

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

Version 2024						Introduction T	ype: New Item		x	Final Version			Date:	11/8	/2024
			PRODUCT INFORMAT	TION						SPECIAL HAN	DLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA								a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/AN	DA/BLA; PMA/510	<b>)(k):</b> 217068	3			NDA 505(b) Type:	NOT APPLICABLE	1	Temperat	ture Range	Controlled Room	- between 20	and 25 C (68	° – 77° F)	
Medical Device Class, if applical	1														
DUNS:	11-856-3719									mperature Range F	Requirement	Excursions	permitted to 1	5° to 30°C (5	59° to 86°F)
Proprietary Name (If Applicable) a Selling Unit NDC:	31722-326-01	ame: Lisdexa	amfetamine Dimesylate Che Unit of Use NDC:	wable lablets t	0 mg	UPC:	331722326018		(wri Notes	te in)					
UDI	51722-520-01		CVX Code:			MVX Code:	331722320016		notes						
Description:	Liedevenfetenie	e Dimesylate Chewabl							la thia are	duatta ha ahiana.				Nie	1
Description:	Lisdexamietamine	e Dimesylate Chewabi	le Tablets 60 mg								d to customers on i d to customers on o			No No	
Active Ingredient(s): Lisdexamfetamine dimesylate															
b. Contact for temperature excursion questions:															
URL for Additional Product Inform		www.camberpharma	i.com						Name:			Soma Raju			
Address:	800 Centennial A Piscataway	ve, Suite 1			State:	Address 2: NJ	7'		Number:			732-529-042			
City: Key Contact:	Customer Service	<u>د</u>			Email:		Zip: 08854 @camberpharma.com		Group E-mail: somaraju@heterousa.com						
Phone Number:	1-866-827-3647				Fax:	732-562-8788	<u>Journoorpriama.com</u>		c. Special regulations for product in any states?					*Yes	1
Product Therapeutic Classificatio	n:	Central nervous sys	tem (CNS) stimulant			I				eturns requirement				*Yes	
-															1
	ADDITI	ONAL PRODUCT INF	ORMATION			PRODUCT [	DESCRIPTION INFORMATI	ION	d. Store product (unit o	f sale) upright?				No	]
The product is?			Is the Product	Direct-Ship C	inly				Protect p	product (unit of sa	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				00		Initial sh	elf life at launch (	if different):				Months
a product kit? if yes, list NDCs of		No	FDA Approval Status			Strength:	60 mg				ORDER INFORM				
component parts			T DA Approval Status				Chewable tablet				ORDER INFORM	ATION			
reverse numbered?		No				Dosage Form	n:		Unit of S	ale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present							Bottle			00 Chewable		
latex-free?		Yes	Corn, Alcohol,	Animal, Suga		Product Sha	pe: Diamond, biconvex	۲		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free? correctional institution block?		No					White to off-white			Ampule Glass		Minimum o	rder quantity	2	Yes
opioid?		No No				Product Cold	or:			Tube		Willing	ruer quantity	ſ	Tes
Cannabinoid?		No	Country of Origin	USA		Product Imp	Debossed 'AT' on one s	side and		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for					Froduct imp	'60' on the other side			Vial Liquid Multi			many of whi	ch package	type?
hospital scanning?			Is this product covered u		1					Vial Powder Sgl		24	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (1	AA)?	Yes					Vial Powder Multi Other: Write In			Inner/Carton Case	/Pack	
			FOR GENERIC DRUG PRO	ODUCTS		1				Other. Write III		1	Case		
			TOR GENERIC DRUG FRO	00013											
					Au	thorized Generic	*If Authorized Generic, oth	ner		PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB						section fields are not applied	icable	Rec. sell unit to custom	ner?		Rx billing u	nit to pharma	acy:	
II. Generic Equivalent to What Bra	ind?:	Vyvanse Chewable											Each	-	
									(Write-in, e.g. 1 Vial)				Gram		
		DRUG SUPPLY	Y CHAIN SECURITY ACT (I	JSCSA) INFOR	MATION				HCPCS J-Code:				Milliliter		
Does supplier meet DSCSA defini	tion of manufactu	rer?	Yes	7	GLN:	0860000397957				ITEM	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No	-											
If yes, select exemption:				_	GCP:						Dimensi	ons (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			iginal product			Item/Each:	0.17	2.02	2.02	3.4	13.80	1
Is product sold by manufacturer's			Yes	_	•	rect from mfr?									
Has FDA granted waiver/exceptio If yes, attach documentation from		roduct?	No		Provide sour	ce manufacturer fo	r repackaged product		Box/Carton/Bundle/ Inner Pack:						
in yes, attach documentation fro									Case:						
		GTIN	I AND HIBCC PRODUCT IN	FORMATION						4.12	12.3	8.3	3.8	387.94	24
									Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)		HIBCC		GTI	N-14	Unit of Use GTIN-	-14							
x Item/Each	N	Quantity 1			003	31722326018									
Box/Carton/Bundle/Inner Pack	IN				003	522020010	-		COST	INFORMATION			WHOLESALI	ER US <u>E ONL</u>	Y:
X Case	N	24			103	31722326015	-								
Pallet									Regular Cost			Vendor #:			
							-		Invoice Cost (WAC) (\$)		\$1,107.95	Whsl. Code			
							-		As of date:	9/18/2024		Fineline Co	ae:		
							-		A S OF GALE.	0,10/2024					
			Attach copy of SAFETY DA	TA SHEET (SD	S) or non haza	rd letter, PACKAGE	INSERT, LABEL AND PHO	DTO OF PR	RODUCT PACKAGING and	BARCODE.					
*Please provide any additional inf	ormation on page	2.				See new p. 3 for	Designated Drop Ship Onl	ıly.	Signatur	e:					

