See Instructions for OMB Statement. FORM APPROVED:OMB No.0910-0543. Expiration Date: 6/30/2020

DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE

## FOOD AND DRUG ADMINISTRATION ESTABLISHMENT REGISTRATION AND LISTING FOR HUMAN CELLS, TISSUES

FFI: 3000838420

1. REGISTRATION NUMBER (FDA Establishment Identifier)

2. REASON FOR SUBMISSION a. INITIAL REGISTRATION / LISTING | VALIDATED BY FDA:17-NOV-2017 b. X ANNUAL REGISTRATION / LISTING DISTRICT: Dallas

VALIDATION--FOR FDA USE ONLY

AND CELLULAR AND TISSUE-BASED PRODUCTS (HCT/P: (See reverse side for instructions)		PEI: 3009838429			=	c. CHANGE IN INFORMATION d. INACTIVE					PRINTED BY FDA:27-JAN-2018			
PART I - ESTABLISHMENT INFORMATION	PART II - PRODUCT INFORMATION							222	<b>≦</b> ₽12	ᄪᄆᄙᇸ				
3. OTHER FDA REGISTRATIONS	10. ESTABLISHMENT FUNCTIONS AND TYPES OF HCT / Ps													
a. BLOOD FDA 2830 NO	Establishment Functions									T/Ps RIBEI 271.1	PAS!	GC SATE	14. PROPRIETARY	
b. DEVICES FDA 2891 NO.	Types of HCT / Ps		Recover	Screen	Test	Package	Process	Store	Label	Distribute	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	Bi <sup>ω</sup> NAME(S)
c. DRUG FDA 2656 NO													š	
<ol> <li>PHYSICAL LOCATION (Include legal name, number and street, city, state, country, and post office code)</li> </ol>	a. Bone													
PHRK Intervention, Inc dba Southside Device LLC	b. Cartilage													
2058 N KImball Ave Southlake, Texas 76092	c. Cornea													
	d. Dura Mater	_												
a. PHONE 972-979-9392 EXT	e. Embryo	SIP Directed Anonymous												
b. ☐ SATELLITE RECOVERY ESTABLISHMENT (MANUFACTURING ESTABLISHMENT FEI NO  C. ☐ TESTING FOR MICRO-ORGANISMS ONLY	f. Fascia													
5. ENTER CORRECTIONS TO ITEM 4	g. Heart Valve													
	h. Ligament													
MAILING ADDRESS OF REPORTING OFFICIAL (Include institution name if applicable, number and street, city, state, country, and post office code)	i. Oocyte	SIP Directed Anonymous												
PHRK Intervention, Inc d/b/a Southside Device LLC Attn: Jennifer L. Dyer	j. Pericardium													
2058 North Kimball ave Southlake, Texas 76092		Autologous Family Related Allogeneic												
	I. Sclera													
a. PHONE 972-979-9392 EXT	m. Semen	SIP Directed Anonymous												
7. ENTER CORRECTIONS TO ITEM 6 b. PHONE	n. Skin													
	Therapy	Autologous Family Related Allogeneic												
8. U.S. AGENT	p. Tendon													
		Autologous Family Related Allogeneic												
a. E-MAIL	r. Vascular Graft													
9. REPORTING OFFICIAL'S SIGNATURE	s. Amniotic Membr	ane						X		X	X			*** See full text on next page
a. TYPED NAME Jennifer L. Dyer	t.													
b. E-MAIL jdyer@southsidedevice.com	u.													
c. TITLE Owner d. DATE 16-NOV-2017	v.													

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. REGISTRATION NUMBER		2
(FDA Establishment Identifier)		
FEI: 3009838429		
(	FDA Establishment Identifier)	REGISTRATION NUMBER FDA Establishment Identifier)

## ADDITIONAL INFORMATION:

## Proprietary Name(s):

abiodry2,ambio5,epifix,biocover,amnioclear,amniofix,purion,hydrofix, amniotic fluid Amniotic

Membrane

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