

## Sandeep Chiplonkar

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**From:** CDRH Registration and Listing <reglist@CDRH.FDA.GOV>  
**Sent:** Tuesday, October 17, 2017 9:39 AM  
**To:** RA\_Licenses  
**Subject:** Registration Number 2249062: Successful 2018 Medical Device Establishment Registration

**Follow Up Flag:** Follow up  
**Flag Status:** Flagged



Dear JOEL OSBORNE:

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2018:

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, 2018. Registration for 2019 will be conducted between October 1 and December 31, 2018.

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  
Office of Compliance  
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)



Dear Reporting Official:

This letter describes how to update your annual establishment registration for human cells, tissues, and cellular and tissue-based products (HCT/Ps). The annual update period begins November 15, 2017 and ends December 31, 2017 (see 21 CFR 1271.21). For this year, a few expectations have changed for the annual registration update using the electronic Human Cell and Tissue Establishment Registration System (eHCTERS). Note that FDA published in the Federal Register of August 31, 2016 (81 FR 60170) a final rule that amended FDA's regulations governing drug establishment registration and drug listing, including establishment registration and listing for HCT/Ps. Access the final rule [here](#).

- 1) You are only required to list HCT/Ps that are regulated solely under Section 361 of the Public Health Service Act and 21 CFR Part 1271 of the regulations (361 HCT/Ps). Annual registration using an appropriate FDA electronic registration system is also required for HCT/Ps regulated as drugs, devices, and/or biological products under 21 CFR Parts 207 or 807, as applicable; however, eHCTERS should no longer be used for these products, and you may deactivate the existing registrations in eHCTERS if you are not manufacturing any 361 HCT/Ps. Please refer to [Drug Establishment Registration and Listing](#) or [Device Registration and Listing](#) websites for additional information.

Note: FDA does not require establishments that manufacture drugs and devices under an investigational new drug application (IND) (21 CFR Part 312) or an investigational device exemption (IDE) (21 CFR Part 812) to register and list their HCT/Ps until the HCT/P is approved through a biologics license application (BLA), a new drug application (NDA), or a premarket approval application (PMA); or cleared through a premarket notification submission (510(k)). Therefore, establishments that only manufacture HCT/Ps currently under an IND or IDE may deactivate or change their registration in eHCTERS as indicated.<sup>1</sup>

- 2) Beginning November 29, 2017, HCT/P establishment and product listing information required under §1271.22 (including the annual update) must be submitted electronically, unless waived in certain circumstances. Establishments may request a waiver from the electronic submission requirement as described in §1271.23.

The following instructions describe how to submit your annual registration update electronically. For your convenience, your current registration information is attached to this email that includes your FDA Establishment Identifier (FEI) # and the validated date of your current registration.

- 1) Access CBER On-Line through a secure web page [here](#).
- 2) If you have forgotten your password, click the link "Forgot your User Name or Password?" and follow the instructions. Once you receive your user name and/or password information, return to the CBER On-Line Login Screen.
- 3) Enter your User Name and Password, select the application: eHCTERS Tissue Establishment Registration from the drop-down list, then press the "LOGIN" button.
- 4) On the Activity Selection screen, select both "Edit Validated Form", and select "Annual Registration/Listing" as "Reason for Submission". Select your establishment from the drop-down list,

then press the “Continue” button. If you do not see your establishment in the drop-down list, follow the instructions for adding your establishment to your account.

- 5) Verify that all registration information is accurate on each screen and update the information where applicable. When your review and update is complete press the “Submit to FDA” button.
- 6) When prompted, sign the form by entering your Reporting Official E-Mail address then press “Continue”.

Please do not submit your registration update too early as the change will not be reflected in the annual update if submitted prior to November 15, 2017.

After receipt of your annual registration update, we will provide the Reporting Official with an updated copy of the registration form, which will reflect a new validation date in the upper right-hand corner of the form. Please keep this copy at the establishment location for inspection purposes.

If you have questions or need assistance in updating the registration, please contact us at 240-402-8369. You may also submit questions about registration to [tissuereg@fda.hhs.gov](mailto:tissuereg@fda.hhs.gov).

