

Bob Ogden

bobogden2005@gmail.com>

Notification of New Device Establishment Registration

1 message

reglist@cdrh.fda.gov <reglist@cdrh.fda.gov> Reply-To: reglist@cdrh.fda.gov To: bobogden2005@gmail.com Fri, Apr 10, 2020 at 4:19 PM

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
Center for Devices and Radiological Health
10903 New Hampshire Ave., WO66 Room 2621
Silver Spring, Maryland 20993-0002

April 10, 2020

Name of Official Correspondent Address of Official Correspondent DAVID LENNARZ 144 RESEARCH DRIVE

HAMPTON, VIRGINIA 23666

UNITED STATES

DAVID.LENNARZ@REGISTRARCORP.COM

Owner Operator Number

10068952

Dear Sir or Madam,

We have received your registration and listing information for the following medical device establishment

Establishment Name

Establishment Address

METRIS INSTRUMENTS EAST, LLC

25

LONGMEADOW PLACE, 21

SOUTH SETAUKET, NEW YORK 11720 UNITED

STATES

The information submitted has been processed and entered into the FDA Registration and Device Listing Database. Your device establishment is now considered registered. You will be notified of your official registration number within 90 days.

Once you receive a registration number, you are required to re-register on an annual basis from October through December. Failure to re-register every year will invalidate your registration and result in your device establishment and listing information being removed from the FDA Medical Device Registration and Listing Web site.

For inquiries about the status of your registration or assignment of your registration number, please contact the Registration and Listing Program Office at reglist@cdrh.fda.gov or calling (301) 796-7400.

If you have any questions regarding FDA policy related to the Registration and Listing program, please contact the Registration and Listing staff at device.reg@fda.hhs.gov or calling (301) 796-7400.