

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

Version 2024						Introduction Type	New Item		x Final Version			Date:	5/17	/2025	
			PRODUCT INFORMAT	ION					SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*			
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA					ANDA	a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/AN	DA/BLA; PMA/510(k):	211487				NDA 505(b) Type:	NOT APPLICABLE		Temperature Range	Controlled Room		and 25 C (68	° – 77° F)		
Medical Device Class, if applicat															
DUNS:	11-856-3719								Other Temperature Range	Requirement					
Proprietary Name (If Applicable) a		: Hydroco	odone Bitartrate and Acetam	inophen Tablets	, USP 7.5 mg/				(write in)						
Selling Unit NDC: UDI	31722-942-01		Unit of Use NDC:			UPC: 331 MVX Code:	722942010		Notes						
			CVX Code:			WVX Code:								1	
Description:	Hydrocodone Bitartrat	te and Acetaminop	hen Tablets, USP 7.5 mg/32	25 mg					Is this product to be shippe				No		
Active Ingradient(a):	A diverse la ma di anticipitati lu Di antergianzian la Constanzianziane lu Di antergianziane														
Active Ingredient(s): Hydrocodone bitartrate, USP, acetaminophen, USP b. Contact for temperature excursion guestions:															
URL for Additional Product Inform	nation: ww	ww.camberpharma.	com						Name:		Soma Raju				
Address:	800 Centennial Ave, S	Suite 1				Address 2:			Number:		732-529-042	23			
City:	Piscataway				State:		p: 08854	Group E-mail: somaraju@hetero				neterousa.cor	ousa.com		
Key Contact:	Customer Service				Email:	customerservice@car	nberpharma.com					1			
Phone Number:	1-866-827-3647				Fax:	732-562-8788			lations for product in any				*Yes	-	
Product Therapeutic Classification	n: Cor	mbination opioid, and non	n-opioid, non-salicylate analgesic and a	antipyretic					Special returns requiremen	ts for this product?			*Yes		
		AL PRODUCT INF				PRODUCT_DES	CRIPTION INFORMATION	d Store produ	ct (unit of sale) upright?				No	1	
The product is 2	ADDITIONA			Direct-Ship On	dv.	T RODUCT DES	IN IN THE ON THE ONE			ala) faam liadato				1	
The product is? a legend device?	No		Is the Product Is the Product	Neither	''y		100 ct	e. Shelf life:	Protect product (unit of s	are/ from light?			No 24	Months	
if yes, enter class #	INC		Orphan Drug Status			Size:	100 01		nitial shelf life at launch	if different):			24	Months	
a product kit?	No)				Cára na thu	7.5 mg/325 mg								
if yes, list NDCs of			FDA Approval Status			Strength:				ORDER INFOR					
component parts						Dosage Form:	Tablet								
reverse numbered?	No							r.	Unit of Sale			NDC selling	unit?		
co-licensed? latex-free?	No		Allergens Present				Capsule		x Bottle Box/Carton		1 Bottle of 1	g. 1 Box of 1			
preservative-free?	Ye		Corn,	, Dye		Product Shape:	Capsule		Ampule		(write-iii, e	g. I box of f	U VIAIS)		
correctional institution block?	No						Off white/white		Glass		Minimum o	rder quantity	?	Yes	
opioid?	Ye					Product Color:			Tube					·	
Cannabinoid?	No	2	Country of Origin	USA		Product Imprint:	Debossed 'T 258' on one side and plain on other side with bisect line		Vial Liquid Sgl						
If Unit Dose, is item bar coded to u	init dose for					i roduct imprint.	plain on other side with bisect line		Vial Liquid Multi			many of whi	ch package	type?	
hospital scanning?			Is this product covered un						Vial Powder Sgl		24	Each	/D I-		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TA	AA)?	Yes				Vial Powder Multi Other: Write In			Inner/Carton Case	/Pack		
			FOR GENERIC DRUG PRC	DUCTS					Other: Write III		_	Case			
			FOR GENERIC DRUG FRC	000013				-							
				Γ	Au	thorized Generic *If	Authorized Generic, other		PI	ARMACY ORDER	/ BILL UNIT				
I. Orange Book Rating:	AA			1			tion fields are not applicable	Rec. sell unit t	o customer?		Rx billing u	nit to pharma	acv:		
II. Generic Equivalent to What Bra		orco										Each			
								(Write-in, e.g.	Vial)	-		Gram			
		DRUG SUPPLY	CHAIN SECURITY ACT (D	DSCSA) INFORM	NATION			HCPCS J-Code	:	-		Milliliter			
Deep our lies most DSCSA definit	tion of monutootunon?		Yes	т	GLN:	0040060447600			ITE	AND PACKING I		M			
Does supplier meet DSCSA definit Is product exempt from DSCSA?	tion of manufacturer?		No	- '	GLN:	0843368117603			1161	AND PACKING I	NFURMATIO	N			
			110					1		Dimens			Malana	0-1	
If yes, select exemption: Other exemption - Write in:					GCP:			1	Weight Lbs.	Dimens	ions (US msr Width	nts.) Height	Volume (Cube)	Saleable # Pieces	
Is product repackaged?			No		f yes, was or	iginal product purchas	ed	Item/Each:							
Is product sold by manufacturer's	exclusive distributor	?	Yes		direct from m				0.14	1.84	1.84	3.23	10.86	1	
Has FDA granted waiver/exception		ict?	No] _	Provide sourc	ce manufacturer for rep	ackaged product	Box/Carton/Bu	ndle/						
If yes, attach documentation from	n FDA.							Inner Pack:							
		GTIN	AND HIBCC PRODUCT IN					Case:	3.8	12.3	8.3	3.8	387.94	24	
		GHN	AND TIBEC PRODUCT IN	IONMATION				Pallet:							
Saleable Unit of Measure	RFID tag(Y/N) Sa	aleable	HIBCC		GTI	N-14	Unit of Use GTIN-14	, anot							
11	Qu	uantity													
x Item/Each	N	1			0033	31722942010									
Box/Carton/Bundle/Inner Pack									COST INFORMATION			WHOLESALI	ER USE ONL	_Y:	
X Case Pallet	N	24			1033	31722942017		Denvilan Cont			Vendor #:				
Pallet								Regular Cost Invoice Cost (V		\$16.00	Whsl. Code	#•			
										φ10.00	Fineline Co				
								As of date:	7/1/2020		1				
							COT LADEL AND DUOTO OF 5								
*Please provide any additional inf			Attach copy of SAFETY DA	TA SHEET (SDS	 or non hazar 		ERT, LABEL AND PHOTO OF F ignated Drop Ship Only.		SING and BARCODE.						

