with identification number:1984

Muhteşem Gökhan Yücel
Head of Notified Body

18 May 2021, Istanbul, Turkey

CERTIFICATE