

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

Version 2021						Introduction <sup>-</sup>	Type:	New Item		x Final Version			Date:	11/20	0/2024	
			PRODUCT INFORMAT	ION						SPECIAL HAND	LING AND STOR	RAGE REQUI	REMENTS*			
Company Name: Camber Pharmaceuticals, Inc.					Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN			vice):	20	14787						Controlled Room		and 25 C (6	8° – 77° F)		
Medical Device Class, if applicat		. , ,	•							,						
DUNS:	11-856-3719								1	Other Temperature Range F	Requirement					
Proprietary Name (If Applicable) a	and Established Nar	me: Lacos	samide Tablets, USP 150 mg							(write in)	·					
Selling Unit NDC:	31722-814-60		Unit of Use NDC:		31722-814-60	UPC:	33172281	14607		Notes						
UDI			CVX Code:			MVX Code:										
Description:	Lacosamide Tablet	s, USP 150 mg							1	Is this product to be shipped	to customers on	ice?		No	1	
•										Is this product to be shipped	to customers on	dry ice?		No		
Active Ingredient(s):		Lacosamide, USP													-	
	b. Contact for temperature excursion questions:															
URL for Additional Product Inform		www.camberphari	ma.com						Name: Soma Raju							
Address:	800 Centennial Ave	e, Suite 1				Address 2:						732-529-0423				
City:	Piscataway				State: Email:	NJ Zip: 08854 customerservice@camberpharma.com				Group E-mail:		somaraju@l	heterousa.co	<u>m</u>		
Key Contact:	Customer Service 1-866-827-3647				Fax:	732-562-8788	e@camberp	narma.com	c. Special regulations for product in any states?						*Yes	
Phone Number:		A - 1' 1 1			rax:	132-302-0100			c. Special reg							
Product Therapeutic Classification	n:	Anticonvulsant								Special returns requirement	s for this product?			No	]	
	ADDITIO	NAL PRODUCT IN	NEOPMATION			PPODUCT	DESCRIPT	ION INFORMATION	d Store produ	uct (unit of sale) upright?				No	1	
	ADDITIO	NALTRODUCTIO		D: . 01: .		TRODUCTI	DESCRIPT	ION IN ORMATION	u. Store prout						]	
The product is?	1		Is the Product	Direct-Ship ( Unit of Use	Only		00	ct	e. Shelf life:	Protect product (unit of sa	le) from light?			No		
a legend device?		No	Is the Product	Unit of Use		Size:	60	Ct	e. Shelf life:	Initial shelf life at launch (	f different):			24	Months Months	
if yes, enter class # a product kit?		No	Orphan Drug Status				15	0 mg		illitiai sileli ille at laulicii (	i dillerent).				WIOTILIS	
if yes, list NDCs of		140	FDA Approval Status			Strength:	13	onig			ORDER INFOR	MATION				
component parts			- Divippioral Gialag				Fil	m coated tablet				-				
reverse numbered?		No				Dosage For	m:			Unit of Sale		What is the	NDC selling	unit?		
co-licensed?		No	Allergens Present							x Bottle		1 Bottle of 6	0 Tablets			
latex-free?	Yes Soy				Product Sha	ane. Ov	al, biconvex		Box/Carton		(Write-in, e	.g. 1 Box of 1	0 Vials)			
preservative-free?		Yes		-,		1 Todact One				Ampule						
correctional institution block?		No				Product Col	lor: Sa	llmon		Glass		Minimum o	rder quantit	y?	Yes	
opioid?		No	0							Tube						
Cannabinoid?		No	Country of Origin	India		Product Imp		bossed with 'J' on one side d '14' on the other side		Vial Liquid Sgl		W. V 1				
If Unit Dose, is item bar coded to u hospital scanning?	init dose for		Is this product covered ur	ador the						Vial Liquid Multi			Each	ich package	type?	
If Unit Dose, indicate NDC here:			Trade Agreements Act (T		No				Vial Powder Sgl Vial Powder Multi			24	Inner/Cartor	/Pack		
ii onit bose, indicate NBC fiere.	I.		Trade rigidements rich (1	701).	IVO					Other: Write In			Case	// ack		
			FOR GENERIC DRUG PRO	DUCTS					1_	Guion William			Jodoo			
			TOR GENERIO DROGTRO	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,												
					Aut	horized Generic	*If Author	ized Generic, other		PH/	ARMACY ORDER	/ BILL UNIT				
I. Orange Book Rating:	AB					section fields are not applicable							Rx billing unit to pharmacy:			
II. Generic Equivalent to What Brand?: Vimpat								Troor com unit	Each							
									(Write-in, e.g. 1 Vial) Gram							
		DRUG SUPPL	LY CHAIN SECURITY ACT (D	SCSA) INFOR	RMATION				( , . 3	,			Milliliter			
													4			
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes		GLN:	0860000397957				ITEM	AND PACKING I	NFORMATIO	N			
Is product exempt from DSCSA?			No													
If yes, select exemption:					GCP:					Weight Lbs.	Dimens	ions (US msn	nts.)	Volume	Saleable #	
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?	[		No		If yes, was or				Item/Each:	0.1	1.6	1.6	3	7.68	1	
Is product sold by manufacturer's			Yes	_	purchased di											
Has FDA granted waiver/exception If yes, attach documentation from		oduct?	No		Provide source	ce manufacturer f	tor repacka	iged product	Box/Carton/B Inner Pack:	undle/						
ir yes, attach documentation from	m FDA.								Case:							
		GTI	IN AND HIBCC PRODUCT IN	FORMATION					Case:	2.84	10	7	4	280	24	
		0	IN AND THEODY NODOGY IN	TORMATION					Pallet:							
Saleable Unit of Measure	Sa	leable Quantity	HIBCC		GTIN	V-14	ι	Jnit of Use GTIN-14	III and							
X Item/Each	Ī	1				31722814607		0331722814607								
Box/Carton/Bundle/Inner Pack			0000						COST INFORMATION			<u> </u>	WHOLESALER USE ONLY:			
x Case		24		2033												
Pallet	_								Regular Cost			Vendor #:				
									Invoice Cost (	(WAC) (\$)	\$17.05	Whsl. Code				
										40/4/0004		Fineline Co	de:			
					-				As of date:	12/1/2024						
									As of date:	12/1/2024						
			Attach copy of SAFETY DAT	A CHEET (CD	S) or non be-ser	Hottor BACKACE	INCEPT	AREL AND BHOTO OF								