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

Version 2024 For Designate	ed Drop Ship Only Products, Please Use Page 3					
MATERIAL HAZ	ARD 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? No d. Does this product require special clean-up instructions? No (If yes, attach SDS with special instructions.) No e. Does the product contain DEHP? No Is this product regulated for shipment by DOT? No (if yes, answer a-e below and provide SDS) No	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:					
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics					
Is this product regulated for shipment by IATA? (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? Yes If Yes, is it managed with a pharmacy registry? No Website URL: https://opioidanalgesicrems.com/home.html					
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) Special Permit; DOT-SP Special Permit; DOT-SP Special Permit; DOT-SP	Yes REMS: Yes REMS Program Manager Name: Murali Kuraku Supplier Manages REMS registry exclusively: Murali Kuraku Wholesale distributor support: DEA #: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:					
ADD'L STORAGE INFORMATION	Registry: No Registry Program Contact Name: Phone: Comments Phone:					
Is the Product Controlled Substance? Yes Controlled Substance Code 9193	RETURN INSTRUCTIONS					
Controlled by State(s)? Yes Listed Chemical (List I or II) No ARCOS Reportable? Yes If yes, indicate which:	Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: Yes URL/Link to returns policy:					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes Restricted to retail pharmacy only: No	contact - customerservice@camberpharma.com Special regulations or returns requirements for this					
Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No Comments:	Product in certain states? Yes If so, which states? Other requirements? Comments? DEA Form 222 or its electronic equivalent is required for all returns in all states.					
MISCELLANEC *Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Part 1	DUS NOTES and/or Image of Product Barcode: 1301.72.					



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

Version 2024	FOR DESIGNATED DROP SHIP PRODUCT ONLY -	if not a designated drop ship, do not complete.
Order Method for	r Designated Drop Ship Product	Standard Order Receipt and Processing
. ,	Fax Number: Fax Number: Phone No.: Site Address:	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:
	hone: es 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:
		Priority Overnight receipt available:
No restriction: Select YES if sold to retail phan Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician o Restricted from US territories? (explain in con Comments:	ffices only: ments)	PO Receipt Cut off time: Image: Constraint of the constr
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:	mation Required to Process PO:	Return Instructions Return Instructions Contact # if product is received damaged: Is product returnable for credit: URL/Link to returns policy: URL/Link to returns policy:
Mis	scellaneous Notes:	
		ADDITIONAL INFORMATION Is product order for scheduled patient procedure? Is product order for restocking purposes?