

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

Version 2021						Introduction	Type:	New Item		x Fina	al Version			Date:	11/26	/2024	
			PRODUCT INFORMAT	ION						S	PECIAL HAND	LING AND STOR	AGE REQUI	REMENTS*			
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA a. T							a Temperatu	a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/AN			vice).	20	09301	7491100		7.11.071	u. remperatu	Temperature		Controlled Room -		and 25 C (68	3° – 77° F)		
Medical Device Class, if applicab			,								9-						
DUNS:	11-856-3719								1	Other Tempe	rature Range F	Requirement	Excursions p	permitted bet	ween 15°C to	30°C (59°F	
Proprietary Name (If Applicable) a	and Established Na	me: Laco	samide Oral Solution, USP 10	mg/mL					1	(write in			to 86°F)			(32.1	
Selling Unit NDC:	31722-627-26				31722-627-26	26 UPC: 331722627269			Notes				Do not freeze				
UDI		CVX Code:			MVX Code:	MVX Code:											
Description: Lacosamide Oral Solution, USP 10 mg/mL Is this product to be shipped to customers on ice? No																	
			•									to customers on d			No		
Active Ingredient(s):		Lacosamide, USP															
·									b. Contact for	r temperature	excursion que	estions:					
URL for Additional Product Inform													ma Raju				
Address:		ennial Ave, Suite 1				Address 2:			Number:				732-529-0423				
City:	Piscataway				State: Email:	NJ	NJ Zip: 08854 customerservice@camberpharma.com			Group E-mail:				somaraju@heterousa.com			
Key Contact: Phone Number:	1-866-827-3647	The state of the s			Fax:	732-562-8788	e@camr	<u>berpnarma.com</u>	c. Special regulations for product in any states?						No		
Product Therapeutic Classification					Fax.	732-302-0700			c. Special reg		-						
Product Therapeutic Classification	n:	Anticonvulsant								Special return	ns requirement	s for this product?			No		
	ADDITIO	NAL PRODUCT II	NEORMATION			PRODUCT	DESCR	RIPTION INFORMATION	d Store prod	luct (unit of sa	le) upright?				No		
<b>T</b>	ADDITIO	MALIKODOOTII		Discret Ohio	0-1-	TRODUCT	DECON	AII TION IN OKIIATION	u. Store prou	-							
The product is? a legend device?		No	Is the Product	Direct-Ship Unit of Use	Only			200 mL	e. Shelf life:	Protect prod	luct (unit of sa	le) from light?			No 24	Months	
if yes, enter class #		INO	Orphan Drug Status	Offic Of Ose		Size:		200 IIIL	e. Sileli ille.	Initial shalf li	ife at launch (i	f different):			Months Months		
a product kit?		No	Orphan Drug Otatus					10 mg/mL		miliai siicii i	ire at laurion (i	r directing.			Months		
if yes, list NDCs of		110	FDA Approval Status			Strength:		10 mg/m2	ORDER INFORMA			ATION					
component parts			•			Dosage For	·m·	Clear, oral solution									
reverse numbered?		No	-			Dosage For				Unit of Sale			What is the	NDC selling	unit?		
co-licensed?		No	Allergens Present							x Bot			1 Bottle of 2				
latex-free?	Yes Gluten, Whey, Soy, Corn, Alcohol, Anima			Animal	Product Shape: N/A			Box/Carton				(Write-in, e.g. 1 Box of 10 Vials)					
preservative-free?		No	Products, S	ugar, Wheat						Am					_		
correctional institution block?		No				Product Co	lor:	Colorless to yellow or		Gla			Minimum o	rder quantity	/?	Yes	
opioid? Cannabinoid?		No No	Country of Origin	India				yellow-brown N/A		Tub	e   Liquid Sgl						
If Unit Dose, is item bar coded to u		INO	Country of Origin	IIIula		Product Imp	print:	INA			Liquid Sgi Liquid Multi		If Yes, how	many of wh	ich nackado	type?	
hospital scanning?	init dosc for		Is this product covered un	nder the							Powder Sgl			Each	ion package	type.	
If Unit Dose, indicate NDC here:			Trade Agreements Act (T		No						Powder Multi			Inner/Carton	/Pack		
											er: Write In			Case			
			FOR GENERIC DRUG PRO	DUCTS													
					Aut	horized Generic		thorized Generic, other	PHARMACY ORDER / BILL UNIT								
I. Orange Book Rating: AA					section fields are not applicable			Rec. sell unit to customer?				Rx billing unit to pharmacy:					
II. Generic Equivalent to What Bra	and?:	Vimpat											Each				
								(Write-in, e.g.	(Write-in, e.g. 1 Vial)				Gram				
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION  Milliliter																	
Does supplier meet DSCSA definition of manufacturer?  Yes  GLN: 0860000397957  ITEM AND PACKING INFORMATION																	
Is product exempt from DSCSA?	mon or manuractur		No	-	JLN.	00000000391951					TILINI	AND I ACKING IN					
If yes, select exemption:					GCP:				1			Dimensio	ons (US msn	nte \	Volume	Saleable #	
Other exemption - Write in:					GUF.				1	V	Veight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No		If yes, was ori	ginal product			Item/Each:					_			
Is product repackaged:	exclusive distribu	itor?	Yes		purchased dir						0.57	2.5	2.5	6.5	40.63	1	
Has FDA granted waiver/exception			No		Provide source	e manufacturer	for repa	ackaged product	Box/Carton/B	Bundle/							
If yes, attach documentation from	m FDA.								Inner Pack:								
									Case:		14.2	15.5	10.5	7.5	1220.63	24	
		GT	IN AND HIBCC PRODUCT IN	FORMATION							1-1.2	10.0	10.0	7.0	1220.00		
II									Pallet:								
Saleable Unit of Measure	Sa	aleable Quantity	HIBCC		GTIN		_	Unit of Use GTIN-14									
X Item/Each	1 00331722627269					00331722627269		COSTLINE	OPMATION			NHOLESALI	ED LISE ON	V·			
Box/Carton/Bundle/Inner Pack  X Case		24 20331722627263			1722627263			COST INFORMATION				WHOLESALER USE ONLY:					
Pallet	24 2033172			11122021203	122621263			Regular Cost									
	T								Invoice Cost			\$111.51	Vendor #: Whsl. Code	#:			
	†											<b>\$111.51</b>	Fineline Co				
									As of date:	5/3	1/2022		1				
													]				
*Please provide any additional inf		•	Attach copy of SAFETY DAT	A SHEET (SD	S) or non hazard			RT, LABEL AND PHOTO OF I	PRODUCT PACK	AGING and BA	RCODE.						



