

FEI: 3005033855

**Other FDA Registrations:**

**Blood:**

**Devices:**

**Drugs:**

Reason For Last Submission: Annual Registration/Listing  
 Last Annual Registration Year: 2020  
 Last Registration Receipt Date: 12/10/2019  
 Summary Report Print Date: 01/07/2020

**Legal Name and Location:**

Fairfax Cryobank, Inc.  
 1305 West 34th Street  
 Suite 210

Austin, Texas 78705  
 USA

Phone: 512-206-0408

**Ext.:**

**Reporting Official:**

Megan Taylor, Document Administrator  
 3015 Williams Drive  
 Suite 110  
 Fairfax, Virginia 22031  
 USA  
 Phone: 800-338-8407 Ext.  
 mtaylor@givf.com

**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	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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Legal Name:

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