DEPARTMENT OF HEALTH AND HUMAN SERVICES
PUBLIC HEALTH SERVICE
FOOD AND DRUG ADMINISTRATION

ESTABLISHMENT REGISTRATION AND LISTING FOR HUMAN CELLS, TISSUES, AND CELLULAR AND TISSUE-BASED PRODUCTS DESCRIBED IN 21 CFR 1271.10

FEI: 3007499718

Other FDA Registrations:
Blood:
Devices: FEI: 3010097171
Drugs:

Reason For Last Submission: Annual Registration/Listing
Last Annual Registration Year: 2026
Last Registration Receipt Date: 12/02/2025
Summary Report Print Date: 12/15/2025

Legal Name and Location:

Millstone Medical Outsourcing
8836 Polk Lane
Suite 100

Olive Branch, Mississippi 38654
USA

Phone: 508-679-8384 **Ext.:**

Reporting Official:

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Satellite Recovery Establishment: No
Parent Manufacturing Establishment FEI No.:
Testing For Micro-Organisms Only: No

Note: FDA acceptance of an establishment registration and HCT/P listing does not constitute a determination that an establishment is in compliance with applicable rules and regulations or that the HCT/P is licensed or approved by FDA (21 CFR 1271.27(b)).

HCT/P(s)	Donor Type(s)	Establishment Functions								Date of Discontinuance	Date of Resumption	Proprietary Name(s)
		Recover	Screen	Donor Testing	Package	Process	Store	Label	Distribute			
Amniotic Membrane							X	X	X			***See full text on next page.
Blood Vessel												
Bone							X	X	X			***See full text on next page.
Cardiac Tissue - non-valved												
Cartilage												
Cornea												
Dura Mater												
Embryo												
Fascia												
Heart Valve												
HPC Apheresis												
HPC Cord Blood												
Ligament							X	X	X			
Nerve Tissue												
Oocyte												
Ovarian Tissue												
Pancreatic Islet Cells - autologous												
Parathyroid												
Pericardium							X	X	X			Puros Pericardium
Peripheral Blood Mononuclear Cells												
Peritoneal Membrane												
Sclera												
Semen												
Skin							X	X	X			***See full text on next page.
Tendon							X	X	X			
Testicular Tissue												
Tooth Pulp												
Umbilical Cord Tissue							X	X	X			***See full text on next page.

FDA information collection OMB Control number: 0910-0543, expiration date: 08/31/2026

Additional Information: No additional information provided.

Proprietary Name(s):		
	Amniotic Membrane	Amniofix; CTM Thin; CTM Thick; CTM Flow; CTM Boost; CTM Paste; CTM Powder; Amnioflo; AMNIOGRAFT; AMNIOGUARD; CLARIX 100; CLARIX CORD; CLARIX FLO; NEOX 100; NEOX CORD; NEOX FLO; NEOX RT; Puros; CopiOs; Biotivity A/C Plus; SimpliMax Amniotic Membrane Allograft
	Bone	Osteoamp; Purebone; Exponent; Allograft Bone Wedge; SeaSpine Capistrano Cervical Spacer; SeaSpine PLIF Allograft; SeaSpine Compressible Bone Matrix; SeaSpine Cervical Allograft; Laminoplasty Cortical Allograft; DBMForm; Allogenix DBM; Bonus CC Matrix; Bonus II DMB; EquivaBone; FiberStack DBM; Fortis ALIF, Composite, Cortical, PLIF, TLIF; Indux Cortical or Cancellous; InterGro DBM; PrimaGen Advanced Allograft; Osteostim Composite; Cortical, ALIF, PLIF; StaGraft Cancellous; StaGraft DBM; StaGraft Fiber; Trinnect; Puros; Puros-S; Fortitude; Bonus Triad; Copios; OSTEOAMP SELECT Flowable, RegenerOss; Puros; RegenaVate; RegenerOssAllograft Putty; Treace Cancellous Wedge; CortiFuse; OssiGraft¿, OssiGraft Prime¿
	Skin	DermaSpan; DermaSpan Mesh; SimpliDerm;
	Umbilical Cord Tissue	CTM Flow; CTM Boost; CTM Paste; CTM Flow LV; CTM X-Thick

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