## **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?  b. CA Prop. 65 Carcinogen or Reproductive Toxicant?  Is the product a CA Prop 65 carcinogen?  No	SDS Hazard Classification  x Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?  No  Does the product label bear a CA Prop 65 warning?  No	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	Does the product have an Aerosol class? If yes, identify NFPA Storage Level:  NFPA Storage Level:						
Is this product regulated for shipment by DOT?  (if yes, answer a-e below and provide SDS)  a. UN/Identification Number  b. Proper Shipping Name	Is the product a NIOSH hazardous drug?  If yes, indicate which:						
c. DOT Hazard Class d. Packing Group	Hazardous Waste Identification  EPA Hazardous Waste Code:  Waste Characteristics						
e. Inhalation Hazard?  Is this product regulated for shipment by IATA?  No	EPA Hazardous waste Code:						
(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?  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 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)	REMS:  REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier:  No Phone:  Phone:  DEA #: NCPDP#: NCPDP#: NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments  Registry:  No						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product  Controlled Substance? Yes Controlled Substance Code  Controlled by State(s)? Yes Listed Chemical (List I or II) No	RETURN INSTRUCTIONS  Contact tel. # if product received damaged: 1-866-827-3647						
ARCOS Reportable? Yes If yes, indicate which: Schedule No. 5 Is it a scheduled listed chemical product?: No	Contact tel. # if product received damaged:  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 product in certain states?						
Restricted to hospital, clinics, and physician offices only:  Restricted from US territories? (explain in comments)  No  Comments:	product in certain states?  If so, which states? Other requirements? Comments?						
CONTRIBUTIO.							
MISCELLAND	OUS NOTES and/or Image of Product Barcode:						
Discard any unused lacosamide oral solution remaining after six (6) months of first opening the bottle. oral solution (equivalent to 20 mL) contains 0.32 mg of phenylalanine.	henylketonurics: A 200 mg dose of lacosamide						



## **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?