Dear MARCO ANCESCHI:

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2022:

Registration Number: 3011306968
Owner Operator Number: 10047761
ANCHIPLAST SRL
Via Briana, 4
Bondeno di Gonzaga, Mantova 46023
ITALY

If you do not see a registration number assigned to the establishment and your establishment previously had one, please send an email to reglist@cdrh.fda.gov and include the registration number you believe is assigned to your establishment. We will review and determine if a duplicate registration has been created for your establishment.

Your registration is valid until December 31, 2022. Registration for 2023 will be conducted between October 1 and December 31, 2022.

Please note that 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 the CDRH Registration and Listing Helpdesk at reglist@cdrh.fda.gov.

CDRH Registration and Listing Helpdesk
Imports & Registration and Listing Team
Division 2 Establishment Support
Office of Regulatory Programs
Office of Product Evaluation and Quality
Center for Devices and Radiological Health
U.S. Food and Drug Administration