

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

Version 2024						Introduction Ty	vpe: New Item	x	Final Version			Date:	3/17/	/2025
			PRODUCT INFORMAT	ION					SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA		a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN	DA/BLA; PMA/510	(k): 218	3110			NDA 505(b) Type:	NOT APPLICABLE		rature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat	ole:													
DUNS:	11-856-3719							Other	Femperature Range	Requirement	Excursions p	ermitted to 1	5°C to 30°C (	59°F to
Proprietary Name (If Applicable) a		me: Phe	enylephrine Hydrochloride Injecti	ion, USP 100 m					write in)		86°F)			
Selling Unit NDC:	31722-345-10		Unit of Use NDC:		31722-345-10		331722345101	Notes						
UDI			CVX Code:			MVX Code:								
Description:	Phenylephrine Hy	drochloride Injectio	ion, USP 100 mg/10 mL (10 mg/r	mL) Pharmacy	Bulk Package			Is this	product to be shippe	d to customers on i	ce?		No	1
								Is this	product to be shippe	d to customers on o	dry ice?		No	
Active Ingredient(s):		Phenylephrine h	ydrochloride, USP											
UDI for Additional Draduat Inform		www.camberpha						b. Contact for temper Name:		estions:	Soma Raju			
URL for Additional Product Inform Address:	800 Centennial Av		ma.com			Address 2:		Numb			732-529-042	3		
	Piscataway	re, Suite i			State:	NJ	Zip: 08854						n	
Key Contact:	Customer Service				Email:		camberpharma.com	Group E-mail: somaraju@heterousa.com			<u></u>			
Phone Number:	1-866-827-3647				Fax:	732-562-8788		c. Special regulations for product in any states?			No	1		
Product Therapeutic Classification		Alpha-1 adrener	gic receptor agonist						I returns requirement				No	
		1												
	ADDITI	ONAL PRODUCT	INFORMATION			PRODUCT D	ESCRIPTION INFORMATION	d. Store product (uni	t of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship C	nlv				t product (unit of sa	ale) from light?			No	1
a legend device?		No	Is the Product	Unit of Use			1 x 10 mL pharmacy bulk	e. Shelf life:	i pi oddor (dinit or ot	,			24	Months
if yes, enter class #		110	Orphan Drug Status			Size:	package vial		shelf life at launch (	if different):				Months
a product kit?		No				Strength:	100 mg/10 mL (10 mg/mL)							1
if yes, list NDCs of			FDA Approval Status			Strength.				ORDER INFORM	MATION			
component parts						Dosage Form	Sterile, clear solution							
reverse numbered?		No				, s		Unit of				NDC selling		
co-licensed?		No	Allergens Present				NI/A		Bottle		1 Box of 1 Ph			
latex-free? preservative-free?		Yes				Product Shap	e: N/A	x	Box/Carton Ampule		(vvrite-in, e.	g. 1 Box of 10	u viais)	
correctional institution block?		Yes No					Colorless	x	Glass		Minimum o	der quantity	2	Yes
opioid?		No				Product Color			Tube		Minimum of	uer quantity	•	103
Cannabinoid?		No	Country of Origin	India		Brederick	N/A		Vial Liquid Sql					
If Unit Dose, is item bar coded to u	nit dose for		, ,			Product Impri	nt:	X	Vial Liquid Multi		If Yes, how	many of whi	ch package	type?
hospital scanning?			Is this product covered ur						Vial Powder Sgl		1	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (T	AA)?	No				Vial Powder Multi			Inner/Carton	/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PRO	DUCTS										
					•		the Authorized Oceansis with an		DL					
				-	Au		*If Authorized Generic, other section fields are not applicable	PHARMACY ORDER / BILL UNIT						
	AP	Manager						Rec. sell unit to customer?		Rx billing unit to pharmacy:				
II. Generic Equivalent to What Bran	nd?:	Vazculep						(Write-in, e.g. 1 Vial)				Each Gram		
		DRUG SUP	PPLY CHAIN SECURITY ACT (	DSCSA) INFOR	MATION			HCPCS J-Code:				Milliliter		
		51100 001						J23	71			Willinger		
Does supplier meet DSCSA definit	tion of manufactur	er?	Yes		GLN:	0331722498975		020		AND PACKING I	NFORMATIO	ا		
Is product exempt from DSCSA?			No											
If yes, select exemption:				_	GCP:					Dimensi	ions (US msn	its.)	Volume	Saleable #
Other exemption - Write in:					-				Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product purcl	hased	Item/Each:	0.07	1.7	1.7	2.6	7.51	1
Is product sold by manufacturer's			Yes		direct from m				0.07	1.7	1.7	2.0	1.01	
Lies CDA suggested sugly and suggestion					Provide source	e manufacturer for	repackaged product	Box/Carton/Bundle/						
Has FDA granted waiver/exception		oduct?	No					Inner Pack:						
If yes, attach documentation from		oduct?	NO					0				6.5	786.5	72
		·						Case:	6.2	11	11	6.5		
		·	TIN AND HIBCC PRODUCT IN	IFORMATION					6.2	11	11	6.5		
If yes, attach documentation from	n FDA.	C	GTIN AND HIBCC PRODUCT IN	IFORMATION	GTI	N-14	Unit of Use GTIN-14	Case: Pallet:	6.2	11	11	6.5		
		G		IFORMATION	GTI	N-14	Unit of Use GTIN-14		6.2	11	11	6.5		
If yes, attach documentation from	n FDA.	C	GTIN AND HIBCC PRODUCT IN	IFORMATION		N-14 81722345101	Unit of Use GTIN-14 00331722345101		6.2	11				
If yes, attach documentation from	RFID tag(Y/N)	G Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION				Pallet:	6.2 ST INFORMATION	11			ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	RFID tag(Y/N)	G Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003			Pallet:		11			ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003	31722345101		Pallet: CC Regular Cost	ST INFORMATION		Vendor #:	WHOLESALI	ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003	31722345101		Pallet:	ST INFORMATION		Vendor #: Whsl. Code	WHOLESAL	ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003	31722345101		Pallet: CC Regular Cost Invoice Cost (WAC) (	ST INFORMATION		Vendor #:	WHOLESAL		Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003	31722345101		Pallet: CC Regular Cost	ST INFORMATION		Vendor #: Whsl. Code	WHOLESAL		Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	GTIN AND HIBCC PRODUCT IN	IFORMATION	003	31722345101		Pallet: CC Regular Cost Invoice Cost (WAC) (	ST INFORMATION		Vendor #: Whsl. Code	WHOLESAL	ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	n FDA. RFID tag(Y/N)	Saleable Quantity	STIN AND HIBCC PRODUCT IN HIBCC		203	31722345101	00331722345101	Pallet: CC Regular Cost Invoice Cost (WAC) ( As of date:	ST INFORMATION \$) 3/11/2025		Vendor #: Whsl. Code	WHOLESAL	ER USE ONL	Y:
If yes, attach documentation from Saleable Unit of Measure	RFID tag(Y/N)	C Saleable Quantity 1 72 72	STIN AND HIBCC PRODUCT IN HIBCC		203	31722345101 31722345105 rd letter, PACKAGE I		Pallet: CC Regular Cost Invoice Cost (WAC) ( As of date:	ST INFORMATION S) 3/11/2025 nd BARCODE.		Vendor #: Whsl. Code	WHOLESAL		Y:

