


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: 3005875823	<b>2. REASON FOR SUBMISSION</b> 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	<b>VALIDATION--FOR FDA USE ONLY</b> VALIDATED BY FDA:23-JUL-2018 DISTRICT: San Francisco PRINTED BY FDA:14-SEP-2018								
<b>PART I - ESTABLISHMENT INFORMATION</b>		<b>PART II - PRODUCT INFORMATION</b>						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	<b>14. PROPRIETARY NAME(S)</b>	
<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>4. PHYSICAL LOCATION</b> (Include legal name, number and street, city, state, country, and post office code) AlloSource  1700 N Chrisman Rd Tracy, California 95304  a. PHONE 925-963-3525 EXT _____ b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____) c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY		<b>Types of HCT / Ps</b>	<b>Establishment Functions</b>									
<b>5. ENTER CORRECTIONS TO ITEM 4</b>		a. Bone						X		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										
		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>6. MAILING ADDRESS OF REPORTING OFFICIAL</b> (Include institution name if applicable, number and street, city, state, country, and post office code) AlloSource Attn: Trevor Wright 6278 S. Troy Circle Centennial, Colorado 80111  a. PHONE 720-873-4733 EXT _____		n. Skin						X		X	X	
<b>7. ENTER CORRECTIONS TO ITEM 6</b>		o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic										
		p. Tendon						X		X	X	X
		q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic										
<b>8. U.S. AGENT</b>  a. E-MAIL		r. Vascular Graft										
<b>9. REPORTING OFFICIAL'S SIGNATURE</b>  a. TYPED NAME Trevor Wright b. E-MAIL twright@allosource.org c. TITLE Director of Regulatory Affairs d. DATE 23-JUL-2018		s. Amniotic Membrane						X		X	X	
		t.										
		u.										
		v.										