

PART I - ESTABLISHMENT INFORMATION

3. OTHER FDA REGISTRATIONS

a. BLOOD FDA 2830 NO. _____

b. DEVICES FDA 2891 NO. _____

c. DRUG FDA 2856 NO. _____

4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code)
Cord For Life
 270 Northlake Boulevard, Suite 1012
 Altamonte Springs, Florida 32701

a. PHONE 407-834-8333 EXT _____

b. SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____)

c. TESTING FOR MICRO-ORGANISMS ONLY

5. ENTER CORRECTIONS TO ITEM 4

6. MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if applicable, number and street, city, state, country, and post office code)
 Cord For Life
 Attn: Deborah A. Sardone, MEd, MLT (ASCP)
 270 Northlake Boulevard Suite 1012
 Altamonte Springs, Florida 32701

a. PHONE 407-834-8333 EXT _____

7. ENTER CORRECTIONS TO ITEM 6

b. PHONE _____

8. U.S. AGENT

9. REPORTING OFFICIAL'S SIGNATURE

a. TYPED NAME Deborah A. Sardone, MEd, MLT (ASCP)

b. E-MAIL dsardone@lifeforcecyobanks.com

c. TITLE QA Manager

d. DATE 26-FEB-2018

PART II - PRODUCT INFORMATION

10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / PS

| Types of HCT / Ps | Establishment Functions | | | | | | 11 HCT/PS DESCRIBED IN 21 CFR 1271.10 | 11 HCT/PS REGULATED AS MEDICAL DEVICES | 12 HCT/PS REGULATED AS BIOLOGICAL DRUGS | 13 HCT/PS REGULATED AS BIOLOGICAL DRUGS | 14. PROPRIETARY NAME(S) | |
|----------------------------------|-------------------------|--------|------|---------|---------|-------|---------------------------------------|--|---|---|-------------------------|-------|
| | Recover | Screen | Test | Package | Process | Store | | | | | | Label |
| a. Bone | | | | | | | | | | | | |
| b. Cartilage | | | | | | | | | | | | |
| c. Cornea | | | | | | | | | | | | |
| d. Dura Mater | | | | | | | | | | | | |
| e. Embryo/ | | | | | | | | | | | | |
| f. Fascia | | | | | | | | | | | | |
| g. Heart Valve | | | | | | | | | | | | |
| h. Ligament | | | | | | | | | | | | |
| i. Oocyte | | | | | | | | | | | | |
| j. Pericardium | | | | | | | | | | | | |
| k. Peripheral Blood Stem | | | | | | | | | | | | |
| l. Sclera | | | | | | | | | | | | |
| m. Semen | | | | | | | | | | | | |
| n. Skin | | | | | | | | | | | | |
| o. Somatic Cell Therapy Products | | | | | | | | | | | | |
| p. Tendon | | | | | | | | | | | | |
| q. Umbilical Cord Blood | | | | | | | | | | | | |
| r. Vascular Graft | | | | | | | | | | | | |
| s. Placenta | | | | | | | | | | | | |
| t. Umbilical Cord | | | | | | | | | | | | |
| u. | | | | | | | | | | | | |
| v. | | | | | | | | | | | | |

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 (HCT/PS)**
(See reverse side for instructions)

1. REGISTRATION NUMBER
(FDA Establishment Identifier)
FEI: 1000525281

2. REASON FOR SUBMISSION
a. INITIAL REGISTRATION / LISTING
b. ANNUAL REGISTRATION / LISTING
c. CHANGE IN INFORMATION
d. INACTIVE

See Instructions for OMB Statement. FORM APPROVED OMB No 0910-0543. Expiration Date: 3/31/2017
VALIDATION -FOR FDA USE ONLY
VALIDATED BY FDA-19-NOV-2016
DISTRICT: Florida
PRINTED BY FDA-15-DEC-2016

PART I - ESTABLISHMENT INFORMATION

3. OTHER FDA REGISTRATIONS
a. BLOOD FDA 2830 NO
b. DEVICES FDA 2891 NO
c. DRUG FDA 2656 NO

4. PHYSICAL LOCATION (include legal name, number and street, city, state, country, and post office code)
LifeForce Cryobank Sciences, Inc.
270 Northlake Boulevard, Suite 1012
Altamonte Springs, Florida 32701

a. PHONE 407-834-8333 EXT
b. SATELLITE RECOVERY ESTABLISHMENT
(MANUFACTURING ESTABLISHMENT FEI NO _____)
c. TESTING FOR MICRO-ORGANISMS ONLY
5. ENTER CORRECTIONS TO ITEM 4

6. MAILING ADDRESS OF REPORTING OFFICIAL (include institution name if applicable, number and street, city, state, country, and post office code)
LifeForce Cryobank Sciences, Inc.
Attn: Denise F. Clifton, M.L.T., AT (ASCP)
270 Northlake Boulevard Suite 1012
Altamonte Springs, Florida 32701

a. PHONE 407-834-8333 EXT
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE

8. U.S. AGENT

a. E-MAIL
9. REPORTING OFFICIAL'S SIGNATURE

a. TYPED NAME Denise F. Clifton, M.L.T., AT (ASCP)
b. E-MAIL dclifton@lifeformcryobanks.com
c. TITLE QA Manager
d. DATE 18-NOV-2016

PART II - PRODUCT INFORMATION

10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / PS

| Types of HCT / Ps | Establishment Functions | | | | | Label | Distribute | 11 HCT/PS DESCRIBED IN 21 CFR 1271.10 | 12 HCT/PS REGULATED AS MEDICAL DEVICES | 13 HCT/PS REGULATED AS BIOLOGICAL DRUGS OR DRUGS | 14 PROPRIETARY NAME(S) |
|----------------------------------|-------------------------|--------|------|---------|---------|-------|------------|---------------------------------------|--|--|------------------------|
| | Recover | Screen | Test | Package | Process | | | | | | |
| a. Bone | | | | | | | | | | | |
| b. Cartilage | | | | | | | | | | | |
| c. Cornea | | | | | | | | | | | |
| d. Dura Mater | | | | | | | | | | | |
| e. Embryo | | | | | | | | | | | |
| f. Fascia | | | | | | | | | | | |
| g. Heart Valve | | | | | | | | | | | |
| h. Ligament | | | | | | | | | | | |
| i. Oocyte | | | | | | | | | | | |
| j. Pericardium | | | | | | | | | | | |
| k. Peripheral Blood Stem | | | | | | | | | | | |
| l. Sclera | | | | | | | | | | | |
| m. Semen | | | | | | | | | | | |
| n. Skin | | | | | | | | | | | |
| o. Somatic Cell Therapy Products | | | | | | | | | | | |
| p. Tendon | | | | | | | | | | | |
| q. Umbilical Cord Blood | | | | | | | | | | | |
| r. Vascular Graft | | | | | | | | | | | |
| s. Placenta | | | | | | | | | | | |
| t. Umbilical Cord | | | | | | | | | | | |
| u. | | | | | | | | | | | |
| v. | | | | | | | | | | | |