

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

Version 2024						Introduction 1	Type: New Item		x Final Version			Date:	5/17	/2025
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: AND/							tion: ANDA	a. Temperatu	a. Temperature – Indicate the USP temperature range for this product.					
Application Number for NDA/AN			9			NDA 505(b) Type			Temperature Range	Controlled Room		and 25 C (68	8° – 77° F)	
Medical Device Class, if applicat														
DUNS:	11-856-3719								Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Na 31722-949-01	ame: Oxyco	done and Acetaminophen T		g/325 mg	1150	004700040040		(write in)					
Selling Unit NDC: UDI	31722-949-01		Unit of Use NDC: CVX Code:			UPC: MVX Code:	331722949019		Notes					
-	A							_			<u> </u>			1
Description:	Oxycodone and A	cetaminophen Tablet	s, USP 5 mg/325 mg						Is this product to be shippe				No No	-
Active Ingredient(s):														
									r temperature excursion q	uestions:				
URL for Additional Product Inform		www.camberpharm	ia.com						Name:		Soma Raju			
Address:	800 Centennial Av	ve, Suite 1			0 4-4-1	Address 2:			Number:		732-529-04			
City: Key Contact:	Piscataway Customer Service				State: Email:	NJ	Zip: 08854 @camberpharma.com		Group E-mail: somaraju@heterousa.com					
Phone Number:	1-866-827-3647				Fax:	732-562-8788	<u>geamberphanna.com</u>	c. Special regulations for product in any states? *Yes						1
Product Therapeutic Classificatio			agonist, and non-opioid, non-salicylate	analgesic and					Special returns requirement				*Yes	-
		antipyretic								··· ··· ··· ··· ···				
	ADDITIC	ONAL PRODUCT IN	FORMATION			PRODUCT	DESCRIPTION INFORMATION	d. Store prod	luct (unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship O	inly				Protect product (unit of	ale) from light?			No	1
a legend device?		No	Is the Product	Neither		Size:	100 ct	e. Shelf life:		-			24	Months
if yes, enter class #			Orphan Drug Status			0.20.		-11	Initial shelf life at launch	(if different):				Months
a product kit? if yes, list NDCs of		No	FDA Approval Status			Strength:	5 mg/325 mg			ORDER INFORM				
component parts			PDA Approval Status				Tablet			ORDER INFORM	ATION			
reverse numbered?		No				Dosage For	n:		Unit of Sale		What is the	NDC selling	y unit?	
co-licensed?		No	Allergens Present					_	x Bottle		1 Bottle of 1	00 Tablets		
latex-free?		Yes	Dye, Corn, A	lcohol, Animal		Product Sha	Round, biconvex		Box/Carton		(Write-in, e	.g. 1 Box of 1	0 Vials)	
preservative-free? correctional institution block?		Yes		,			White to off white	-	Ampule Glass		Minimum			Vee
opioid?		No Yes				Product Col	or: White to off-white		Tube		winimum d	rder quantity	y?	Yes
Cannabinoid?		No	Country of Origin	USA			Break line on one side and deboss	d	Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for		, ,	1		Product Imp	with 'T 192' on other side		Vial Liquid Multi		If Yes, how	many of wh	ich package	type?
hospital scanning?			Is this product covered u						Vial Powder Sgl		24	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TAA)?	Yes				Vial Powder Mult	i		Inner/Cartor	n/Pack	
			FOR GENERIC DRUG PR						Other: Write In		1	Case		
			FOR GENERIC DRUG PR	000015										
					A	uthorized Generic	*If Authorized Generic, other		Pł	ARMACY ORDER	BILL UNIT			
I. Orange Book Rating:	AA						section fields are not applicable	Rec. sell uni	to customer?		Rx billing u	nit to pharm	acy:	
II. Generic Equivalent to What Bra	and?:	Percocet							Each					
								(Write-in, e.g. 1 Vial) Gram						
		DRUG SUPPL	Y CHAIN SECURITY ACT (DSCSA) INFOR	MATION			HCPCS J-Co	de:			Milliliter		
Does supplier meet DSCSA defini	ition of manufactu	rer?	Yes		GLN:	0843368117603			ITEN	AND PACKING IN	FORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:				147-1-641	Dimensio	ons (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			riginal product		Item/Each:	0.14	1.84	1.84	3.23	10.86	1
Is product sold by manufacturer's			Yes			lirect from mfr?	or repackaged product	Box/Carton/	Dundle/					
Has FDA granted waiver/exceptio If yes, attach documentation fro			110		FI OVIUE SOU		or repackaged product	Inner Pack:	Sulfule/					
,,								Case:	3.11	12.3	8.3	3.8	387.94	24
		GTIN	I AND HIBCC PRODUCT II	NFORMATION					3.11	12.3	0.5	3.0	307.94	24
Onlandski klasti of Managers								Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)		HIBCC		GT	IN-14	Unit of Use GTIN-14							
X Item/Each	N	Quantity 1			003	331722949019		11						
									COST INFORMATION			WHOLESAL	ER USE ONI	LY:
Box/Carton/Bundle/Inner Pack					103	331722949016	1							
	N	24												
Box/Carton/Bundle/Inner Pack	N	24						Regular Cos			Vendor #:			
Box/Carton/Bundle/Inner Pack	N	24					-	Regular Cos Invoice Cost		\$10.07	Whsl. Code			
Box/Carton/Bundle/Inner Pack	N	24					-	Invoice Cost	(WAC) (\$)	\$10.07				
Box/Carton/Bundle/Inner Pack	N	24								\$10.07	Whsl. Code			
Box/Carton/Bundle/Inner Pack	N	24						Invoice Cost	(WAC) (\$)	\$10.07	Whsl. Code			
Box/Carton/Bundle/Inner Pack			Attach copy of SAFETY DA	TA SHEET (SDS	i) or non haza		INSERT, LABEL AND PHOTO (Designated Drop Ship Only.	Invoice Cost As of date:	(WAC) (\$) 3/14/2019	\$10.07	Whsl. Code			

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Version 2024 For De	ignated Drop Ship Only Products, Please Use Page 3
MATERI	L HAZARD CLASSIFICATION and TRANSPORTATION
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?	x Organic Corrosive 0 Inorganic Oxidizer 0 Steroid/Androgen Contact Hazard
d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP?	Io Does the product have an Aerosol class? If yes, indicate which: No Io NFPA Storage Level: No Io Is the product a NIOSH hazardous drug? No
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 Characteristics
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:
Passenger Cargo Passenger & Cargo	Image: Ned Guide Required Yes Limited Distribution Requirement No Comments / Details: (For example, iPledge program?) Image: 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 Perovision (listed in Column 7 of 49 CFR 172.101);	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
Controlled Substance? Yes Controlled Substance Code 9143 Controlled by State(s)? Yes Listed Chemical (List I or II) ARCOS Reportable? Yes If yes, indicate which:	lo Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy:
Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only:	es contact - customerservice@camberpharma.com Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments? DEA Form 222 or its electronic equivalent is required for all returns in all states.
MISCEI *Storage of this product must abide by the federally mandated DEA requirements outlined in 21 0	ANEOUS NOTES and/or Image of Product Barcode: FR Part 1301.72.



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Version 2024 FOR DESIGNATED DROP SHIP 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:	Overnight receipt available: PO Receipt cut off time: Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
Class of Trade Restriction:	Priority Overnight receipt available:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Saturday Overnight receipt available: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?