

DEPARTMENT OF HEALTH AND HUMAN SERVICES FOOD AND DRUG ADMINISTRATION ESTABLISHMENT REGISTRATION AND LISTING FOR HUMAN CELLS, TISSUES, AND CELLULAR AND TISSUE-BASED PRODUCTS (HCT/PS) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier): FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY  VALIDATED BY FDA: 12-DEC-2005 PRINTED BY FDA: 12-DEC-2005 DISTRICT OFFICE: San Francisco
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PART I - ESTABLISHMENT INFORMATION

3. OTHER FDA REGISTRATIONS

a. BLOOD FDA 2830 NO. _____

b. DEVICES FDA 2891 NO. _____

c. DRUG FDA 2656 NO. _____

PART II - PRODUCT INFORMATION

10. ESTABLISHMENT FUNCTIONS:

a. RECOVER c. TEST e. PROCESS g. LABEL

b. SCREEN d. PACKAGE f. STORE h. DISTRIBUTE

4. PHYSICAL LOCATION *(include legal name, number and street, city, state, country, and post office code)*

Community Blood Center dba Community Tissue Services
 3425 N. First Street., Suite 103
 Fresno, California 93726

PHONE 559-224-1168 EXT

TYPES OF HCT/PS	11. HCT/PS DESCRIBED IN 21 CFR 1271.10	12. HCT/PS REGULATED AS MEDICAL DEVICES, DRUGS, OR BIOLOGICAL DRUGS	13. PROPRIETARY NAME(S)
a. Bone	X		
b. Cartilage	X		
c. Cornea			
d. Dura Mater			
e. Embryo			
f. Fascia	X		
g. Heart Valve	X		
h. Ligament	X		
i. Oocyte			
j. Pericardium	X		
k. Peripheral Blood Stem Cells			
l. Sclera			
m. Semen			
n. Skin	X		
o. Somatic Cells			
p. Tendon	X		
q. Umbilical Cord Blood Stem Cells			
r. Vascular Graft	X		
s.			
t.			
u.			
v.			

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)*

Community Blood Center/Community Tissue Services
 349 S. Main Street
 Dayton, Ohio 45402-2715

PHONE 937-461-3450 EXT 3288

7. ENTER CORRECTIONS TO ITEM 6

8. U.S. AGENT

a. E-MAIL b. PHONE

9. REPORTING OFFICIAL'S SIGNATURE

a. TYPED NAME Judith E. Woll, MD
 b. E-MAIL jwoll@cbocls.org
 c. TITLE CEO d. DATE 29-NOV-2005

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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY * 3001238554 * VALIDATED By FDA:12/10/07 PRINTED By FDA:12/17/07 DISTRICT: San Francisco
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	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous											
	f. Fascia	X	X			X	X	X	X			
	g. Heart Valve	X	X							X		
	h. Ligament	X	X				X	X	X	X		
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous											
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	t.											
	u.											
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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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:30-DEC-2008 DISTRICT: San Francisco PRINTED BY FDA:05-JAN-2009
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a. E-MAIL _____	g. Heart Valve <input checked="" type="checkbox"/> SIP h. Ligament <input checked="" type="checkbox"/> Directed i. Oocyte <input type="checkbox"/> Anonymous j. Pericardium k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous l. Sclera m. Semen <input type="checkbox"/> Family Related n. Skin o. Somatic Cell Therapy Products <input type="checkbox"/> Allogeneic p. Tendon q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Allogeneic r. Vascular Graft	X	X				X	X	X	X	X		
a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO/Medical Director d. DATE 08-DEC-2008	g. Heart Valve <input checked="" type="checkbox"/> SIP h. Ligament <input checked="" type="checkbox"/> Directed i. Oocyte <input type="checkbox"/> Anonymous j. Pericardium k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous l. Sclera m. Semen <input type="checkbox"/> Family Related n. Skin o. Somatic Cell Therapy Products <input type="checkbox"/> Allogeneic p. Tendon q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Allogeneic r. Vascular Graft	X	X				X	X	X	X	X		
	g. Heart Valve <input checked="" type="checkbox"/> SIP h. Ligament <input checked="" type="checkbox"/> Directed i. Oocyte <input type="checkbox"/> Anonymous j. Pericardium k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous l. Sclera m. Semen <input type="checkbox"/> Family Related n. Skin o. Somatic Cell Therapy Products <input type="checkbox"/> Allogeneic p. Tendon q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Allogeneic r. Vascular Graft	X	X				X	X	X	X	X		
	g. Heart Valve <input checked="" type="checkbox"/> SIP h. Ligament <input checked="" type="checkbox"/> Directed i. Oocyte <input type="checkbox"/> Anonymous j. Pericardium k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous l. Sclera m. Semen <input type="checkbox"/> Family Related n. Skin o. Somatic Cell Therapy Products <input type="checkbox"/> Allogeneic p. Tendon q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Allogeneic r. Vascular Graft	X	X				X	X	X	X	X		

