

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: 0001077605

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

14. PROPRIETARY NAME(S)

11. HCT/PS DESCRIBED IN 21 CFR 1271.10

12. HCT/PS REGULATED AS MEDICAL DEVICES

13. HCT/PS REGULATED AS DRUGS OR BIOLOGICAL DRUGS

PART II - PRODUCT INFORMATION	Establishment Functions				11. HCT/PS DESCRIBED IN 21 CFR 1271.10	12. HCT/PS REGULATED AS MEDICAL DEVICES	13. HCT/PS REGULATED AS DRUGS OR BIOLOGICAL DRUGS	14. PROPRIETARY NAME(S)
	Recover	Screen	Test	Package				
10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps								
Types of HCT / Ps								
a. Bone								
b. Cartilage								
c. Cornea								
d. Dura Mater								
e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous								
f. Fascia								
g. Heart Valve								
h. Ligament								
i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous								
j. Pericardium								
k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input checked="" type="checkbox"/> Allogeneic			X				X	
l. Sclera								
m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous								
n. Skin								
o. Somatic Cell Therapy Products <input checked="" type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic				X			X	
p. Tendon								
q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic								
r. Vascular Graft								
s. Therapeutic Cells			X				X	
t.								
u.								
v.								

3. OTHER FDA REGISTRATIONS
a. BLOOD FDA 2830 NO. FEI: 0001077605
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)
Medic Regional Blood Center
1601 Ailor Avenue
Knoxville, Tennessee 37921

a. PHONE 865-524-3074 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)
MEDIC Regional Blood Center
Attn: Martha S. Cox, MPH, MT
1601 Ailor Avenue
Knoxville, Tennessee 37921

a. PHONE 865-524-3074 EXT 668
b. PHONE

7. ENTER CORRECTIONS TO ITEM 6

8. U.S. AGENT

a. E-MAIL

9. REPORTING OFFICIAL'S SIGNATURE
Matthew Cox
2/3/2015

a. TYPED NAME Martha S. Cox, MPH, MT

b. E-MAIL mcox@medicblood.org

c. TITLE Chief Quality Officer

d. DATE 02-JAN-2015