



2021

CERTIFICATE OF REGISTRATION

This certifies that:

VIVOSENS BIYOTEKNOLOJI AR-GE, SAN. & TIC. LTD. STI
Etiler Mah. Yanarsu Sok
Dogan Apt. No:44/4
Besiktas Istanbul, TR 34467

is registered with the U.S. Food and Drug Administration for FY 2021 pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations:

Establishment Registration:	3015512211
Device Classification Name:	DIAZONIUM COLORIMETRY, UROBILINOGEN (URINARY, NON-QUANT.)
Product Code:	CDM
Regulation Number:	862.1785
Official Correspondent and U.S. Agent:	Registrar Corp 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

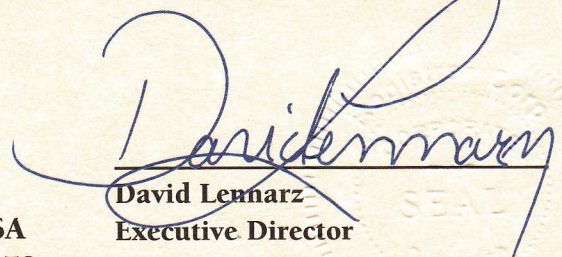
Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does 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. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Registrar Corp assumes no liability to any person or entity in connection with the foregoing.

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Registrar Corp

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David Lennarz
Executive Director
Registrar Corp
Dated: October 27, 2020