FDA

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Establishment Registration & Device Listing

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Establishment:
TISSUETECH, INC.
Business Trade Name:
BIO-TISSUE, INC.

8305 NW 27th St Ste 101
Doral, FL 33122
Registration Number: 3003415347
FEI Number*: 3003415347
Status: Active
Date Of Registration Status: 2022

Owner/Operator:
TISSUETECH, INC.[6]
7300 Corporate Center Drive
Suite 700
Miami, FL US 33126
Owner/Operator Number: 9067198[7]

Official Correspondent:
Lorraine Chua
7300 Corporate Center Drive
Suite 700
Miami, FL 33126
Phone: 786-4567719

* Firm Establishment Identifier (FEI) should be used for identification of entities within the imports message set

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Page Last Updated: 11/29/2021

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Silver Spring, MD 20993
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Delhy Arias

Subject: FW: Registration Number 3003415347: Successful 2022 Medical Device Establishment Registration
Attachments: Header.jpg; SignatureBlockLogo.png

From: CDRH Registration and Listing <reglist@CDRH.FDA.GOV>
Sent: Tuesday, October 12, 2021 5:01:50 AM
To: Lorraine Chua <lchua@tissuetechinc.com>
Subject: Registration Number 3003415347: Successful 2022 Medical Device Establishment Registration

Dear Lorraine Chua:

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

Registration Number: 3003415347
Owner Operator Number: 9067198
TISSUETECH, INC.
8305 NW 27th St Ste 101
Doral, FL 33122
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, 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

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