4/10/2020

FDA | U.S. Food and Drug Administration

Registration Confirmation

Facility: METRIS INSTRUMENTS EAST, LLC, South Setauket, New York, UNITED STATES

You have successfully entered your facility registration and device listing information. You should print a copy of this page for your records. Listing numbers appear below for the products manufactured, developed, or processed at this facility.

As a manufacturer, specification developer, or single-use device reprocessor, you are required to pay an annual fee for medical device facility registration.

You will receive another e-mail providing you with your registration number in approximately 30 to 90 days. Until your registration number is assigned, reference your Owner/Operator number in any correspondence with the Center for Devices and Radiological Health.

Your registration will be valid through Dec 31, 2020. An e-mail will be sent to the Owner/Operator and the Official Correspondent 90 days before the facility is required to re-register for Fiscal Year 2020 with instructions on how and when to re-register.

Note: Registering your device facility and listing your devices does not, in any way, constitute FDA approval of your facility or your devices.

Should you have any questions, please send an e-mail to reglist@cdrh.fda.gov (mailto:reglist@cdrh.fda.gov).

The Owner/Operator Number for this Registration is: 10068952

Facility Information

Initial Importer:	**Y**
Facility Name:	**METRIS INSTRUMENTS EAST, LLC**
Address:	**25 Longmeadow Place, 21 South Setauket, New York, 11720, UNITED STATES**
Foreign Trade Zone:	**N**

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