



TURKISH ACCREDITATION AGENCY

ACCREDITATION CERTIFICATE

As a "Type A" Inspection Body

VALTECH VALİDASYON LABORATUVAR TEST HİZMETLERİ TİCARET LİMİTED ŞİRKETİ

Central Address: YENİ BATI MAH. MURATDAĞI SK. No:10/3 YENİMAHALLE/ANKARA Ankara / Türkiye

is accredited in accordance with TS EN ISO/IEC 17020:2012 standard within the scope given in Annex following the assessment conducted by TURKAK.

Accreditation Number : AB-0694-M

Accreditation Date : 16.03.2022

Revision Date / Number : 09.04.2026 / 04


This certificate shall remain in force until 15.03.2030, subject to continuing compliance with the standard TS EN ISO/IEC 17020:2012, related regulations and requirements.

Gülden Banu Müderrisoğlu
Secretary General



Turkish Accreditation Agency (TURKAK) is a signatory to the European co-operation for Accreditation (EA) Multilateral Agreement (MLA) and International Laboratory Accreditation Cooperation (ILAC) Mutual Recognition Agreement (MRA) in the scope of ISO/IEC 17020.

This document has been signed by Gülden Banu Müderrisoğlu with a secure electronic signature in accordance with the electronic signature law numbered 5070. Use the QR code to verify the e-signed document.


 <p>Inspection TS EN ISO/IEC 17020 AB-0694-M</p>	VALTECH VALİDASYON LABORATUVAR TEST HİZMETLERİ TİCARET LİMİTED ŞİRKETİ	
	Accreditation Nr : AB-0694-M Revision Nr: 04 Date: 09.04.2026	
"Type A" Inspection Body		
Address : YENİ BATI MAH. MURATDAĞI SK. No:10/3 YENİMAHALLE/ANKARA Ankara / Türkiye	Phone : +90 312 502 2509 Fax : - Email : info@valtech.com.tr Website : www.valtech.com.tr	

Environment/Clean Rooms, Related Controlled Environments and Air Conditioning Systems

Inspection Field	Inspection Type	Standard/Specification
HVAC Systems of Health Care Rooms or Facilities / Class Ib Operation Rooms	Initial Inspection	ISO 14644-1 ISO 14644-2 ISO 14644-3 DIN 1946-4 ASHRAE 170
HVAC Systems of Health Care Rooms or Facilities / Class Ib Operation Rooms	Periodic Inspection	ISO 14644-1 ISO 14644-2 ISO 14644-3 DIN 1946-4 ASHRAE 170
HVAC Systems of Health Care Rooms or Facilities / Class II Intensive Care Units and Other Hospital Hygienic Areas	Initial Inspection	ISO 14644-1 ISO 14644-2 ISO 14644-3 DIN 1946-4 ASHRAE 170
HVAC Systems of Health Care Rooms or Facilities / Class II Intensive Care Units and Other Hospital Hygienic Areas	Periodic Inspection	ISO 14644-1 ISO 14644-2 ISO 14644-3 DIN 1946-4 ASHRAE 170
HVAC Systems of Manufacturing Areas, Laboratories and Laminar Air Flow Areas	Initial Inspection	The Rules Governing Medicinal Products in the European Union Volume 4 EU Guidelines for Good Manufacturing Practice for Medicinal Products for Human and Veterinary Use Annex 1- Manufacture of Sterile Medicinal Products FDA Guidance for Industry Sterile Drug Products Produced by Aseptic Processing- Current Good Manufacturing Practice TS EN ISO 14644-1 TS EN ISO 14644-2 TS EN ISO 14644-3 IEST-RP-CC006 IEST-RP-CC034
HVAC Systems of Manufacturing Areas, Laboratories and Laminar Air Flow Areas	Periodic Inspection	The Rules Governing Medicinal Products in the European Union Volume 4 EU Guidelines for Good Manufacturing Practice for Medicinal Products for Human and Veterinary Use Annex 1- Manufacture of Sterile Medicinal Products FDA Guidance for Industry Sterile Drug Products Produced by Aseptic Processing- Current Good Manufacturing Practice TS EN ISO 14644-1 TS EN ISO 14644-2 TS EN ISO 14644-3 IEST-RP-CC006 IEST-RP-CC034
Biosafety Cabinets	Periodic Inspection	TS EN 12469 ANSI/NSF-49
Unidirectional Airflow (UDAF) Cabinets	Initial Inspection	TS EN ISO 14644-1 TS EN ISO 14644-2 TS EN ISO 14644-3 IEST-RP-CC034 IEST-RP-CC002
Unidirectional Airflow (UDAF) Cabinets	Periodic Inspection	TS EN ISO 14644-1 TS EN ISO 14644-2 TS EN ISO 14644-3 IEST-RP-CC002 IEST-RP-CC034



Accreditation Scope

 TÜRKAK Inspection TS EN ISO/IEC 17020 AB-0694-M	VALTECH VALİDASYON LABORATUVAR TEST HİZMETLERİ TİCARET LİMİTED ŞİRKETİ	
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Biocontamination Control	Initial Inspection	TS EN 17141
Biocontamination Control	Periodic Inspection	TS EN 17141
Temperature and Humidity Performance Verification- Storage Areas, Cold Rooms	Initial Inspection	EU Guideline on Good EU Guideline on Good Distribution Practice of Medicinal Products For Human Use WHO Technical Report Series No.961, Annex 9 Supplement 7,8 FDA 21 CFR Part 11
Temperature and Humidity Performance Verification- Storage Areas, Cold Rooms	Periodic Inspection	EU Guideline on Good EU Guideline on Good Distribution Practice of Medicinal Products For Human Use WHO Technical Report Series No.961, Annex 9 Supplement 7,8 FDA 21 CFR Part 11
Temperature and Humidity Performance Verification- Stability Rooms and Air Conditioning Cabinets	Initial Inspection	EudraLex Volume 4 Annex 15 Good Manufacturing Practice(GMP) for Human and Veterinary Use EU Guidelines on Good Distribution Practice (GDP) of Products For Human Use Medicinal WHO Technical Report Series No.961, Annex 9 Supplement 7,8 FDA 21 CFR Part 11
Temperature and Humidity Performance Verification- Stability Rooms and Air Conditioning Cabinets	Periodic Inspection	EudraLex Volume 4 Annex 15 Good Manufacturing Practice(GMP) for Human and Veterinary Use EU Guidelines on Good Distribution Practice (GDP) of Products For Human Use Medicinal WHO Technical Report Series No.961, Annex 9 Supplement 7,8 FDA 21 CFR Part 11

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Environment and Health/Sterilizers

Inspection Field	Inspection Type	Standard/Specification
Steam Sterilizers	Initial Inspection	TS EN ISO 17665 TS EN 285 TS EN 13060
Steam Sterilizers	Periodic Inspection	TS EN ISO 17665 TS EN 285 TS EN 13060

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