

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

Version 2021						Introduction 1	Туре:	Post Launch Change	x	Final Version			Date:	6/23/	2024
			PRODUCT INFORMA	TION						SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*	m	
Company Name:	Camber Pharmaceuti	icals, Inc.				Applica	tion:	ANDA	a. Temperature – In	dicate the USP temp	erature range for t	his product.			
Application Number for NDA/ANI									ntrolled Room – between 20 and 25 C (68° – 77° F)						
Medical Device Class, if applicable:															
DUNS:	11-856-3719								Other	Temperature Range	Requirement		permitted to 1	5°C to 30°C (	59°F to
Proprietary Name (If Applicable) and		e: Oxcarba	zepine Tablets, USP 150 r	ng						(write in)		86°F)			
Selling Unit NDC:	31722-023-01		Unit of Use NDC:			UPC:	33172202	23016	Notes	5					
UDI	CVX Code: MVX Code:														
Description: Oxcarbazepine Tablets, USP 150 mg ls this product to be shipped to customers on ice? No															
Active Ingredient(s): Oxcarbazepine, USP Is this product to be shipped to customers on dry ice? No															
b. Contact for temperature excursion questions:															
URL for Additional Product Inform								Name: Soma Raju							
Address:	300 Centennial Ave, Suite 1			Address 2:				ber:		732-529-042					
City: Key Contact:	Piscataway State: Customer Service Email:				NJ Zip: 08854 customerservice@camberpharma.com			p E-mail:		somaraju@r	eterousa.cor	<u>n</u>			
Phone Number:	1-866-827-3647				732-562-8788			c Special regulation	ns for product in any	states?			No		
Product Therapeutic Classification		ntiepileptic				102 002 0100	562-8788 c. Special regulations for product in any states? Special returns requirements for this								
Troduct merupeutic olassification		naophopho							Opec					140	
	ADDITION	AL PRODUCT INF	ORMATION			PRODUCT	DESCRIPT	ION INFORMATION	d. Store product (ur	nit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	nlv					ect product (unit of sa	ale) from light?			No	
a legend device?	No	0	Is the Product	Neither			10	0 ct	e. Shelf life:	for product (unit of st	ic) non ngin.			24	Months
if yes, enter class #		-	Orphan Drug Status			Size:				I shelf life at launch (	if different):				Months
a product kit?	No	0				Strength:	15	0 mg							
if yes, list NDCs of			FDA Approval Status			Strength.					ORDER INFORM	IATION			
component parts						Dosage Form	m: Fili	m-coated tablet							
reverse numbered?	No		All			-				of Sale			NDC selling	unit?	
co-licensed? latex-free?	No		Allergens Present				0	val, biconvex	x	Bottle Box/Carton		1 Bottle of 1	g. 1 Box of 1	) //iolo)	
preservative-free?	Ye					Product Sha	ape:	al, Diconvex		Ampule		(write-iii, e.	g. 1 Dox of 10	5 viais)	
correctional institution block?	No						Bro	own		Glass		Minimum o	der quantity	? [	Yes
opioid?	No					Product Cole	or:			Tube					
Cannabinoid?	No	0	Country of Origin	India		Product Imp	Debo	ossed with 'V' on one side and '7' and '6' nother side seperated by a score line		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	nit dose for					i roudet imp	(func	nother side seperated by a score line ctional scoring) on both sides		Vial Liquid Multi				ch package t	ype?
hospital scanning?			Is this product covered u							Vial Powder Sgl		24	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (	IAA)?	No					Vial Powder Multi Other: Write In			Inner/Carton	/Pack	
			FOR GENERIC DRUG PR	ODUCTS						Other. White In			Case		
			FOR GENERIC DRUG PR	000013					-						
					Au	uthorized Generic	*If Authori	ized Generic, other		PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB						section fie	elds are not applicable	Rec. sell unit to cus	tomer?		Rx billing u	nit to pharma	acv:	
I. Generic Equivalent to What Brand?: Trileptal								Each							
-	·								(Write-in, e.g. 1 Vial	)	_		Gram		
		DRUG SUPPLY	CHAIN SECURITY ACT (	DSCSA) INFOR	MATION								Milliliter		
Does supplier meet DSCSA definit	ion of manufacturor?		Yes	_	GLN:	0331722498975					I AND PACKING I		N		
Is product exempt from DSCSA?		· I	No	-	JLN.	3331722490975					ANDTACKING				
If yes, select exemption:					GCP:				1		Dimensi	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		lf yes, was o	riginal product pur	chased		Item/Each:	0.1	1	1.58		7.79	1
Is product sold by manufacturer's			Yes		direct from n	nfr?					1.58	1.56	3.14	1.19	
Has FDA granted waiver/exception		uct?	No		Provide sour	rce manufacturer fo	or repackag	ged product	Box/Carton/Bundle/						
If yes, attach documentation from	n FDA.								Inner Pack:						
		GTIN	AND HIBCC PRODUCT II						Case:	2.9	9.75	6.75	4.25	279.70	24
		GIN	AND HIDCO PRODUCT II	- OKMATION					Pallet:						
Saleable Unit of Measure	Salea	able Quantity	HIBCC		GT	IN-14	U	Jnit of Use GTIN-14							
X Item/Each		1				331722023016			L	OST INFORMATION					
Box/Carton/Bundle/Inner Pack								C	WHOLESALER USE ONLY:						
X Case		24			203	331722023010	_								
Pallet							-		Regular Cost			Vendor #:			
					_		-		Invoice Cost (WAC)	(\$)	\$30.00	Whsl. Code Fineline Co			
	-						-		As of date:	3/28/2022		Fineline Co	ue:		
	-						-		no or uale.	0,20,2022		1			
												1			
			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza	ard letter, PACKAGE	E INSERT, L	ABEL AND PHOTO OF P	RODUCT PACKAGING	and BARCODE.					
*Please provide any additional info	ormation on page 2.			, -				d Drop Ship Only.		ature:					

## HDA🔾

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

Version 2021 For Designated Drop Ship Only Products, Please Use Page 3							
MATERIAL HAZARD 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? 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 No No No No No No N	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?       Yes						
Is this product regulated for shipment by DOT? No (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class	If yes, indicate which: Hazardous Waste Identification						
d. Packing Group							
e. Inhalation Hazard? Is this product regulated for shipment by IATA? No	EPA Hazardous Waste Code: Waste Characteristics						
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	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:           Passenger           Cargo           Passenger & Cargo	Med Guide Required     No       Limited Distribution Requirement     Image: 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)	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 #:						
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Registry: No						
ADD'L STORAGE INFORMATION	Registry Program Contact Name:     Phone:       Comments						
Is the Product							
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:       If yes, indicate which:       No         Schedule No.       Is it a scheduled listed chemical product?:       No	RETURN INSTRUCTIONS       Contact tel. # if product received damaged:     1-866-827-3647       Is product returnable for credit:     Yes						
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:						
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	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:         No           Restricted from US territories? (explain in comments)         No	product in certain states?     No       If so, which states? Other requirements? Comments?						
Comments:							
MISCELLANE	OUS NOTES and/or Image of Product Barcode:						



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

Version 2021 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: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available:       Image: Comparison of the co
Class of Trade Restriction:	PO Receipt Cut off time:
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?