

FEI: 3005287828

**Other FDA Registrations:**

**Blood:**

**Devices:**

**Drugs:**

Reason For Last Submission: Annual Registration/Listing  
 Last Annual Registration Year: 2023  
 Last Registration Receipt Date: 11/18/2022  
 Summary Report Print Date: 12/01/2022

**Legal Name and Location:**

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**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												
Blood Vessel												
Bone												
Cardiac Tissue - non-valved												
Cartilage												
Cornea												
Dura Mater												
Embryo	Anonymous, Directed, SIP						X	X	X			
Fascia												
Heart Valve												
HPC Apheresis												
HPC Cord Blood												
Ligament												
Nerve Tissue												
Oocyte	Anonymous, Directed, SIP						X	X	X			
Ovarian Tissue							X	X	X			
Pancreatic Islet Cells - autologous												
Parathyroid												
Pericardium												
Peripheral Blood Mononuclear Cells												
Peritoneal Membrane												
Sclera												
Semen	Anonymous, Directed, SIP	X	X		X	X	X	X	X			
Skin												
Tendon												
Testicular Tissue					X	X	X	X	X			
Tooth Pulp												
Umbilical Cord Tissue												

**Additional Information:** No additional information provided.

**Proprietary Name(s):**

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