

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: 3009122697

2. REASON FOR SUBMISSION
 a. INITIAL REGISTRATION / LISTING
 b. ANNUAL REGISTRATION / LISTING
 c. CHANGE IN INFORMATION
 d. INACTIVE

VALIDATION-FOR FDA USE ONLY
 VALIDATED BY FDA:20-DEC-2017
 DISTRICT: San Francisco
 PRINTED BY FDA:27-JAN-2018

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		
		Types of HCT / Ps		Recover	Screen	Test	Package	Process	Store	Label	Distribute							
a. BLOOD FDA 2830 NO.		a. Bone																
b. DEVICES FDA 2891 NO.		b. Cartilage																
c. DRUG FDA 2656 NO.		c. Cornea		X			X	X	X	X	X							
		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		X			X	X	X	X	X							
		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. Amniotic Membrane							X									
		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)
 SightLife
 Attn: Thomas D. Miller, MS, CEBT
 1200 6th Ave
 Ste 300
 Seattle, Washington 98101

a. PHONE 206-838-4630 EXT
 b. PHONE

7. ENTER CORRECTIONS TO ITEM 6

8. U.S. AGENT

a. E-MAIL
 b. PHONE

9. REPORTING OFFICIAL'S SIGNATURE
 Thomas D Miller
 Thomas D. Miller, MS, CEBT
 tom.miller@sightlifesurgical.com
 VP of Quality and Regulatory Affairs

a. TYPED NAME
 b. E-MAIL
 c. TITLE
 d. DATE 19-DEC-2017