From: CDRH Registration and Listing <reglist@CDRH.FDA.GOV>
Subject: **Registration Number 2029386: Successful 2021 Medical Device Establishment Registration**
Date: December 19, 2020 10:22:09 AM PST
To: "LMI@EARTHLINK.NET" <LMI@EARTHLINK.NET>
▶ 2 Attachments, 51.5 KB

Dear DANIEL LUCAS, JR.:

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2021:

Registration Number: 2029386
Owner Operator Number: 9012199
LUCAS MEDICAL, INC.
1751 SOUTH DOUGLASS RD.
ANAHEIM, CA 92806
UNITED STATES

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, 2021. Registration for 2022 will be conducted between October 1 and December 31, 2021.

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

Tel: 301-796-7400, Option 1
Email: reglist@cdrh.fda.gov