

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

Version 2021						Introduction Ty	/pe: New Item		x Final Version			Date:	7/22/	2024
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STOF	AGE REQUIR	EMENTS*		
Company Name:	Camber Pharmac	ceuticals, Inc.				Applicati	on: ANDA	a Temperature -	Indicate the USP temper	erature range for t	nis product			
Application Number for NDA/AN			ce):	21:	3709				mperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applica			,-											
DUNS:	11-856-3719							Oth	her Temperature Range I	Requirement				
Proprietary Name (If Applicable)		ame: Dextroamph	netamine Saccharate, Amphetamine Aspartate, Dextro	amphetamine Sulfate and Amp	hetamine Sulfate Tablets (Mix	xed Salts of a Single-Entity Amphetamine	Product) 30 mg		(write in)					
Selling Unit NDC:	31722-164-01		Unit of Use NDC:			UPC:	331722164016	No	ites					
UDI			CVX Code:			MVX Code:								
Description:	Devtroamphetam	ine Saccharate Amn	hetamine Aspartate Devtro	amphotomine Su	Ifate and Amph	etamine Sulfate Tabl	ets (Mixed Salts of a Single-Entity	le t	this product to be shipped	d to customers on i	202		No	
Description.	Amphetamine Pro		netamine Aspartate, Dextros	amprietamine ou	illate and Ampri	letarrille Sullate Tabl	ets (Mixed Saits of a Siligle-Effility		this product to be shipped				No	
Active Ingredient(s):			e saccharate, amphetamine	aspartate, dextr	oamphetamine	sulfate, USP, and an	ophetamine sulfate. USP		and product to be emppe	a to odotomoro on t	.,		110	
Active Ingredient(s): Dextroamphetamine saccharate, amphetamine aspartate, dextroamphetamine sulfate, USP, and amphetamine sulfate, USP						b. Contact for temperature excursion questions:								
URL for Additional Product Inform	mation:	www.camberpharm	a.com						me:		Soma Raju			
Address:	800 Centennial A	ve, Suite 1				Address 2:		Nu	ımber:		732-529-042			
City:	Piscataway				State:	NJ	Zip : 08854	Gre	oup E-mail:		somaraju@h	eterousa.cor	<u>n</u>	
Key Contact:	Customer Service	9			Email:		camberpharma.com							
Phone Number:	732-529-0430				Fax:	732-562-8788			tions for product in any				*Yes	
Product Therapeutic Classification	on:	Central nervous sys	stem (CNS) stimulant					Sp	ecial returns requirement	ts for this product?			*Yes	
								_						
	ADDITI	IONAL PRODUCT IN	IFORMATION			PRODUCT D	ESCRIPTION INFORMATION	d. Store product	(unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	Only			Pro	otect product (unit of sa	ale) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	100 ct	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status			Size.		Init	tial shelf life at launch (if different):				Months
a product kit?		No				Strength:	30 mg							
if yes, list NDCs of			FDA Approval Status			ou chigan.				ORDER INFORM	IATION			
component parts						Dosage Form	Tablet							
reverse numbered?		No						Un	it of Sale		What is the		unit?	
co-licensed?		No	Allergens Present				5 1016		x Bottle		1 Bottle of 10			
latex-free?		Yes	Dye, Corn,	Alcohol, Sugar		Product Shap	Round, flat faced, beveled edge		Box/Carton Ampule		(Write-in, e.o	j. 1 Box of 10) Vials)	
preservative-free? correctional institution block?		No					Light to dark people	-	Glass		Minimum or	dor augntitu		Yes
opioid?		No				Product Colo	r: Light to dark peach		Tube		William Of	uer quantity	f	162
Cannabinoid?		No	Country of Origin	USA			One full bisect and two partial bisect		Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for	140	Country or origin	COA		Product Impri	int: lines on one side, and debossed with 'T' over '376' on other side		Vial Liquid Multi		If Yes, how I	nany of whi	ch package t	vpe?
hospital scanning?	driit dosc for		Is this product covered u	inder the			over 376 on other side	' II	Vial Powder Sql			Each	on paonago i	. , po .
If Unit Dose, indicate NDC here:			Trade Agreements Act (Yes				Vial Powder Multi			Inner/Carton	/Pack	
•									Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS										
					Au	thorized Generic	*If Authorized Generic, other		P⊦	IARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB						section fields are not applicable	Rec. sell unit to c	ustomer?		Rx billing ur	it to pharma	icv:	
II. Generic Equivalent to What Bra		Adderall								T		Each	,.	
								(Write-in, e.g. 1 V	ial)	1		Gram		
		DRUG SUPPI	LY CHAIN SECURITY ACT	(DSCSA) INFOR	RMATION							Milliliter		
Does supplier meet DSCSA defin		rer?	Yes		GLN:	0860000397957			ITEN	I AND PACKING II	NFORMATION 1			
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:				Waight I 5-	Dimensi	ons (US msm	ts.)	Volume	Saleable #
Other exemption - Write in:								.	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	riginal product purcl	hased	Item/Each:	0.11	1.52	1.52	2.69	6.21	1
					direct from m	nfr?						2.00	0.21	·
Is product sold by manufacturer's			Yes											
Has FDA granted waiver/exception	on/exemption for pr						repackaged product	Box/Carton/Bund	lle/					
	on/exemption for pr		Yes				repackaged product	Inner Pack:	lie/					24
Has FDA granted waiver/exception	on/exemption for pr	roduct?	Yes No	NEODMATION			repackaged product		lle/	9.8	6.5	3	191.1	24
Has FDA granted waiver/exception	on/exemption for pr	roduct?	Yes	NFORMATION			repackaged product	Inner Pack: Case:		9.8	6.5	3	191.1	24
Has FDA granted waiver/exception if yes, attach documentation fro	on/exemption for pr om FDA.	GT	Yes No IN AND HIBCC PRODUCT	NFORMATION	Provide source	ce manufacturer for		Inner Pack:		9.8	6.5	3	191.1	24
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure	on/exemption for pr om FDA.	GT Saleable Quantity	Yes No	NFORMATION	Provide source	ce manufacturer for	repackaged product Unit of Use GTIN-14	Inner Pack: Case:		9.8	6.5	3	191.1	24
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure x Item/Each	on/exemption for pr om FDA.	GT	Yes No IN AND HIBCC PRODUCT	NFORMATION	Provide source	ce manufacturer for		Inner Pack: Case:	3	9.8			191.1	
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure	on/exemption for pr om FDA.	GT Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	ce manufacturer for		Inner Pack: Case:		9.8				
Has FDA granted waiver/exception if yes, attach documentation from the state of the	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	N-14 31722164016		Inner Pack: Case:	3	9.8				
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	N-14 31722164016		Inner Pack: Case: Pallet:	3 COST INFORMATION		\	VHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	N-14 31722164016		Inner Pack: Case: Pallet: Regular Cost	COST INFORMATION		Vendor #:	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	N-14 31722164016		Inner Pack: Case: Pallet: Regular Cost	3 COST INFORMATION		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT	NFORMATION	GTII 003	N-14 31722164016		Inner Pack: Case: Pallet: Regular Cost Invoice Cost (WA	COST INFORMATION		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception if yes, attach documentation from the saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT HIBCC		GTII 003	N-14 31722164016 31722164013	Unit of Use GTIN-14	Inner Pack: Case: Pallet: Regular Cost Invoice Cost (WA As of date:	3 COST INFORMATION C) (\$) 8/12/2021		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X	on/exemption for pr om FDA.	GTI Saleable Quantity	Yes No IN AND HIBCC PRODUCT HIBCC		GTII 003	N-14 31722164016 31722164013		Inner Pack: Case: Pallet: Regular Cost Invoice Cost (WA As of date:	3 COST INFORMATION C) (\$) 8/12/2021		Vendor #: Whsl. Code	WHOLESALI		



