

EC Certificate**Production Quality Assurance System according to
Medical Devices Directive 93/42/EEC Annex-V****Certificate Number: 1984-MDD-17-459**

We hereby declare that an examination has been carried out following the requirements of the national legislation to which the undersigned is subject, transposing Annex-V of the Directive 93/42/EEC on medical devices. We certify that the production quality system conforms with the relevant provisions of the aforementioned legislation.

Organization:**BIOTEX MEDİKAL TEKSTİL İTHALAT İHRACAT
SANAYİ VE TİCARET ANONİM ŞİRKETİ**

İslampaşa mahallesi 2 Nolu Şehitler Caddesi No: 41/G Merkez, Rize, Turkey

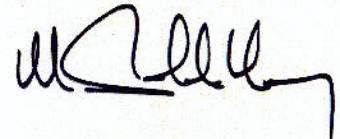
Products: Sterile Disposable Surgical Drapes, Gowns and Sets

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

Report Number: M.5022.02
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Revision Number: 01
Expiry Date: 27 May 2024

Kiwa Belgelendirme Hizmetleri A.Ş. has audited the quality system restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions in accordance with MDD Annex V and found that the quality system meets the applicable requirements in MDD Annex V.

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



Muhtesem Gökhan Yücel
Head of Notified Body

05 March 2020, Istanbul, Turkey