

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: 3010664398

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: Los Angeles
PRINTED BY FDA:27-JAN-2018

PART I - ESTABLISHMENT INFORMATION		PART II - PRODUCT INFORMATION						PART III - ESTABLISHMENT FUNCTIONS		PART IV - REGULATORY INFORMATION					
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		14. PROPRIETARY NAME(S)	
4. PHYSICAL LOCATION (include legal name, number and street, city, state, country, and post office code)		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				
850 Health Sciences Road Suite 2020 Irvine, California 92617		d. Dura Mater													
a. PHONE (949) 854-0800 EXT _____		e. Embryo <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
b. <input type="checkbox"/> SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO. _____)		f. Fascia					X		X		X			TUTOPLAST	
c. <input type="checkbox"/> TESTING FOR MICRO-ORGANISMS ONLY		g. Heart Valve													
5. ENTER CORRECTIONS TO ITEM 4		h. Ligament													
6. MAILING ADDRESS OF REPORTING OFFICIAL (include institution name if applicable, number and street, city, state, country, and post office code)		i. Oocyte <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
SightLife Attn: Thomas D. Miller, MS, CEPT 1200 6th Ave Ste 300 Seattle, Washington 98101		j. Pericardium													
a. PHONE 206-838-4630 EXT _____		k. Peripheral Blood Stem <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic					X	X	X	X	X				
7. ENTER CORRECTIONS TO ITEM 6		l. Sclera		X											
b. PHONE _____		m. Semen <input type="checkbox"/> SIP <input type="checkbox"/> Directed <input type="checkbox"/> Anonymous													
8. U.S. AGENT		n. Skin							X		X			FlexHD Structural	
a. E-MAIL		o. Somatic Cell Therapy Products <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
9. REPORTING OFFICIAL'S SIGNATURE		p. Tendon													
Thomas D. Miller 11291 2018		q. Umbilical Cord Blood <input type="checkbox"/> Autologous <input type="checkbox"/> Family Related <input type="checkbox"/> Allogeneic													
Thomas D. Miller, MS, CEPT tom.miller@sightlifesurgical.com		r. Vascular Graft													
VP of Quality and Regulatory Affairs		s. Amniotic Membrane							X		X			*** See full text on next page	
DATE 19-DEC-2017		t. Umbilical Cord							X		X			*** See full text on next page	
		u. _____													
		v. _____													

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ADDITIONAL INFORMATION:

Proprietary Name(s):

Amniotic Membrane AmnioGraft, PROKERA, Neox Flo, Clarix Flo, Neox 100, Clarix 100
Umbilical Cord Neox Cord 1k, Neox Cord RT, Clarix Cord 1k