Timothy Alcid

From: CDRH Registration and Listing <reglist@CDRH.FDA.GOV>
Sent: Wednesday, November 17, 2021 9:59 PM
To: RA_Licenses
Subject: Registration Number 2249062: Successful 2022 Medical Device Establishment Registration
Attachments: Header.jpg; SignatureBlockLogo.png

Dear JOEL OSBORNE:

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

Registration Number: 2249062
Owner Operator Number: 9032400
MUSCULOSKELETAL TRANSPLANT FOUNDATION
125 MAY ST, STE 300
EDISON CORP CTR
EDISON, NJ 08837
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

Help (./help/index.html)

DRLM Home (mainMenu.htm)

✔ Facility

✔ Products Listing

Annual Registration Successful

Facility: MUSCULOSKELETAL TRANSPLANT FOUNDATION, EDISON, New Jersey, UNITED STATES

You have successfully updated your registration and listing information for 2022.

Your registration will be valid through Dec 31, 2022.

Be sure to print this page for your records.

The next registration renewal period is October 1 - December 31, 2022.

Registering your facility and listing devices does not, in any way, constitute FDA approval of your facility or devices.

You may contact the FDA with any questions at reglist@cdrh.fda.gov.

The Owner/Operator Number for this Registration is: 9032400.

Facility Information

Registration Number:
2249062

Initial Importer:
N

Facility Name:
MUSCULOSKELETAL TRANSPLANT FOUNDATION

Legal Name:

Address:
125 MAY ST, STE 300, EDISON CORP CTR
EDISON, New Jersey, 08837, UNITED STATES

DUNS Number:

Foreign Trade Zone:
N

Facility URL:

Other Business Trade Name(s):

Establishment located on a campus:

Owner/Operator Information

Owner/Operator Number:
9032400

Contact Name:
JOEL - OSBORNE

Company:
MUSCULOSKELETAL TRANSPLANT FOUNDATION

Address: **125 MAY ST., SUITE 300 , --**
Edison, NEW JERSEY, 08837, UNITED STATES

Telephone:
732 - 6610202

Fax:
732 - 6612189

E-mail: **RA_Licenses@MTF.org**

DUNS Number:

Official Correspondent Information

Contact Name:
JOEL - OSBORNE

Company:
MUSCULOSKELETAL TRANSPLANT FOUNDATION

Address: **125 MAY ST., SUITE 300 , --**
Edison, NEW JERSEY, 08837, UNITED STATES

Telephone:
732 - 6610202

Fax:
732 - 6612189

E-mail: **RA_Licenses@MTF.org**

DUNS Number:

Device Listings

Listing Number	Premarket Submission Number/Type	Product Code(s)	Device Name(s)	Activities
D209115	K042125	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
D209124	K091218	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D208635	K040262	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
		GXP	METHYL METHACRYLATE FOR CRANIOPLASTY	

D209127	K040501	NUN	BONE GRAFTING MATERIAL, HUMAN SOURCE	Manufacturer
D422442	510(k) exempt	QBO	Instruments designed for press-fit osteochondral implants	Repackager/Relabeler
D351843	K181633	GAT	SUTURE, NONABSORBABLE, SYNTHETIC, POLYETHYLENE	Manufacturer
D227432	K100754	FMF	Syringe, piston	Repackager/Relabeler
D327564	K110771	FMF	Syringe, piston	Repackager/Relabeler
D371590	510(k) exempt	KYZ	SYRINGE, IRRIGATING (NON DENTAL)	Manufacturer

Date of Initial Registration: 1997-12-08 15:45:51.0