ISO 13485:2016

Certificate of Registration

This is to Certify that

Medical Device Quality Management System of

GROTEX LIMITED LIABILITY COMPANY (GROTEX LLC)

71/2, INDUSTRIALNY AVE., LIT A, 195279, SAINT - PETERSBURG, RUSSIAN FEDERATION (RUSSIA)

100, LIT. B, INDUSTRIAL AREA CENTRAL DEPARTMENT, 193149, SVERDLOVSK URBAN LOCATION, VSEVOLOZHSKY MUNICIPAL DISTRICT, LENINGRAD REGION, RUSSIAN FEDERATION (RUSSIA)

has been assessed and found to conform to the requirements of

ISO 13485:2016

for the following scope :

DEVELOPMENT, MANUFACTURE AND DISTRIBUTION OF LIQUID STERILE AND NON-STERILE (SOLUTIONS, AEROSOLS, GELS, SPRAYS, DROPS, INJECTIONS) MEDICAL PRODUCTS.

Certificate No	: **22IMJW97**		
Initial Registration Date	: 10/06/2022	Issuance Date	: 10/06/2022
Date of Expiry	: 09/06/2025		
1st Surve. Due	: 10/05/2023	2nd Surve. Due	: 10/05/2024

Director

ACCREDITED Management Systems Certification Body MSCB-119

AQC MIDDLE EAST LLC

Head Office: Office No. 02, Ground Floor, Sharjah Media City, Sharjah, UAE. e-mail: info@aqcworld.com,

**Validity of the Certificate is subject to successful completion of surveillance audit on or before of due date. (in case surveillance audit is not allowed to be conducted, this certificate shall be suspended/withdrawal).*

Certificate Verification: Please Re-check the validity of certificate at http://www.aqcworld.com/activeclients.aspx or www.aqcworld.com at Active Clients.

Certificate is the property of AQC Middle East LLC and shall be returned immediately when demanded