Sincerely yours,

Aletha Davis-Knight  
Human Tissue Establishment Registration Coordinator  
Division of Human Tissues  
Office of Tissues and Advanced Therapies

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<sup>1</sup> [Guidance for Industry: Regulation of Human Cells, Tissues, and Cellular and Tissue-Based Products \(HCT/Ps\) - Small Entity Compliance Guide](#)

[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 2018.

Your registration will be valid through Dec 31, 2018.

Be sure to print this page for your records.

The next registration renewal period is October 1 - December 31, 2018.

**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:****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
D209127	K040501	NUN	BONE GRAFTING MATERIAL, HUMAN SOURCE	Manufacturer
D095116	Exempt	HXL	MALLET	Specification Developer Remanufacturer
D209124	K091218	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D209119	K110003	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer

		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D227434	K980987	FMF	Syringe, piston	Repackager/Relabeler
D227432	K100754	FMF	Syringe, piston	Repackager/Relabeler
D209115	K042125	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
D227435	K921520	JKA	Tubes, vials, systems, serum separators, blood collection	Repackager/Relabeler
D227436	K931367	JKA	Tubes, vials, systems, serum separators, blood collection	Repackager/Relabeler
E379492	K043260	MTJ	WAX,BONE	Manufacturer
E381064	K042829	MQV	FILLER, BONE VOID, CALCIUM COMPOUND	Manufacturer
		MBP	Filler, bone void, osteoinduction (w/o human growth factor)	
D016309	K053339	GAT	SUTURE, NONABSORBABLE, SYNTHETIC, POLYETHYLENE	Manufacturer
		JDW	PIN, FIXATION, THREADED	
		HWC	Screw, fixation, bone	
D151442	Exempt	KDC	INSTRUMENT, SURGICAL, DISPOSABLE	Specification Developer Repackager/Relabeler
D208635	K040262	GXP	METHYL METHACRYLATE FOR CRANIOPLASTY	Manufacturer
		MQV	FILLER, BONE VOID, CALCIUM COMPOUND	
D016312	K061498	JDW	PIN, FIXATION, THREADED	Manufacturer
		MAI	Fastener, fixation, biodegradable, soft tissue	
A978688	Exempt	LXH	ORTHOPEDIC MANUAL SURGICAL INSTRUMENT	Manufacturer
D095120	Exempt	HXG	TAMP	Remanufacturer
D142437	BK020051	KSS	SUPPLIES, BLOOD-BANK	Manufacturer
D142438	BK060006	KSS	SUPPLIES, BLOOD-BANK	Manufacturer
D097840	Exempt	HTO	REAMER	Remanufacturer Specification Developer

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

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125 MAY STREET | EDISON, NJ 08837 | [MTFBIOLOGICS.ORG](http://MTFBIOLOGICS.ORG)

November 27, 2017

**RE: Removal of HCT/Ps Regulated as Medical Devices from FDA Annual Registration**

To Whom It May Concern:

2017 Annual Registration for 2018; FDA has provided a letter (attached), which states that only HCT/Ps solely regulated under Section 361 of the Public Health Service Act and 21 CFR Part 1271 of the regulations (361 HCT/Ps), should be listed on FDA HCT/P Registrations; and any HCT/Ps regulated as Medical Devices should be removed. The following MTF products which are HCT/Ps that are regulated as Medical Devices, all of which have associated 510ks (attached); have been/will be removed.

- DBX Mix
- DBX Paste
- DBX Putty
- DBX
- DBX Strip
- DBX Inject
- AFT

Please note that these products now fall under MTFs FDA Medical Device Registration, and no changes have been made with regards to status; these products are FDA approved.

If you have any questions, please do not hesitate to contact me at (732)-661-0202, extension 2209 or email MTF Regulatory Affairs at [Regulatory\\_Customer\\_Inquiry@mtf.org](mailto:Regulatory_Customer_Inquiry@mtf.org).

Sincerely,

A handwritten signature in black ink, appearing to read "Joel C. Osborne".

Joel C. Osborne  
Vice President, Regulatory Affairs