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

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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZ	ZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic?  No	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	3D3 Hazaru Giassiilvativii						
Is the product a CA Prop 65 carcinogen?	x Organic Corrosive						
Is the product a CA Prop 65 cereinogen:  No  No	Inorganic Oxidizer						
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard						
c. Contact Hazard?	Does the product have an Aerosol class? If yes, No						
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:						
(If yes, attach SDS with special instructions.)	NFPA Storage Level:						
e. Does the product contain DEHP?							
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug?						
(if yes, answer a-e below and provide SDS)	If yes, indicate which:						
a. UN/Identification Number							
b. Proper Shipping Name							
c. DOT Hazard Class	Hazardous Waste Identification						
d. Packing Group							
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA?							
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number							
b. Proper Shipping Name	Is there a REMS on this product?						
c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?						
d. Packing Group	Website URL:						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required No						
Passenger	Limited Distribution Requirement						
Cargo	Comments / Details: (For example, iPledge program?)						
Passenger & Cargo							
Is this a reportable quantity? No	REMS: No						
RQ Threshold:	REMS Program Manager Name: Phone:						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:						
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:						
No (if yes, identify method below)  Limited Quantity	Provider Name:  Site Enrollment Number assigned  DEA #:  NCPDP#:						
Consumer Commodity, ORM-D	Site Enrollment Number assigned by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)	by Supplier.						
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);							
SP#	Registry: No						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product							
Controlled Substance? Yes Controlled Substance Code 2746	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						
Schedule No. 5 Is it a scheduled listed chemical product?: No	Is product returnable for credit:  Yes						
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:						
	contact - customerservice@camberpharma.com						
	Contact - Customerservice@Camberpharma.com						
Restricted to retail pharmacy only:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:	product in certain states?						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:							
MISCELLANE	OUS NOTES and/or Image of Product Barcode:						
	-						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Pa	II 1301.12.						



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

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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  Fax Number:	Purchase order daily receipt cut off time by supplier Cut off time:
c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number:	Shipping lead time of PO:  Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:
Contracted 3PL company / contact #: Name: Phone:	
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:
Drop Ship service fee billed with each order:	PO Receipt cut off time:
Drop Ship miscellaneous fees billed:  Comments:	Days of week overnight is available:  Monday Tuesday Wednesday Thursday Friday
	Priority Overnight receipt available:
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:  Phone: Fax: EDI:  Overnight Fees apply: Other fees apply:
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?