

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

Version 2021						Introduction Ty	/pe:	New Item		x Final Version			Date:	6/23	3/2024
			PRODUCT INFORMA	TION						SPECIAL HAN	IDLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA						a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 213812							Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)								
Medical Device Class, if applicable:															
DUNS:	11-856-3719									Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Na	me: Eplei	renone Tablets 25 mg						I	(write in)					
Selling Unit NDC:	31722-049-90		Unit of Use NDC		31722-049-90		331722049	900		Notes					
UDI			CVX Code:			MVX Code:									
Description: Eplerenone Tablets 25 mg Is this product to be shipped to customers on ice? No								1							
_										Is this product to be shippe				No	1
Active Ingredient(s):		Eplerenone													
							b. Contact for temperature excursion questions:								
URL for Additional Product Inform		www.camberpharr	na.com		1					Name:		Soma Raju			
Address:	800 Centennial A	ve, Suite 1			State:	Address 2:	71	054		Number:		732-529-042			
City:	Piscataway Customer Service				Email:	customerservice@	Zip: 088			Group E-mail:		somaraju@f	heterousa.coi	<u>n</u>	
Key Contact: Phone Number:	1-866-827-3647	*			Fax:	732-562-8788	camperpria	illia.com	a Special re	gulations for product in any	ctotoc?			No	7
Product Therapeutic Classification		Salactive aldoster	one receptor antagonist antih	vnertensive	I ax.	732-302-0700			c. Special re	Special returns requiremen				No	-
Product Therapeutic Classification	on:	Selective aldoster	one receptor antagonist antii	yperterisive						Special returns requiremen	is for this product?			INO	_
	ADDITI	ONAL PRODUCT II	NEORMATION			PRODUCT DE	ESCRIPTIO	ON INFORMATION	d Store pro	luct (unit of sale) upright?				No	7
	ADDIII	OWAL I RODUCT II		Direct-Ship	Omles	I RODUCI DI	_ookii TiC	AT MI ORMATION	u. Store prod						1
The product is?		NI.	Is the Product	Unit of Use	Jniy		00 -		. 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	Offit of Ose		Size:	90 ct	t	e. Shelf life:	Initial shelf life at launch	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				25 m	na		initial shell life at launch	ir amerent):				Wonths
if yes, list NDCs of		INO	FDA Approval Status			Strength:	2011	19			ORDER INFORM	MATION			
component parts			1 2717 pp. oral olatao				Film	coated tablet							
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 Tablets		
latex-free?		Yes	Dairy Lac	tose, Casein		Product Shape	Roui	nd, biconvex		Box/Carton		(Write-in, e.	.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Dull y, Luc	tose, ouseni		1 Todatet Griape	·.			Ampule					
correctional institution block?		No				Product Color	r: Light	t yellow		Glass		Minimum o	rder quantity	/?	Yes
opioid?		No								Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprii		ossed with 'V' on one side 68' on the other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for		to the overdent consent	or don't be			anu	bo on the other side		Vial Liquid Multi				ich package	type?
hospital scanning?			Is this product covered Trade Agreements Act		No					Vial Powder Sgl Vial Powder Multi		24	Each	/Deels	
If Unit Dose, indicate NDC here:			Trade Agreements Act	IAA)!	INO					Other: Write In			Inner/Cartor Case	/Pack	
			FOR GENERIC DRUG PF	ODUCTS					<u> </u>	Other: Write III			Odsc		
			TOK GENERIC DROG FI	1000013											
					Aut	horized Generic *	*If Authorize	ed Generic, other		PI	HARMACY ORDER	/ BILL UNIT			
L Oranga Baak Batings	AB			_	7.00			ds are not applicable	Pac sell uni	to customer?					
I. Orange Book Rating: II. Generic Equivalent to What Bra		Inspra						.,	itec. sen um	to customer:		KX billing u	nit to pharm Each	acy:	
ii. Generic Equivalent to what Bra	anu r.	шэрга							(Write-in, e.g	1 Vial)			Gram		
		DRUG SUPF	PLY CHAIN SECURITY ACT	(DSCSA) INFO	RMATION				(**************************************				Milliliter		
				,											
Does supplier meet DSCSA defini	ition of manufactur	rer?	Yes		GLN:	0331722498975				ITEI	II AND PACKING II	NFORMATIO	N		
Is product exempt from DSCSA?			No							· · · · · · · · · · · · · · · · · · ·					
If yes, select exemption:					GCP:				1		Dimensi	ons (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:									· [Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product purch	hased		Item/Each:	0.07	1.57	1.57	3.13	7.72	1
Is product sold by manufacturer's			Yes		direct from m						1.07	1.07	3.13	1.12	'
Has FDA granted waiver/exception		oduct?	No		Provide source	e manufacturer for	repackage	d product	Box/Carton/	Bundle/					
If yes, attach documentation fro	m FDA.								Inner Pack:						
		67	TIN AND LUDGE BRODUCT	NEODMATION					Case:	2.1	10	6.75	4.25	286.88	24
		G	TIN AND HIBCC PRODUCT	NEORWATION					Pallet:						
Saleable Unit of Measure	c	Saleable Quantity	HIBCC		GTIN	.1.1.4	Llei	it of Use GTIN-14	Pallet:						
X Item/Each	3	1	THECC			31722049900		331722049900							
Box/Carton/Bundle/Inner Pack					0000		300			COST INFORMATION			WHOLESAL	ER USE ONL	Y:
X Case		24			3033	31722049901									
Pallet									Regular Cos	t		Vendor #:			
									Invoice Cost	(WAC) (\$)	\$75.00	Whsl. Code	#:		
									[]			Fineline Co	de:		
									As of date:	8/28/2023		ļ			
									[]						
ļ.									Ц			<u> </u>			
			Attach copy of SAFETY D	ATA SHEET (SI	DS) or non hazar				PRODUCT PACK						
*Please provide any additional inf		2				See new p. 3 for D	hotennian	Dron Shin Only		Signature:					



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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:						



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

Order Method for Designated Drop S	nip 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: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	Shipping lead time of PO: Hours Days Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:
Expedited Freight Charges or Other Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order:		Overnight receipt available: 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, hospitals, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess 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?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?