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

Version 2024 Fo	or Designated Drop Ship Only Products, Please Use Page 3						
MA	TERIAL 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? Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning?	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? Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/dentification Number	No     Does the product have an Aerosol class? If yes, identify     No       No     NFPA Storage Level:     NFPA Storage Level:       No     Is the product a NIOSH hazardous drug?     No       If yes, indicate which:     If yes, indicate which:						
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics						
Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	No  REMS or REGISTRY RESTRICTIONS  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	No     Med Guide Required       Limited Distribution Requirement     No       Comments / Details: (For example, iPledge program?)     No						
Is this a reportable quantity? No RQ Threshold: No 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:     DEA #:       Provider Name:     DEA #:       Site Enrollment Number assigned     NCPDP#:       by Supplier:     NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments     No       Registry:     No       Registry Program Contact Name:     Phone:       Comments     Phone:						
Is the Product							
Controlled Substance?         Yes         Controlled Substance Code         1205           Controlled by State(s)?         Yes         Listed Chemical (List I or II)         1           ARCOS Reportable?         Yes         If yes, indicate which:         I           Schedule No.         2         Is it a scheduled listed chemical product?:         Is it a scheduled listed chemical product?:           CLASS OF TRADE RESTRICTION:           No restriction:         Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	No     Image: Contact tel. # if product received damaged:     1-866-827-3647       No     Is product returnable for credit:     Yes       URL/Link to returns policy:     contact - customerservice@camberpharma.com						
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:	Yes     Contact - customerservice@camberpharma.com       No     Special regulations or returns requirements for this product in certain states?     Yes       No     If so, which states? Other requirements? Comments?       DEA Form 222 or its electronic equivalent is required for all returns in all states.						
	SCELLANEOUS NOTES and/or Image of Product Barcode:						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Part 1301.72.							

Release DATE



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

Version 2024 FOR DESIGNATED DROP SHIP PRODUCT ONLY - in	f 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         Site Address:    Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name:	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:
Phone:	
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 second
	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:         Order receipt method:       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?