## **HDA** Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2024 For Designa	ted Drop Ship Only Products, Please Use Page 3
MATERIAL HA	ZARD CLASSIFICATION and TRANSPORTATION
Is this product (check all that apply): a. Cytotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? No Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? 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? Is this product regulated for shipment by DOT? No	x       Organic       Corrosive         Inorganic       Oxidizer         Steroid/Androgen       Contact Hazard         Does the product have an Aerosol class? If yes, identify NFPA Storage Level:       No         NFPA Storage Level:       Image: Contact Hazard
(if yes, answer a-e below and provide SDS)         a. UN/Identification Number         b. Proper Shipping Name         c. DOT Hazard Class         d. Packing Group         e. Inhalation Hazard?	Is the product a NIOSH hazardous drug? If yes, indicate which: Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics
Is this product regulated for shipment by IATA? No (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	REMS or REGISTRY RESTRICTIONS         Is there a REMS on this product?       No         If Yes, is it managed with a pharmacy registry?       No         Website URL:       Image: Colored and Col
Is the product restricted for air shipment? If so, indicate restriction:           Passenger         No           Cargo         Passenger & Cargo           Is this a reactable quantity?         No	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)  REMS: No
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) Special Permit; DOT-SP Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	REMS:     No       REMS Program Manager Name:     Phone:       Supplier Manages REMS registry exclusively:     Phone:       Wholesale distributor support:     Provider Name:       Site Enrollment Number assigned     DEA #:       by Supplier:     NPI #:
SP#	Registry:     No       Registry Program Contact Name:     Phone:
ADD'L STORAGE INFORMATION Is the Product Controlled Substance Code	Comments RETURN INSTRUCTIONS
Controlled Substance?         No         Controlled Substance Code           Controlled by State(s)?         No         Listed Chemical (List I or II)         No           ARCOS Reportable?         No         If yes, indicate which:         No           Schedule No.         Is it a scheduled listed chemical product?:         No	KEI URN INSTRUCTIONS       Contact tel. # if product received damaged:     1-866-827-3647       Is product returnable for credit:     Yes       URL/Link to returns policy:     Yes
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	contact - customerservice@camberpharma.com
Restricted to retail pharmacy only:     No       Restricted to hospital, clinics, and physician offices only:     No       Restricted from US territories? (explain in comments)     No	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?
MISCELLANE	OUS NOTES and/or Image of Product Barcode:



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

Version 2024	FOR DESIGNATED DROP SHIP PRODUCT ONLY - if r	ot a designated drop ship, do not complete.	
Order Method for Des	signated Drop Ship Product	Standard Order Receipt and Processing	
Purchase orders may be accepted by: a. EDI b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name Phone Expedited Freight Charges o		Purchase order daily receipt cut off time by supplier         Cut off time:         Shipping lead time of PO:         Hours         Da         Ships same day for next day receipt:         Ships for second day receipt:         Ships regular ground for 3-10 days receipt:         Overnight and Priority Overnight PO Processing	ays
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:		Days of week overnight is available: Monday	
Comments:	s only:	Days of week overnight is available:       Tuesday         Tuesday       Wednessi         Priority Overnight receipt available:       Friday         PO Receipt Cut off time:       PO Receipt Cut off time:         Order receipt method:       Phone:         Fax:       EDI:         Overnight Fees apply:       Policity	/ day
Other Data Informati	ion 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?	
Miscell	aneous Notes:		
		ADDITIONAL INFORMATION	
		Is product order for scheduled patient procedure? Is product order for restocking purposes?	