

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

Version 2021				Introduction	Type: Post Launch Ch	hange	X	Final Version			Date:	6/23/	/2024
PRODUCT INFORMATION						SPECIAL HANDLING AND STOR			AGE REQUIR	EMENTS*			
Company Name: Camber Pharmaceuticals, Inc. Application:							a. Temperature – Indica	ate the USP tempe	erature range for t	his product.			
Application Number for NDA/AN		ed device):	040901						Controlled Room		and 25 C (68	8° – 77° F)	
Medical Device Class, if applicable:													
DUNS:	11-856-3719						Other Te	mperature Range I	Requirement				
Proprietary Name (If Applicable) and		Hydralazine Hydrochloride Tablets					(wr	ite in)					
J	31722-521-01	Unit of Use NDC:		UPC:	331722521017		Notes						
UDI		CVX Code:		MVX Code:									
Description:	Hydralazine Hydrochloride Tab	lets, USP 50 mg						oduct to be shipped				No	
Is this product to be shipped to customers on dry ice? No													
Active Ingredient(s): Hydralazine hydrochloride, USP b. Contact for temperature excursion questions:													
URL for Additional Product Inform	action: www.comb	erpharma.com					b. Contact for tempera Name:	ture excursion qu	estions:	Soma Raju			
	800 Centennial Ave, Suite 1	erphama.com		Address 2:			Number			732-529-042	3		
	Piscataway State:			NJ	Zip: 08854		Group E			somaraju@h		n	
	Customer Service Email:			customerservice	omerservice@camberpharma.com								
Phone Number:	1-866-827-3647		Fax:	732-562-8788			c. Special regulations for product in any states?					No	
Product Therapeutic Classification													
	ADDITIONAL PROD	UCT INFORMATION		PRODUCT	DESCRIPTION INFORMAT	TION	d. Store product (unit o	of sale) upright?				No	
The product is?		Is the Product	Direct-Ship Only					product (unit of sa	ale) from light?			No	
a legend device?	No	Is the Product	Neither	Size:	100 ct		e. Shelf life:					24	Months
if yes, enter class #		Orphan Drug Status					Initial sh	elf life at launch (if different):				Months
a product kit? if yes, list NDCs of	No	FDA Approval Status		Strength:	50 mg				ORDER INFORM				
component parts		FDA Approval Status			Tablet				ORDER INFORM				
reverse numbered?	No			Dosage For	rm:		Unit of S	Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Present						Bottle		1 Bottle of 10			
latex-free?	Yes			Product Sh	Round			Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?	Yes			Floudet Sh	ape.			Ampule					
correctional institution block?	No			Product Co	Orange			Glass		Minimum or	der quantity	/?	Yes
opioid?	No				Unscored, debossed with '			Tube					
Cannabinoid?	No	Country of Origin	India	Product Im	print: side and '40' on the other s			Vial Liquid Sgl Vial Liquid Multi		K V = = 1 =			
If Unit Dose, is item bar coded to un hospital scanning?	nit dose for	Is this product covered u	inder the					Vial Powder Sol		If Yes, how 24	Each	сп раскаде	type?
If Unit Dose, indicate NDC here:		Trade Agreements Act (1						Vial Powder Multi			Inner/Carton	/Pack	
		3	/					Other: Write In			Case		
		FOR GENERIC DRUG PRO	ODUCTS										
										_			
				Authorized Generic	*If Authorized Generic, oth			PH	ARMACY ORDER	/ BILL UNIT			
	AA				section fields are not appl	licable	Rec. sell unit to customer? Rx billing unit to pharmacy:						
II. Generic Equivalent to What Brand?: Hydralazine Hydrochloride Tablets (Pliva)							Each						
(Write-in, e.g. 1 Vial) Gram DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION Milliliter													
	DRUG	SUPPLY CHAIN SECURITY ACT (I	DSCSA) INFORMATION				-				Milliliter		
Does supplier meet DSCSA definit	tion of manufacturer?	Yes	GLN:	0331722498975				ITEM	AND PACKING IN	FORMATION			
Is product exempt from DSCSA?		No	-										
If yes, select exemption:			GCP:						Dimensi	ons (US msm	ts.)	Volume	Saleable #
Other exemption - Write in:								Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		No	If yes, was	original product			Item/Each:	0.11	1.75	1.75	3.5	10.72	1
Is product sold by manufacturer's		Yes		direct from mfr?				0.11	1.75	1.75	3.0	10.72	
Has FDA granted waiver/exception		No	Provide so	urce manufacturer	for repackaged product		Box/Carton/Bundle/						
If yes, attach documentation from	m FDA.						Inner Pack:						
-		GTIN AND HIBCC PRODUCT IN					Case:	3.25	11	7.5	4.5	371.25	24
							Pallet:						
Saleable Unit of Measure	Saleable Qua	ntity HIBCC	G	TIN-14	Unit of Use GTIN	I-14							
X Item/Each	1		0	0331722521017									
Box/Carton/Bundle/Inner Pack							COS	INFORMATION		٧	VHOLESALI	ER USE ONL	.Y:
X Case	24		2	0331722521011									
Pallet					_		Regular Cost			Vendor #:	4.		
		-			-		Invoice Cost (WAC) (\$)		\$7.62	Whsl. Code Fineline Co			
							As of date:	4/15/2024		i menne co			
· · · · · · · · · · · · · · · · · · ·													
		Attach copy of SAFETY DAT	TA SHEET (SDS) or non haz	ard letter, PACKAGE	E INSERT, LABEL AND PHO	OTO OF F	PRODUCT PACKAGING an	d BARCODE.					
*Please provide any additional info	ormation on page 2.				or Designated Drop Ship O		Signatu						

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

Version 2021 For Designate	ed Drop Ship Only Products, Please Use Page 3						
MATERIAL HAZ	ZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? 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? 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? No Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS)	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Storage Level: Is the product a NIOSH hazardous drug? No If yes, indicate which: Image: Storage Level:						
a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Code:						
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? Website URL:						
Is the product restricted for air shipment? If so, indicate restriction: No 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) 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: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:						
SP#	Registry: No						
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone: Phone: 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: Is it a scheduled listed chemical product?: No Schedule No. Is it a scheduled listed chemical product?: No No No	Contact tel. # if product received damaged: 1-866-827-3647 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: 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?						
	OUS NOTES and/or Image of Product Barcode:						



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

Version 2021 FOR DESIGNATED DROP SH	IP PRODUCT ONLY - if not a designated drop ship, do not complete.
Order Method for Designated 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:	Purchase order daily receipt cut off time by supplier Cut off time: 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 Designated Drop Ship	Fees: Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available: Image: Comparison of time: PO Receipt cut off time: Image: Comparison of time: Days of week overnight is available: Image: Comparison of time: Days of week overnight is available: Image: Comparison of time: Image: Comparison of time: Image: Comparison of time: Image: Compari
	Priority Overnight receipt available:
Class of Trade Restriction: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	PO Receipt Cut off time: n offices Saturday Overnight receipt available: Order receipt method: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Image: Content of the state of the stat
Other Data Information 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?
Miscellaneous Notes:	
	ADDITIONAL INFORMATION Is product order for scheduled patient procedure? Is product order for restocking purposes?