



BioDerm, Inc. FDA Establishment Registration & Device Listing 2022

Establishment: BIODERM, INC.
12320 73rd Court North
Largo, FL 33773

Registration Number: 1063299

FEI Number*: 3001949129

Status: Active

Initial Distributor/Importer: Yes

*Note Firm May Have Additional Establishment Types.
Please Review Listings For Further Information.

Date Of Registration Status: 2022

Owner/Operator: BIODERM, INC.
12320 73rd Court North
Largo, FL 33773

Owner/Operator Number: 9032576

Proprietary Name: **CathGrip Double Strap; CathGrip Double Strap with Wing Seal; CathGrip Oval; CathGrip PEG Protect; CathGrip Single Strap; CathGrip Single Strap with Wing Seal; Tube Securement Device; Low Profile CathGrip Single Strap; Low Profile CathGrip Double Strap; Large LVAD Securement Device; Medium LVAD Securement Device**

Classification Name: DEVICE, INTRAVASCULAR CATHETER SECUREMENT

Product Code: KMK

Device Class: 1

Regulation Number: 880.5210

Medical Specialty: General Hospital

Registered Establishment Name: BIODERM, INC.

Registered Establishment Number: 1063299

Owner/Operator: BIODERM, INC.

Owner/Operator Number: 9032576

Establishment Operations: Contract Manufacturer; Manufacturer



Proprietary Name: FreeDerm Adhesive Remover; FreeDerm Bottles; FreeDerm Wipes

Classification Name: SOLVENT, ADHESIVE TAPE
Product Code: KOX
Device Class: 1
Regulation Number: 878.4730
Medical Specialty: General & Plastic Surgery
Registered Establishment Name: BIODERM, INC.
Registered Establishment Number: 1063299
Owner/Operator: BIODERM, INC.
Owner/Operator Number: 9032576
Establishment Operations: Specification Developer

Proprietary Name: Faceplate Seals; Men's Liberty Acute with CathGrip; Men's Liberty Acute with CathGrip and Towel; Men's Liberty Acute without CathGrip; Men's Liberty; Men's Liberty Acute; Men's Liberty Classic; BioDerm XLS (Oval)

Classification Name: DEVICE, PASTE-ON FOR INCONTINENCE, NON-STERILE
Product Code: NOA
Device Class: 1
Regulation Number: 876.5250
Medical Specialty: Gastroenterology
Registered Establishment Name: BIODERM, INC.
Registered Establishment Number: 1063299
Owner/Operator: BIODERM, INC.
Owner/Operator Number: 9032576
Establishment Operations: Contract Manufacturer; Manufacturer

Proprietary Name: BioDerm Penis Clamp; KindKlamp

Classification Name: CLAMP, PENILE
Product Code: FHA
Device Class: 1
Regulation Number: 876.5160
Medical Specialty: Gastroenterology
Registered Establishment Name: BIODERM, INC.
Registered Establishment Number: 1063299
Owner/Operator: BIODERM, INC.
Owner/Operator Number: 9032576
Establishment Operations: Manufacturer



Proprietary Name:

Classification Name:	BioPlus+ Barrier Film
Product Code:	TAPE AND BANDAGE, ADHESIVE
Device Class:	KGX
Regulation Number:	1
Medical Specialty:	880.5240
Registered Establishment Name:	General Hospital
Registered Establishment Number:	BIODERM, INC.
Owner/Operator:	1063299
Owner/Operator Number:	BIODERM, INC.
Establishment Operations:	9032576
	Specification Developer

Proprietary Name:

Classification Name:	Faceplate Strip
Product Code:	BANDAGE, ELASTIC
Device Class:	FQM
Regulation Number:	1
Medical Specialty:	880.5075
Registered Establishment Name:	General Hospital
Registered Establishment Number:	BIODERM, INC.
Owner/Operator:	1063299
Owner/Operator Number:	BIODERM, INC.
Establishment Operations:	9032576
	Manufacturer