From: reglist@cdrh.fda.gov <reglist@cdrh.fda.gov>
Sent: Thursday, December 30, 2021 7:39 AM
To:
Subject: Notification of New Device Establishment Registration

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Center for Devices and Radiological Health

10903 New Hampshire Ave., WO66 Room 1423

Silver Spring, Maryland 20993-0002

December 30, 2021

Name of Official Correspondent	DAVID LENNARZ
Address of Official Correspondent	144 RESEARCH DRIVE HAMPTON, VIRGINIA 23666 UNITED STATES DAVID.LENNARZ@REGISTRARCORP.COM
Owner Operator Number	10083918

Dear Sir or Madam,

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

Establishment Name	PACIFIC BIOSUPPLY LLC
Establishment Address	530-B HARKLE ROAD
	SUITE 100
	SANTA FE, NEW MEXICO 87505
	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.