

DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE FOOD AND DRUG ADMINISTRATION <b>ESTABLISHMENT REGISTRATION AND LISTING FOR HUMAN CELLS, TISSUES,                  AND CELLULAR AND TISSUE-BASED PRODUCTS (HCT/Ps)</b> (See reverse side for instructions)	<b>1. REGISTRATION NUMBER</b> (FDA Establishment Identifier)  FEI: 3009192463	<b>2. REASON FOR SUBMISSION</b> 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	<b>VALIDATION--FOR FDA USE ONLY</b> VALIDATED BY FDA:24-NOV-2016 DISTRICT: Detroit PRINTED BY FDA:15-DEC-2016
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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)								
<b>3. OTHER FDA REGISTRATIONS</b> a. BLOOD FDA 2830 NO. _____ b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	<b>10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps</b>												
	<b>Types of HCT / Ps</b>	<b>Establishment Functions</b>											
		Recover	Screen	Test	Package	Process	Store	Label	Distribute				
<b>4. PHYSICAL LOCATION</b> (Include legal name, number and street, city, state, country, and post office code) Life Line Stem Cell  720 Broadway New Haven, Indiana 46774  a. PHONE 260-417-4246 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY	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												
<b>5. ENTER CORRECTIONS TO ITEM 4</b>	n. Skin												
	o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
<b>6. MAILING ADDRESS OF REPORTING OFFICIAL</b> (Include institution name if applicable, number and street, city, state, country, and post office code) Life Line Stem Cell Attn: Terri A. Tibbot, MS CTBS 720 Broadway New Haven, Indiana 46774  a. PHONE 260-417-4246 EXT _____	p. Tendon												
	q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input checked="" type="checkbox"/> Allogeneic	X	X		X			X		X			
<b>7. ENTER CORRECTIONS TO ITEM 6</b>	r. Vascular Graft												
	s. Amniotic Membrane	X	X		X			X		X			Placenta
<b>8. U.S. AGENT</b>  a. E-MAIL	t. Placenta	X	X		X			X		X			
<b>9. REPORTING OFFICIAL'S SIGNATURE</b>  a. TYPED NAME Terri A. Tibbot, MS CTBS b. E-MAIL ttibbot@lifelinestemcell.org c. TITLE CEO	u. Amniotic Fluid	X	X		X			X		X			
d. DATE 23-NOV-2016	v. Umbilical Cord	X	X		X			X		X			