## Sandeep Chiplonkar

**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



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 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.

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 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.

**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 | 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 |
| 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**