From: CDRH Registration and Listing [mailto:reglist@CDRH.FDA.GOV]
Sent: Friday, December 30, 2016 8:17 PM
To: Tanya DiSalvo <tdisalvo@criteriontool.com>
Subject: Registration Number 1528668: Successful 2017 Medical Device Establishment Registration

Dear TANYA DISALVO:

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2017:

Registration Number: 1528668
Owner Operator Number: 9005596
CRITERION TOOL & DIE INC
5349 W 161st St
BROOK PARK, OH 44142
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, 2017. Registration for 2018 will be conducted between October 1 and December 31, 2017.

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
Office of Compliance
Center for Devices and Radiological Health
U.S. Food and Drug Administration
Tel: 301-796-7400, Option 1
Email: reglist@cdrh.fda.gov