

## Sandeep Chiplonkar

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**From:** CDRH Registration and Listing <reglist@CDRH.FDA.GOV>  
**Sent:** Friday, October 23, 2020 6:44 PM  
**To:** RA\_Licenses  
**Subject:** Registration Number 2249062: Successful 2021 Medical Device Establishment Registration  
**Attachments:** Header.jpg; SignatureBlockLogo.png  
**Follow Up Flag:** Follow up  
**Flag Status:** Flagged

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Dear Keith Alderman:

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

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](mailto: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](mailto: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](mailto: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, EDISON, New Jersey, UNITED STATES

You have successfully updated your registration and listing information for 2021.

Your registration will be valid through Dec 31, 2021.

Be sure to print this page for your records.

The next registration renewal period is October 1 - December 31, 2021.

**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](mailto: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**

**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):**

### 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:**  
**Keith Alderman**

**Company:**  
**Musculoskeletal Transplant Foundation**

**Address:** **125 May St Ste 300**  
**Edison, NEW JERSEY, 08837, UNITED STATES**

**Telephone:**  
**732 - 6610202 - 2592**

**Fax:**  
**-**

**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	<input type="text" value="Manufacturer"/>
D209124	K091218	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	<input type="text" value="Manufacturer"/>
		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D208635	K040262	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	<input type="text" value="Manufacturer"/>
		GXP	METHYL METHACRYLATE FOR CRANIOPLASTY	

D209127	K040501	NUN	BONE GRAFTING MATERIAL, HUMAN SOURCE	Manufacturer
D351843	K181633	GAT	SUTURE, NONABSORBABLE, SYNTHETIC, POLYETHYLENE	Manufacturer
D227432	K100754	FMF	Syringe, piston	Repackager/Relabeler
D327564	K110771	FMF	Syringe, piston	Repackager/Relabeler
D371590	Exempt	KYZ	SYRINGE, IRRIGATING (NON DENTAL)	Manufacturer

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