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION							
Is this product (check all that apply):							
a. Cytotoxic?	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?							
Is the product a CA Prop 65 carcinogen?	x Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?	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? No	identify NFPA Storage Level: NFPA Storage Level:						
(If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No	NFFA Storage Level:						
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	Hazardous Waste Identification						
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	KEIRO O REGISTRI RESTRICTIONS						
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)	Provider Name: DEA #:						
Limited Quantity	Site Enrollment Number assigned NCPDP#:						
Consumer Commodity, ORM-D	by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)							
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);	Postura						
SP#	Registry: No						
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone:						
	Comments						
Is the Product	RETURN INSTRUCTIONS						
Controlled Substance? Yes Controlled Substance Code 1100	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						
ARCOS Reportable? Yes If yes, indicate which: Schedule No. 2 Is it a scheduled listed chemical product?: No							
	Is product returnable for credit:						
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:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only: No	product in certain states? Yes						
Restricted to hospital, clinics, and physician offices only. Restricted from US territories? (explain in comments) No	If so, which states? Other requirements? Comments?						
Comments:	DEA Form 222 or its electronic equivalent is required for all returns in all states.						
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 Part	1301.72.						
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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 Process	sing			
Purchase orders may be accepted by:		Purchase order daily receipt cut off time by supplier				
a. EDI		Cut off time:				
b. Autofax	Fax Number:					
c. Fax	Fax Number:	Shipping lead time of PO: Hours	Days			
d. Phone only	Phone No.:					
e. Supplier Web Site only	Site Address:	Ships same day for next day receipt:				
Minimum Order Quantity:		Ships for second day receipt:				
Supplier's Customer Service Number:		Ships regular ground for 3-10 days receipt:				
Contracted 3PL company / contact #:	Name:					
	Phone:					
Expedited Freight Cha	rges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Pro	cessing			
Expedited freight fees billed with each orde	er:	Overnight receipt available:				
Drop Ship service fee billed with each orde	r:	PO Receipt cut off time:				
Drop Ship miscellaneous fees billed:		Days of week overnight is available:	Monday			
Comments:		,	Tuesday			
			Wednesday			
			Thursday			
			Friday			
		Priority Overnight receipt available:				
Cla	ss of Trade Restriction:	PO Receipt Cut off time:				
No restriction: Select VES if sold to retail of	narmacy, hospitals, clinics and physician offices	Saturday Overnight receipt available:				
Restricted to retail pharmacy only:	larmacy, nospitals, clinics and physician offices	PO Receipt Cut off time:				
Restricted to hospital, clinics, and physician	o offices only:	Phone: Phone #				
Restricted from US territories? (explain in c		Order receipt method: Fax: Fax #:				
Comments:		EDI:				
		Overnight Fees apply:				
		Other fees apply:				
Other Data Inf	ormation Required to Process PO:	Return Instructions				
Patient Procedure Date:		Contact # if product is received damaged:				
Physician Name:		Is product returnable for credit:				
Physician/Clinic Phone #		URL/Link to returns policy:				
Physician State License #						
Physician/Clinic DEA #:		Special regulations or returns requirements for this product in certa	ain states?			
Physician/Clinic Specialty:		If so, which states? Other requirements? Comments?				
	Miscellaneous Notes:					
		ADDITIONAL INFORMATION				
		Is product order for scheduled patient procedure?				
		Is product order for restocking purposes?				
		13 product order for restocking purposes:				