

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: 3009122697	<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	VALIDATION--FOR FDA USE ONLY VALIDATED BY FDA:25-NOV-2015 DISTRICT: San Francisco PRINTED BY FDA:03-DEC-2015										
<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) SightLife  150 North Hill Dr #23 Brisbane, California 94005  a. PHONE 415-330-0900 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>										
		Recover	Screen	Test	Package	Process	Store	Label	Distribute					
<b>5. ENTER CORRECTIONS TO ITEM 4</b>		a. Bone												
		b. Cartilage												
<b>6. MAILING ADDRESS OF REPORTING OFFICIAL</b> (Include institution name if applicable, number and street, city, state, country, and post office code) SightLife Attn: Thomas D. Miller, B.S. CEPT 1200 6th Ave Ste 300 Seattle, Washington 98101  a. PHONE 206-838-4630 EXT _____		c. Cornea	X	X	X	X	X	X	X	X	X			
		d. Dura Mater												
<b>7. ENTER CORRECTIONS TO ITEM 6</b>		e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
		f. Fascia												
<b>8. U.S. AGENT</b>  a. E-MAIL _____		g. Heart Valve												
		h. Ligament												
<b>9. REPORTING OFFICIAL'S SIGNATURE</b> a. TYPED NAME Thomas D. Miller, B.S. CEPT b. E-MAIL tom.miller@sightlife.org c. TITLE VP of Quality and Regulatory Affairs d. DATE 24-NOV-2015		i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
		j. Pericardium												
a. E-MAIL _____		k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
		l. Sclera	X	X	X	X	X	X	X	X	X			
a. E-MAIL _____		m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous												
		n. Skin												
a. E-MAIL _____		o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
		p. Tendon												
a. E-MAIL _____		q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic												
		r. Vascular Graft												
a. E-MAIL _____		s. Amniotic Membrane						X			X	X		AmnioGraft, PROKERA
		t.												
a. E-MAIL _____		u.												
		v.												