

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

Version 2024						Introduction 1	ype: New Item		x	Final Version			Date:	5/17	7/2025
			PRODUCT INFORMAT	ION						SPECIAL HAN	DLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA							a.	a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/AN			207419			NDA 505(b) Type				erature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applicat										0					
DUNS:	11-856-3719								Other	Temperature Range	Requirement				
Proprietary Name (If Applicable) a		ame: C	Dxycodone and Acetaminophen Ta	ablets, USP 10 m	ıg/325 mg					(write in)					
Selling Unit NDC:	31722-951-01		Unit of Use NDC:			UPC:	331722951012		Notes	3					
UDI			CVX Code:			MVX Code:									
Description:	Oxycodone and A	cetaminophen 1	Tablets, USP 10 mg/325 mg						Is this	s product to be shippe	d to customers on i	ce?		No	
		1							Is this	s product to be shippe	d to customers on o	try ice?		No	
Active Ingredient(s): Oxycodone hydrochloride, USP, acetaminophen, USP									0						
URL for Additional Product Inform	nation:	www.camberg	abarma com					D.	. Contact for temp Nam	erature excursion qu	lestions:	Soma Raju			
Address:	800 Centennial Av		<u>Sharma.com</u>			Address 2:			Num			732-529-042	23		
City:	Piscataway	o, outo i			State:	NJ	Zip: 08854			p E-mail:			heterousa.com	n	
Key Contact:	Customer Service	•			Email:	customerservice	@camberpharma.com								
Phone Number:	1-866-827-3647				Fax:	732-562-8788		c. :	c. Special regulations for product in any states?					*Yes	
Product Therapeutic Classificatio	on:	Combination ful antipyretic	Il opioid agonist, and non-opioid, non-salicylate a	nalgesic and					Special returns requirements for this product?					*Yes	
															_
	ADDITIC	ONAL PRODUC	CT INFORMATION			PRODUCT	DESCRIPTION INFORMATIO	ON d.	Store product (ur	nit of sale) upright?		No			
The product is?			Is the Product	Direct-Ship Or	nly				Prote	ect product (unit of s	ale) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	100 ct	е.	Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status						Initia	I shelf life at launch	(if different):				Months
a product kit?		No				Strength:	10 mg/325 mg				ORDER INFORM				
if yes, list NDCs of component parts			FDA Approval Status				Tablet					ATION			
reverse numbered?		No				Dosage For	n:		Unit	of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						x			1 Bottle of 1			
latex-free?		Yes		cohol, Animal		Product Sha	Capsule			Box/Carton			.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Dye, com, A	conol, Animai		FIGURE	•			Ampule					
correctional institution block?		No				Product Col	White to off-white			Glass		Minimum o	rder quantit	y?	Yes
opioid?		Yes		1104			Debossed with 'T 194' on o			Tube					
Cannabinoid?	wit doop for	No	Country of Origin	USA		Product Imp	rint: side and plain on other side			Vial Liquid Sgl Vial Liquid Multi		If Yos how	many of wh	ich nackada	tuno?
If Unit Dose, is item bar coded to u hospital scanning?	unit dose for		Is this product covered u	nder the						Vial Powder Sgl		24	Each	ісп раскаўе	e type :
If Unit Dose, indicate NDC here:			Trade Agreements Act (1		Yes					Vial Powder Multi			Inner/Cartor	/Pack	
										Other: Write In			Case		
			FOR GENERIC DRUG PRO	DUCTS									-		
				_											
					Au	thorized Generic	*If Authorized Generic, othe			PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AA						section fields are not applic	able Re	ec. sell unit to cus	tomer?	_	Rx billing u	nit to pharm	acy:	
II. Generic Equivalent to What Bra	and?:	Percocet							Each						
									Write-in, e.g. 1 Vial				Gram		
		DRUG SU	JPPLY CHAIN SECURITY ACT (I	JSCSA) INFORM	ATION			HC	CPCS J-Code:				Milliliter		
Does supplier meet DSCSA defini	ition of manufactu	rer?	Yes		GLN:	0843368117603				ITEM	AND PACKING IN	IFORMATIO	N		
Is product exempt from DSCSA?			No	- `											
If yes, select exemption:					GCP:						Dimensi	ons (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		lf yes, was o	riginal product		Ite	em/Each:	0.14	1.84			10.86	4
Is product sold by manufacturer's			Yes		purchased d	irect from mfr?					1.04	1.84	3.23	10.00	1
Has FDA granted waiver/exceptio		roduct?	No		Provide sour	rce manufacturer f	or repackaged product		ox/Carton/Bundle/						
If yes, attach documentation fro	m FDA.								ner Pack:						
			GTIN AND HIBCC PRODUCT IN	FORMATION					ase:	3.86	12.3	8.3	3.8	387.94	24
				- entitient entitient				Pa	allet:						
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBCC		GTI	N-14	Unit of Use GTIN-1		-						
		Quantity													
X Item/Each	N	1			003	31722951012									
Box/Carton/Bundle/Inner Pack									C	OST INFORMATION			WHOLESAL	ER USE ONI	LY:
X Case	N	24			103	31722951019	-	_							
Pallet							-		egular Cost voice Cost (WAC)	(\$)	¢00.44	Vendor #: Whsl. Code	. #.		
							-	inv	voice cost (WAC)	(*)	\$∠0.14	Fineline Co			
							1	As	s of date:	3/14/2019					
							1								
			Attach copy of SAFETY DAT	TA SHEET (SDS)) or non haza				OUCT PACKAGING	and BARCODE.					
	formation on page	•				0	Designated Drop Ship Onl		Sign	ature:					

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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:				
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:	contact - customerservice@camberpharma.com Special regulations or returns requirements for this 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.				
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?