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CERTIFICATE

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 MOUTHPİECES, 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

General Manager

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Certificate is valid till expiration date, subject to successful completion of periodical surveillance audits.
Please contact above numbers for detailed information.

Last Modified: 03 October 2019 - R 00

TÜRKAK
Medical Device Q.M.S.
TS EN ISO/IEC 17021-1
AB-0006-YS

TÜRKAK BDS NO
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