

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: 0002275064	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:29-NOV-2017 DISTRICT: New Jersey PRINTED BY FDA:27-JAN-2018
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PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION	11. HCT/Ps DESCRIBED IN 21 OFK 12/1/10	12. HCT/Ps REGULATED AS MEDICAL DEVICES	13. HCT/Ps REGULATED AS DRUGS OR BIOLOGICAL DRUGS	14. PROPRIETARY NAME(S)									
3. OTHER FDA REGISTRATIONS a. BLOOD FDA 2830 NO. FEI: 0002275064 b. DEVICES FDA 2891 NO. _____ c. DRUG FDA 2656 NO. _____	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) Bergen Community Regional Blood Center dba Community Blood Services Elie Katz Umbilical Cord Blood Program New Jersey Cord Blood Bank (NJCBB) 102 Chestnut Ridge Road Montvale, New Jersey 07645 a. PHONE 201-444-3900 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 checked="" type="checkbox"/> Autologous <input checked="" type="checkbox"/> Family Related <input checked="" type="checkbox"/> Allogeneic	X	X	X	X	X	X	X	X	X		X		
	l. Sclera													
	m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
5. ENTER CORRECTIONS TO ITEM 4	n. Skin													
	o. Somatic Cell Therapy Products <input checked="" type="checkbox"/> Autologous <input checked="" type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic	X	X	X	X	X	X	X		X		X		
6. MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if applicable, number and street, city, state, country, and post office code) Blood Systems, Inc. Attn: Gina Ramirez 6210 East Oak Street P.O.Box 1867 Scottsdale, Arizona 85252-1867 a. PHONE 303-363-2221 EXT _____	p. Tendon													
	q. Umbilical Cord Blood <input checked="" type="checkbox"/> Autologous <input checked="" type="checkbox"/> Family Related <input checked="" type="checkbox"/> Allogeneic	X	X	X	X	X	X	X	X	X		X		
7. ENTER CORRECTIONS TO ITEM 6	r. Vascular Graft													
8. U.S. AGENT a. E-MAIL _____	s. Therapeutic Cells	X	X					X					X	
9. REPORTING OFFICIAL'S SIGNATURE a. TYPED NAME Gina Ramirez b. E-MAIL gramirez@bloodsystems.org c. TITLE Regulatory Manager	t.													
	u.													
	v.													
	d. DATE 29-NOV-2017													