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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input checked="" type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY 1 VALIDATED BY FDA:20-NOV-2009 DISTRICT: San Francisco PRINTED BY FDA:02-DEC-2009
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION									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)
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / PS												
	Types of HCT / PS	Establishment Functions											
		Recover	Screen	Test	Package	Process	Store	Label	Distribute				
4. PHYSICAL LOCATION <i>(Include legal name, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services 3425 N. First Street., Suite 103 Fresno, California 93726 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	a. Bone	X	X				X	X	X	X			
	b. Cartilage	X	X				X	X	X	X			
	c. Cornea	X	X								X		
5. ENTER CORRECTIONS TO ITEM 4	d. Dura Mater												
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
	f. Fascia	X	X				X	X	X	X			
6. MAILING ADDRESS OF REPORTING OFFICIAL <i>(Include institution name if applicable, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610	g. Heart Valve	X	X							X			
	h. Ligament	X	X				X	X	X	X			
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	j. Pericardium	X	X				X	X	X	X			
	k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
	l. Sclera	X	X								X		
8. U.S. AGENT a. E-MAIL _____	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
	n. Skin	X	X				X	X	X	X			
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbcts.org c. TITLE CEO d. DATE 19-NOV-2009	p. Tendon	X	X				X	X	X	X			
	q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
	r. Vascular Graft	X	X								X		
	s. Parathyroid						X		X	X			
	t. Peritoneal Membrane	X	X				X	X	X	X			
	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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:05-JAN-2010 DISTRICT: San Francisco PRINTED BY FDA:23-FEB-2010
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION									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)	
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps													
a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	<i>Establishment Functions</i>													
<i>Types of HCT / Ps</i>	Recover	Screen	Test	Package	Process	Store	Label	Distribute	Distribute					
4. PHYSICAL LOCATION <i>(Include legal name, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services 3425 N. First Street., Suite 103 Fresno, California 93726 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	a. Bone	X	X				X	X	X	X				
	b. Cartilage	X	X				X	X	X	X				
	c. Cornea	X	X							X				
	d. Dura Mater													
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
	f. Fascia	X	X				X	X	X	X				
5. ENTER CORRECTIONS TO ITEM 4	g. Heart Valve	X	X							X				
	h. Ligament	X	X				X	X	X	X				
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
6. MAILING ADDRESS OF REPORTING OFFICIAL <i>(Include institution name if applicable, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610	j. Pericardium	X	X				X	X	X	X				
	k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
	l. Sclera	X	X							X				
	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	n. Skin	X	X				X	X	X	X				
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
8. U.S. AGENT a. E-MAIL _____	p. Tendon	X	X				X	X	X	X				
	q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
	r. Vascular Graft	X	X							X				
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO d. DATE 29-DEC-2009	s. Parathyroid						X		X	X				
	t. Peritoneal Membrane	X	X				X	X	X	X				
	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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input checked="" type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY 1 VALIDATED BY FDA:01-JUN-2010 DISTRICT: San Francisco PRINTED BY FDA:01-JUN-2010
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION								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)
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / PS											
	Types of HCT / Ps	Establishment Functions										
		Recover	Screen	Test	Package	Process	Store	Label	Distribute			
3. OTHER FDA REGISTRATIONS a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	a. Bone					X	X	X	X			
	b. Cartilage					X	X	X	X			
	c. Cornea											
4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services 7100 N. Financial Drive, Suite 105 Fresno, California 93720 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	d. Dura Mater											
	e. Embryo											
	f. Fascia						X	X	X	X		
	g. Heart Valve											
	h. Ligament						X	X	X	X		
	i. Oocyte											
5. ENTER CORRECTIONS TO ITEM 4	j. Pericardium					X	X	X	X			
	k. Peripheral Blood Stem Cells											
	l. Sclera											
	m. Semen											
	n. Skin						X	X	X	X		
	o. Somatic Cell Therapy Products											
6. MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if applicable, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610	p. Tendon					X	X	X	X			
	q. Umbilical Cord Blood Stem Cells											
	r. Vascular Graft											
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	s. Parathyroid					X		X	X			
	t. Peritoneal Membrane					X	X	X	X			
	u.											
8. U.S. AGENT a. E-MAIL _____	v.											
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO d. DATE 13-MAY-2010												

