



Certificate of Registration

This is to certify that

Quality Management System

for Medical Devices
of

FATİH MEDİKAL TIBBİ CİHAZ SİSTEMLERİ-FATİH KAÇMAZ

ESENLER MAHALLESİ İTİR SOKAK NO:2/6
PENDİK / İSTANBUL / TÜRKİYE.

complies with the requirements of

EN ISO 13485:2012

This certificate is valid concerning all activities related to:

**MANUFACTURING, EXPORTING AND SELLING OF MOBILE
HEALTH DEVICES, X-RAY DEVICES, AUDIOMETRY BOOTHS,
AUDIOMETERS, CR SYSTEMS AND BLOOD COLLECTION SEATS.**

**MOBİL SAĞLIK ARAÇLARI, RÖNTGEN CİHAZLARI,
ODYOMETRİ KABİNLERİ, ODYOMETRELER, CR SİSTEMLERİ
VE KAN ALMA KOLTUKLARI İMALATI, İHRACATI VE SATIŞI.**

MD-1016
Certificate No.

Mar. 26, 2018
Date of this Certificate

Mar. 25, 2019
*Next Audit Due Date

Mar. 26, 2018
Date of Initial Registration

Mar. 25, 2021
Certification Expiry Date


Managing Director/Director



TRANSPACIFIC CERTIFICATIONS LIMITED

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Accreditation by Joint Accreditation System of Australia and New Zealand (Accreditation No.M2640303IN)
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<http://www.jas-anz.org/our-directory/certified-organisations>

This certificate is only valid if it is available/valid on TCL website at <http://tclcertifications.com/client-register/>.

The certificate of Registration remains the property of Transpacific Certifications Limited and shall be returned immediately upon request.
*In case if Surveillance Audit is not allowed to be conducted on or before the specified date; the Certificate shall be Suspended/Withdrawn.