

Electronic Product Certificate

Annual Report

Certificate Expires: September 1, 2022

This certifies that:

Agop Havluciyan X-Ray Guvenlik Sistemleri Merkezi Sanayi Ticaret Ltd. Beylikduzu OSB Mahallesi, 10 Cadde, NO. 14 Beylikduzu, Istanbul TR-34524 Turkey

has submitted its Radiation Safety Report to the U.S. Food and Drug Administration pursuant to part 1002 of Title 21, US Code of Federal Regulations, such submission having been verified as currently effective on the date hereof by Registrar Corp:

Product Trade Name: XRC with model(s) XRC 50-30, XRC 60-40,

XRC 60-40DV, XRC 60-40P, XRC 100-100,

XRC 100-100DV, XRC 100-100LC, XRC 150-165,

XRC 150-165DV, XRC 75-55, XRC 75-55DV, XRC 180-180, XRC 180-180DV, XRC 65-85,

XRC 90-75, XRC 125-125DV, XRC 85-85, XRC 50-30SE,

XRC 65-45, XRC 180-180DV 320KV

Accession Number: 2130195-000 Date Submitted: July 2, 2021 U.S. Agent / Registrar Corp

Registrant Contact: 144 Research Drive, Hampton Virginia, 23666 USA

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Registrar Corp will confirm that such submission has been completed upon request and presentation of this certificate until September 1, 2021, unless terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Registrar Corp, waives, and will hold harmless and indemnify Registrar Corp from any and all claims in connection with this product, its labeling, radiation safety reporting, commerce or use. Submission of a Radiation Safety Report does not constitute approval of the firm,

document or device by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of report submission or the assignment of an accession number is misleading. The U.S. Food and Drug Administration of the assignment of an accession number is misleading. The U.S. Food and Drug Administration of the assignment of an accession number is misleading. The U.S. Food and Drug Administration.

Oavid Lemarz

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Registrar-Corp

Executive Director