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) <i>(See reverse side for instructions)</i>	1. REGISTRATION NUMBER (Field Establishment Identifier) FEI: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY 1 VALIDATED BY FDA:29-DEC-2010 DISTRICT: San Francisco PRINTED BY FDA:05-JAN-2011
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION														
3. OTHER FDA REGISTRATIONS a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps									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)		
	Types of HCT / Ps	Recover	Screen	Test	Package	Process	Store	Label	Distribute						
4. PHYSICAL LOCATION <i>(Include legal name, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services 7100 N. Financial Drive, Suite 105 Fresno, California 93720 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	a. Bone						X	X	X	X					
	b. Cartilage						X	X	X	X					
	c. Cornea														
	d. Dura Mater														
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous														
	f. Fascia						X	X	X	X					
5. ENTER CORRECTIONS TO ITEM 4	g. Heart Valve														
	h. Ligament						X	X	X	X					
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous														
	j. Pericardium						X	X	X	X					
	k. Peripheral Blood Stem Cells <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic														
	l. Sclera														
6. MAILING ADDRESS OF REPORTING OFFICIAL <i>(Include institution name if applicable, number and street, city, state, country, and post office code)</i> Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous														
	n. Skin						X	X	X	X					
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic														
	p. Tendon						X	X	X	X					
8. U.S. AGENT a. E-MAIL _____	q. Umbilical Cord Blood Stem Cells <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic														
	r. Vascular Graft														
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO d. DATE 21-DEC-2010	s. Parathyroid						X		X	X					
	t. Peritoneal Membrane						X	X	X	X					
	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: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:21-NOV-2012 DISTRICT: San Francisco PRINTED BY FDA:06-DEC-2012
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION														14. PROPRIETARY NAME(S)		
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps										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				
4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code)	Establishment Functions																
5. ENTER CORRECTIONS TO ITEM 4	Types of HCT / Ps	Recover	Screen	Test	Package	Process	Store	Label	Distribute								
a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	a. Bone						X		X	X							
4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services 7100 N. Financial Drive, Suite 105 Fresno, California 93720 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	b. Cartilage						X		X	X							
	c. Cornea																
	d. Dura Mater																
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous																
	f. Fascia							X		X	X						
	g. Heart Valve																
	h. Ligament									X		X	X				
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous																
	j. Pericardium							X		X	X						
	k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic																
l. Sclera																	
m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous																	
6. MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if applicable, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610	n. Skin								X		X	X					
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic																
8. U.S. AGENT a. E-MAIL _____	p. Tendon								X		X	X					
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO d. DATE 20-NOV-2012	q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic																
	r. Vascular Graft																
	s. Parathyroid								X		X	X					
	t. Peritoneal Membrane								X		X	X					
	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: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:27-NOV-2013 DISTRICT: San Francisco PRINTED BY FDA:09-DEC-2013
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION								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)
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps											
	Types of HCT / Ps	Establishment Functions										
		Recover	Screen	Test	Package	Process	Store	Label	Distribute			
a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	a. Bone							X	X	X		
	b. Cartilage							X	X	X		
	c. Cornea											
	d. Dura Mater											
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous											
	f. Fascia							X	X	X		
	g. Heart Valve											
	h. Ligament							X	X	X		
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous											
	j. Pericardium							X	X	X		
	k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic											
	l. Sclera											
	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous											
	n. Skin							X	X	X		
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic											
	p. Tendon							X	X	X		
	q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic											
	r. Vascular Graft											
	s. Parathyroid							X	X	X		
	t. Peritoneal Membrane							X	X	X		
	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: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:02-DEC-2014 DISTRICT: San Francisco PRINTED BY FDA:22-DEC-2014
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION										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)						
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps																			
	Types of HCT / Ps	Establishment Functions													Recover	Screen	Test	Package	Process	Store
a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____		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 type="checkbox"/> Allogeneic																			
	l. Sclera																			
	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous																			
	n. Skin																			
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic																			
	p. Tendon																			
	q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic																			
	r. Vascular Graft																			
	s. Parathyroid																			
	t. Peritoneal Membrane																			
	u.																			
	v.																			
4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services 7100 N. Financial Drive, Suite 105 Fresno, California 93720 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> 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) Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610																				
7. ENTER CORRECTIONS TO ITEM 6																				
8. U.S. AGENT a. E-MAIL																				
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO																				
	d. DATE 01-DEC-2014																			

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: 3001238554	2. REASON FOR SUBMISSION a. <input type="checkbox"/> INITIAL REGISTRATION / LISTING b. <input checked="" type="checkbox"/> ANNUAL REGISTRATION / LISTING c. <input type="checkbox"/> CHANGE IN INFORMATION d. <input type="checkbox"/> INACTIVE	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:17-NOV-2015 DISTRICT: San Francisco PRINTED BY FDA:03-DEC-2015
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION									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)	
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps													
	Types of HCT / Ps	Establishment Functions												
		Recover	Screen	Test	Package	Process	Store	Label	Distribute					
4. PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code) Community Blood Center dba Community Tissue Services 7100 N. Financial Drive, Suite 105 Fresno, California 93720 a. PHONE 559-224-1168 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	a. Bone							X		X	X			
	b. Cartilage							X		X	X			
	c. Cornea													
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) Community Blood Center dba Community Tissue Services Attn: David M. Smith, MD 349 S. Main Street Dayton, Ohio 45402-2715 a. PHONE 937-461-3450 EXT 3610 7. ENTER CORRECTIONS TO ITEM 6 b. PHONE _____	d. Dura Mater													
	e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
	f. Fascia							X		X	X			
	g. Heart Valve													
	h. Ligament							X		X	X			
	i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
	j. Pericardium							X		X	X			
	k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
	l. Sclera													
	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
8. U.S. AGENT a. E-MAIL _____	n. Skin							X		X	X			
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
	p. Tendon							X		X	X			
	q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
	r. Vascular Graft													
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME David M. Smith, MD b. E-MAIL dsmith@cbccts.org c. TITLE CEO d. DATE 16-NOV-2015	s. Parathyroid							X		X	X			
	t. Peritoneal Membrane							X		X	X			
	u.													
	v.													