

MEDBAR TIBBİ MALZEMELER TURİZM SANAYİ VE TİCARET ANONİM ŞİRKETİ

FATİH MAH. 1142 SOK. NO: 35 SARNIÇ GAZİEMİR – İZMİR – TURKEY

with a scope of

SURGICAL DRAPE PRODUCTS, IV FLOW REGULATOR PRODUCTS,
KARMAN CANNULA PRODUCTS, ENDOSCOPY MOUTHPIECES, URINE
COLLECTION PRODUCTS, MUCOUS ASPIRATION PRODUCTS,
ARTROCOPY SETS, VOMIT/EMESIS BAG PRODUCTS, SCRUB HAND
BRUSHES, FILTERED MOUTHPIECE PRODUCTS, CERVICAL BRUSH
PRODUCTS, AMNIOTIC POUCH PERFORATORS, FECAL PARASITE
CONCENTRATION PRODUCTS, INTENSIVE CARE PRODUCTS
PROCESSES: PRODUCTION, PACKAGING, STERILIZATION, STORING
DISTRIBUTION AND ETHYLENE OXIDE STERILIZATION SERVICES ARE
UNDER THE SCOPE OF EN ISO 11135 STANDARD

Medical devices - Quality management systems - Requirements for
regulatory purposes

"Following elements of the standard are excluded"

"7.5.3" "7.5.4" "7.5.9.2"

EN ISO 13485:2016

Certificate No : M 10326
Initial Certification Date : 03 October 2019
Certification Date : 03 October 2019
Expiration Date : 02 October 2022

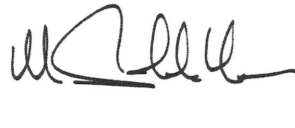


Medical Device Q.M.S.
TS EN ISO/IEC 17021-1

AB-0006-YS



TÜRKAK BDS NO
YS-694E-E7CB



General Manager

Kiwa Certification Services Inc.

ITOSB 9. Cadde No. 15 Tepeören Tuzla - Istanbul - Turkey

Tel: + 90 216 593 25 75 Faks : + 90 216 593 25 74

Web: www.kiwa.com.tr E-mail: info@kiwa.com.tr

Certificate is valid till expiration date, subject to successful completion of periodical surveillance audits.

Please contact above numbers for detailed information.