

FEI: 3003651454

Other FDA Registrations:

Blood:

Devices:

Drugs:

Reason For Last Submission: Annual Registration/Listing
 Last Annual Registration Year: 2021
 Last Registration Receipt Date: 12/02/2020
 Summary Report Print Date: 12/14/2020

Legal Name and Location:

ReproTech, Ltd.
 33 Fifth Avenue NW
 STE 900

St Paul, Minnesota 55112
 USA

Phone: 651-489-0827

Ext.:

Reporting Official:

Amy S Erickson Hagen, QA Officer
 18 South Ninth Street
 STE 201
 Columbia, Missouri 65201
 USA
 Phone: 612-747-4154 Ext.
 aehagen@reprotech.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)).

Establishment Functions

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			
Skin												
Tendon												
Testicular Tissue							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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