Timothy Alcid

From: CDRH Registration and Listing <reglist@CDRH.FDA.GOV>
Sent: Wednesday, November 17, 2021 10:46 PM
To: RA_Licenses
Subject: Registration Number 1000307092: Successful 2022 Medical Device Establishment Registration
Attachments: Header.jpg; SignatureBlockLogo.png

CAUTION: This email originated from outside of the organization. Do not click links or open attachments unless you recognize the sender and know the content is safe.

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: 1000307092
Owner Operator Number: 9032400
MUSCULOSKELETAL TRANSPLANT FOUNDATION
1232 MIDVALLEY DRIVE
JESSUP, PA 18434
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

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Help (./help/index.html)

DRLM Home (mainMenu.htm)

✔ Facility

✔ Products Listing

Annual Registration Successful

Facility: MUSCULOSKELETAL TRANSPLANT FOUNDATION, JESSUP, Pennsylvania, 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:
1000307092

Initial Importer:
N

Facility Name:
MUSCULOSKELETAL TRANSPLANT FOUNDATION

Legal Name:

Address:
1232 MIDVALLEY DRIVE,
JESSUP, Pennsylvania, 18434, 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
D371590	510(k) exempt	KYZ	SYRINGE, IRRIGATING (NON DENTAL)	Manufacturer
D327564	K110771	FMF	Syringe, piston	Repackager/Relabeler
D422442	510(k) exempt	QBO	Instruments designed for press-fit osteochondral implants	Repackager/Relabeler
D227432	K100754	FMF	Syringe, piston	Repackager/Relabeler
D209124	K091218	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer

		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D351843	K181633	GAT	SUTURE, NONABSORBABLE, SYNTHETIC, POLYETHYLENE	Manufacturer
D209115	K042125	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
D208635	K040262	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
		GXP	METHYL METHACRYLATE FOR CRANIOPLASTY	
D209127	K040501	NUN	BONE GRAFTING MATERIAL, HUMAN SOURCE	Manufacturer

Date of Initial Registration: 2007-08-01 09:41:50.0