

Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2021						Introduction Type	e: Post Launch Change		4 Final Version			Date:	6/24	4/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STOR	RAGE REQUI	REMENTS*		
Company Name:	Camber Pharmace	uticals, Inc.				Application	n: ANDA	a. Temperati	ure - Indicate the USP temp	erature range for t	his product.			
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 204793						1	Temperature Range	Controlled Room		and 25 C (68	3° – 77° F)			
Medical Device Class, if applica														
DUNS:	11-856-3719							_	Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Nai	me: Silodo	osin Capsules 8 mg					I	(write in)					
Selling Unit NDC:	31722-636-90		Unit of Use NDC:		31722-636-90		31722636902		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Silodosin Capsules	s 8 mg						T	Is this product to be shippe	d to customers on i	ce?		No	7
_	·								Is this product to be shippe				No	1
Active Ingredient(s):		Silodosin												
								b. Contact fo	r temperature excursion qu	estions:				
URL for Additional Product Inform		www.camberpharm	a.com					1	Name:		Soma Raju			
Address:	800 Centennial Av	e, Suite 1			State:	Address 2:	00054	-	Number:		732-529-042			
City: Key Contact:	Piscataway Customer Service				Email:	customerservice@ca	Zip: 08854	Group E-mail: somaraju@heterousa.com						
Phone Number:	1-866-827-3647				Fax:	732-562-8788	amberphama.com	c Special re	gulations for product in any	states?			No	٦
Product Therapeutic Classification		Alpha-1 adrenergic	receptor antagonist		ı ux.	102 002 0100		C. Opeciai ie	Special returns requiremen				No	-
Troduct Therapeutic Glassification	л.	/ lipria adienergie	receptor antagonist						opecial returns requiremen	is for this product:			140	_
	ADDITIO	NAL PRODUCT IN	IFORMATION			PRODUCT DES	SCRIPTION INFORMATION	d Store prod	duct (unit of sale) upright?				No	٦
	7,551110			Direct-Ship On	le c	1 1100001 021		u. otore prot						4
The product is?		NI.	Is the Product	Unit of Use	ily		20 -1	. 01-1/17	Protect product (unit of s	ale) from light?			No	
a legend device? if yes, enter class #		No	Is the Product Orphan Drug Status	Offic of Ose		Size:	90 ct	e. Shelf life:	Initial shelf life at launch	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				8 mg		illitiai Sileli ille at iaulicii	ii dinerent).				Months
if yes, list NDCs of		140	FDA Approval Status			Strength:	5g			ORDER INFOR	MATION			
component parts						B	Hard gelatin capsule							
reverse numbered?		No				Dosage Form:			Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						x Bottle		1 Bottle of 9	0 Capsules		
latex-free?		Yes				Product Shape:	Capsule		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes				i roduct onapc.			Ampule					
correctional institution block?		No				Product Color:	White cap and white body		Glass		Minimum o	rder quantity	y?	Yes
opioid?		No							Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint	Imprinted with 'H' on cap		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for		In this was dead account do	and an thin		•	and 'S2' on body		Vial Liquid Multi				ich package	type?
hospital scanning?			Is this product covered under the Trade Agreements Act (No				Vial Powder Sgl Vial Powder Multi		24	Each Inner/Cartor	-/Dools	
If Unit Dose, indicate NDC here:			Trade Agreements Act (IAA)!	NO				Other: Write In			Case	1/Pack	
			FOR GENERIC DRUG PR	ODUCTS					Other: Write III			Ousc		
			TOR GENERIC DROG FR	000013										
					Aut	norized Generic *If	Authorized Generic, other		PI	ARMACY ORDER	/ BILL UNIT			
L Oranga Baak Batings	AB				7100		ection fields are not applicable	Pac sall uni	t to customer?					
I. Orange Book Rating: II. Generic Equivalent to What Bra		Rapaflo						Nec. sen um	t to customer:		Rx billing u	Each	acy:	
ii. Generic Equivalent to what Bra	anur.	Карапо						(Write-in, e.g	ı 1 Vial)			Gram		
		DRUG SUPPI	LY CHAIN SECURITY ACT (DSCSA) INFORM	MATION			(**************************************				Milliliter		
				•										
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes	¬ •	GLN:	0331722498975			ITE	I AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No						·					
If yes, select exemption:					GCP:			1		Dimens	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:								-	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		f yes, was ori	ginal product purcha	sed	Item/Each:	0.1	1.75	1.75	4	12.25	1
Is product sold by manufacturer's	s exclusive distribut	or?	Yes		direct from mf	r?			0.1	1.75	1.75	7	12.25	
Has FDA granted waiver/exception		oduct?	No		Provide sourc	e manufacturer for re	epackaged product	Box/Carton/l	Bundle/					
If yes, attach documentation fro	m FDA.							Inner Pack:						
		0.71	IN AND LUDGO DECELICE I	NEODMATION				Case:	2.8	11	7.5	5	412.50	24
		GII	IN AND HIBCC PRODUCT I	NFORMATION				Pallet:						
Saleable Unit of Measure	c,	aleable Quantity	HIBCC		GTIN	14	Unit of Use GTIN-14	Pallet:						
X Item/Each	36	1	TIBOO			1722636902	00331722636902							
Box/Carton/Bundle/Inner Pack					0000		00001122000002		COST INFORMATION			WHOLESAL	ER USE ONL	LY:
X Case		24			3033	1722636903								
Pallet						-		Regular Cos	t		Vendor #:			
								Invoice Cost		\$90.00	Whsl. Code	#:		
											Fineline Co			
								As of date:	8/20/2020					
	_													
								11			ļ			
*Please provide any additional inf			Attach copy of SAFETY DA	ATA SHEET (SDS	s) or non hazar		SERT, LABEL AND PHOTO OF signated Drop Ship Only.	PRODUCT PACK	AGING and BARCODE. Signature:					



Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2021

For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? No	SDS Hazard Classification					
b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive toxicant? No Does the product label bear a CA Prop 65 warning? No	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard					
c. Contact Hazard? d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level:					
Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name	Is the product a NIOSH hazardous drug? If yes, indicate which:					
c. DOT Hazard Class d. Packing Group	Hazardous Waste Identification					
e. Inhalation Hazard? Is this product regulated for shipment by IATA? No	EPA Hazardous Waste Code: Waste Characteristics					
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS					
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:					
Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)					
Is this a reportable quantity? No RQ Threshold: Is this a marine pollutant? No Is this product shipped utilizing an authorized DOT exception or Special Permit? No (if yes, identify method below) Limited Quantity Consumer Commodity, ORM-D Small Quantity (49 CFR 173.4)	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: No Phone: DEA #: NCPDP#: NPI #:					
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments					
ADD'L STORAGE INFORMATION	Registry: Registry Program Contact Name: Comments No Phone:					
Is the Product						
Controlled Substance Code Controlled by State(s)? ARCOS Reportable? Schedule No. No Controlled Substance Code Listed Chemical (List I or II) No If yes, indicate which: Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION:	RETURN INSTRUCTIONS Contact tel. # if product received damaged: I-866-827-3647 Is product returnable for credit: Yes					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	URL/Link to returns policy: contact - customerservice@camberpharma.com					
Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: No	Special regulations or returns requirements for this product in certain states?					
Restricted from US territories? (explain in comments) No Comments:	If so, which states? Other requirements? Comments?					
MISCELLANE	DUS NOTES and/or Image of Product Barcode:					



Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2021

FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method fo	r Designated Drop Ship Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity:	Fax Number: Fax Number: Phone No.: Site Address:	Shipping lead time of PO: Hours Days Ships same day for next day receipt: Ships for second day receipt:
F	Name: Phone:	Ships regular ground for 3-10 days receipt:
Expedited Freight Charg	ges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order:		Overnight receipt available:
Drop Ship service fee billed with each order:		PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:		Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
		Priority Overnight receipt available:
Class	of Trade Restriction:	PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy only: Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician of Restricted from US territories? (explain in conficulty) Comments:	offices only:	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Infor	rmation Required to Process PO:	Return Instructions
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:		Contact # if product is received damaged: Is product returnable for credit: URL/Link to returns policy: Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?
Mis	scellaneous Notes:	
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?